DEPARTMENT OF HEALTH AND HUMAN SERVICES

CENTERS FOR MEDICARE & MEDICAID SERVICES

					AND TRANSMITTAL TE SURVEY AGENCY	ID: MTA3 Facility ID: 00933	
1. MEDICARE/MEDICAID PROVII (L1) 245336 2.STATE VENDOR OR MEDICAID (L2) 655371100	DER NO.	3. NAME AND AD (L3) THE ESTAT (L4) 433 COUNT (L5) DELANO, M	DDRESS OF FAC TES AT DELAN Y ROAD 30	CILITY	(L6) 55328	4. TYPE OF ACTION: 7 (L8) 1. Initial 2. Recertification 3. Termination 4. CHOW 5. Validation 6. Complaint	
5. EFFECTIVE DATE CHANGE OF (L9) 03/01/2017 6. DATE OF SURVEY 10/2 8. ACCREDITATION STATUS: 0 Unaccredited 1 TJC 2 AOA 3 Other 11. LTC PERIOD OF CERTIFICATION	27/2020 (L34) (L10)	7. PROVIDER/SU 01 Hospital 02 SNF/NF/Dual 03 SNF/NF/Distinct 04 SNF	05 HHA 06 PRTF 07 X-Ray 08 OPT/SP	09 ESRD 10 NF 11 ICF/IID 12 RHC	02 (L7) 13 PTIP 22 CLIA 14 CORF 15 ASC 16 HOSPICE	7. On-Site Visit 9. Other 8. Full Survey After Complaint FISCAL YEAR ENDING DATE: (L35) 12/31	
From (a): To (b): 12.Total Facility Beds 13.Total Certified Beds	46 (L18) 46 (L17)	B. Not in Com	quirements	-	And/Or Approved Waivers Of 2. Technical Personne 3. 24 Hour RN 4. 7-Day RN (Rural SI 5. Life Safety Code * Code: A	7. Medical Director	
14. LTC CERTIFIED BED BREAKD 18 SNF 18/19 SNF 46 (L37) (L38)		ICF (L42)	IID (L43)		15. FACILITY MEETS 1861 (e) (1) or 1861 (j) (1):	(L15)	
STATE SURVEY AGENCY REN SURVEYOR SIGNATURE Susie Haben, Unit Super	`	Date :	1/24/2020		18. STATE SURVEY AGENCY Douglas Larson, Enforce	ment Specialist 11/24/2020	
		COMPLETED E	BY HCFA RE	(L19) EGIONAI	OFFICE OR SINGLE S		(L20
19. DETERMINATION OF ELIGIBI _X 1. Facility is Eligible to 2. Facility is not Eligib	Participate		PLIANCE WITH	H CIVIL		ancial Solvency (HCFA-2572) rol Interest Disclosure Stmt (HCFA-1513) re:	
22. ORIGINAL DATE OF PARTICIPATION 07/01/1986 (L24) 25. LTC EXTENSION DATE: (L27)	-	DATE	ENDING DATE		26. TERMINATION ACTION VOLUNTARY 01-Merger, Closure 02-Dissatisfaction W/ Reimburs 03-Risk of Involuntary Terminati 04-Other Reason for Withdrawal	0 INVOLUNTARY 05-Fail to Meet Health/Safety sement 06-Fail to Meet Agreement on OTHER	
		1	(L45)				
28. TERMINATION DATE:	29	. INTERMEDIARY/	CARRIER NO.		30. REMARKS		
	(L28)	01111		(L31)			

32. DETERMINATION OF APPROVAL DATE

(L33)

DETERMINATION APPROVAL

11/06/2020

(L32)

31. RO RECEIPT OF CMS-1539



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered November 24, 2020 CMS Certification Number (CCN): 245336

Administrator The Estates At Delano Llc 433 County Road 30 Delano, MN 55328

Dear Administrator:

The Minnesota Department of Health assists the Centers for Medicare and Medicaid Services (CMS) by surveying skilled nursing facilities and nursing facilities to determine whether they meet the requirements for participation. To participate as a skilled nursing facility in the Medicare program or as a nursing facility in the Medicaid program, a provider must be in substantial compliance with each of the requirements established by the Secretary of Health and Human Services found in 42 CFR part 483, Subpart B.

Based upon your facility being in substantial compliance, we are recommending to CMS that your facility be recertified for participation in the Medicare and Medicaid program.

Effective November 8, 2020 the above facility is certified for:

46 Skilled Nursing Facility/Nursing Facility Beds

Your facility's Medicare approved area consists of all 46 skilled nursing facility beds.

We have recommended CMS approve the waivers that you requested for the following Life Safety Code Requirements: K521.

If you are not in compliance with the above requirements at the time of your next survey, you will be required to submit a Plan of Correction for this deficiency or renew your request for waiver in order to continue your participation in the Medicare and Medicaid Program.

You should advise our office of any changes in staffing, services, or organization, which might affect your certification status. If, at the time of your next survey, we find your facility to not be in substantial compliance your Medicare and/or Medicaid provider agreement may be subject to non-renewal or termination.

Please contact me if you have any questions.

Sincerely,

Douglas Larson, Enforcement Specialist

Minnesota Department of Health

Licensing and Certification Program

Program Assurance Unit

1 Julius Stapson

The Estates At Delano Llc November 24, 2020 Page 2

Health Regulation Division

Telephone: 651-201-4118 Fax: 651-215-9697

Email: doug.larson@state.mn.us

cc: Licensing and Certification File



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered November 24, 2020

Administrator The Estates At Delano Llc 433 County Road 30 Delano, MN 55328

RE: CCN: 245336

Cycle Start Date: September 3, 2020

Dear Administrator:

On September 23, 2020, we notified you a remedy was imposed. On October 27, 2020 the Minnesota Department(s) of Health and Public Safety completed a revisit to verify that your facility had achieved and maintained compliance. We have determined that your facility has achieved substantial compliance as of November 8, 2020.

As authorized by CMS the remedy of:

• Discretionary denial of payment for new Medicare and Medicaid admissions effective October 23, 2020 be discontinued as of November 8, 2020. (42 CFR 488.417 (b))

However, as we notified you in our letter of September 23, 2020, in accordance with Federal law, as specified in the Act at § 1819(f)(2)(B)(iii)(I)(b) and § 1919(f)(2)(B)(iii)(I)(b), we notified you that your facility is prohibited from conducting Nursing Aide Training and/or Competency Evaluation Programs (NATCEP) for two years from October 23, 2020. This does not apply to or affect any previously imposed NATCEP loss.

The CMS Region V Office may notify you of their determination regarding any imposed remedies.

Your request for a continuing waiver involving the deficiency(ies) cited under K521 at the time of the September 3, 2020 standard survey has been forwarded to CMS for their review and determination. Your facility's compliance is based on pending CMS approval of your request for waiver.

Feel free to contact me if you have questions.

Sincerely,

Douglas Larson, Enforcement Specialist

The Estates At Delano Llc November 24, 2020 Page 2

Minnesota Department of Health Licensing and Certification Program Program Assurance Unit Health Regulation Division

Telephone: 651-201-4118 Fax: 651-215-9697

Email: doug.larson@state.mn.us

cc: Licensing and Certification File

DEPARTMENT OF HEALTH AND HUMAN SERVICES

CENTERS FOR MEDICARE & MEDICAID SERVICES

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PART I - TO RE (OMPLETED R	V THE STATE	SHDVEV ACENCY

Facility ID: 00933

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5. EFFECTIVE DATE CHANGE OF (L9) 03/01/2017	OWNERSHIP	7. PROVIDER/SU	JPPLIER CATEO	GORY 09 ESRD	02 (L7) 13 PTIP 22 CLIA	7. On-Site Visit 8. Full Survey A	
6. DATE OF SURVEY 09/03 8. ACCREDITATION STATUS: 0 Unaccredited 1 TJC 2 AOA 3 Other	/ 2020 (L34) (L10)	02 SNF/NF/Dual 03 SNF/NF/Distinct 04 SNF	06 PRTF 07 X-Ray 08 OPT/SP	10 NF 11 ICF/IID 12 RHC	14 CORF 15 ASC 16 HOSPICE	FISCAL YEAR EN	NDING DATE: (L35)
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14 LTG CERTIFIED DED DREAMDO	WAI	Requirements	and/or Applied	waivers:	* Code: B *	(L12)	
14. LTC CERTIFIED BED BREAKDO 18 SNF 18/19 SNF 46	WN 19 SNF	ICF	IID		15. FACILITY MEETS 1861 (e) (1) or 1861 (j) (1):	(L15)	
(L37) (L38)	(L39)	(L42)	(L43)				
16. STATE SURVEY AGENCY REM	ARKS (IF APPLICA	ABLE SHOW LTC CA	ANCELLATION	DATE):			
17. SURVEYOR SIGNATURE		Date :			18. STATE SURVEY AGENCY	Y APPROVAL	Date:
Lynn Moeller, HFE NE		1	0/07/2020	(L19)	Alison Helm, Enforce	ement Specialis	11/04/2020 (L20
PAI	RT II - TO BE	COMPLETED I	BY HCFA RI	EGIONAI	OFFICE OR SINGLE S	STATE AGENCY	
DETERMINATION OF ELIGIBIL	articipate		MPLIANCE WIT HTS ACT:	H CIVIL	21. 1. Statement of Fina 2. Ownership/Contr 3. Both of the Abov	ol Interest Disclosure S	
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OF PARTICIPATION 07/01/1986	BEGINNING		ENDING DA		VOLUNTARY 00-Merger, Closure	<u>INVO</u>	LUNTARY to Meet Health/Safety
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25. LTC EXTENSION DATE:	27. ALTERNATI				03-Risk of Involuntary Termination 04-Other Reason for Withdrawal	OTHE	<u>R</u> vider Status Change
(L27)		n of Admissions: uspension Date:	(L44) (L45)			00-Act	-
28. TERMINATION DATE:	29). INTERMEDIARY	/CARRIER NO.		30. REMARKS		
		01111					
	(L28)	01111		(L31)			
31. RO RECEIPT OF CMS-1539	32	2. DETERMINATION	N OF APPROVAL	L DATE			
	(L32)			(L33)	DETERMINATION APP	ROVAL	



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered September 23, 2020

Administrator The Estates At Delano LLC 433 County Road 30 Delano, MN 55328

RE: CCN: 245336

Cycle Start Date: September 3, 2020

Dear Administrator:

On September 3, 2020, a survey was completed at your facility by the Minnesota Department(s) of Health and Public Safety to determine if your facility was in compliance with Federal participation requirements for skilled nursing facilities and/or nursing facilities participating in the Medicare and/or Medicaid programs.

This survey found the most serious deficiencies in your facility to be widespread deficiencies that constitute no actual harm with potential for more than minimal harm that is not immediate jeopardy (Level F), as evidenced by the electronically delivered CMS-2567, whereby significant corrections are required.

REMEDIES

As a result of the survey findings and in accordance with survey and certification memo 16-31-NH, this Department recommended the enforcement remedy(ies) listed below to the CMS Region V Office for imposition. The CMS Region V Office concurs and is imposing the following remedy and has authorized this Department to notify you of the imposition:

- Discretionary Denial of Payment for new Medicare and/or Medicaid Admissions, Federal regulations at 42 CFR § 488.417(a), effective October 23, 2020.
- Directed plan of correction (DPOC), Federal regulations at 42 CFR § 488.424. Please see electronically attached documents for the DPOC.

The CMS Region V Office will notify your Medicare Administrative Contractor (MAC) that the denial of payment for new admissions is effective October 23, 2020. They will also notify the State Medicaid Agency that they must also deny payment for new Medicaid admissions effective October 23, 2020.

You should notify all Medicare/Medicaid residents admitted on, or after, this date of the restriction. The remedy must remain in effect until your facility has been determined to be in substantial compliance or your provider agreement is terminated. Please note that the denial of payment for

new admissions includes Medicare/Medicaid beneficiaries enrolled in managed care plans. It is your obligation to inform managed care plans contracting with your facility of this denial of payment for new admissions.

This Department is also recommending that CMS impose:

• Civil money penalty (42 CFR 488.430 through 488.444). You will receive a formal notice from the CMS RO only if CMS agrees with our recommendation.

NURSE AIDE TRAINING PROHIBITION

Please note that Federal law, as specified in the Act at §§ 1819(f)(2)(B) and 1919(f)(2)(B), prohibits approval of nurse aide training and competency evaluation programs and nurse aide competency evaluation programs offered by, or in, a facility which, within the previous two years, has operated under a § 1819(b)(4)(C)(ii)(II) or § 1919(b)(4)(C)(ii) waiver (i.e., waiver of full-time registered professional nurse); has been subject to an extended or partial extended survey as a result of a finding of substandard quality of care; has been assessed a total civil money penalty of not less than \$11,160; has been subject to a denial of payment, the appointment of a temporary manager or termination; or, in the case of an emergency, has been closed and/or had its residents transferred to other facilities.

If you have not achieved substantial compliance by October 23, 2020, the remedy of denial of payment for new admissions will go into effect and this provision will apply to your facility. Therefore, The Estates At Delano LLC will be prohibited from offering or conducting a Nurse Aide Training and/or Competency Evaluation Program (NATCEP) for two years from October 23, 2020. You will receive further information regarding this from the State agency. This prohibition is not subject to appeal. Further, this prohibition may be rescinded at a later date if your facility achieves substantial compliance prior to the effective date of denial of payment for new admissions. However, under Public Law 105-15, you may contact the State agency and request a waiver of this prohibition if certain criteria are met.

ELECTRONIC PLAN OF CORRECTION (ePOC)

Within ten (10) calendar days after your receipt of this notice, you must submit an acceptable ePOC for the deficiencies cited. An acceptable ePOC will serve as your allegation of compliance. Upon receipt of an acceptable ePOC, we will authorize a revisit to your facility to determine if substantial compliance has been achieved. The failure to submit an acceptable ePOC can lead to termination of your Medicare and Medicaid participation (42 CFR 488.456(b)).

To be acceptable, a provider's ePOC must include the following:

- How corrective action will be accomplished for those residents found to have been affected by the deficient practice.
- How the facility will identify other residents having the potential to be affected by the same deficient practice.

- What measures will be put into place, or systemic changes made, to ensure that the deficient practice will not recur.
- How the facility will monitor its corrective actions to ensure that the deficient practice is being corrected and will not recur.
- The date that each deficiency will be corrected.
- An electronic acknowledgement signature and date by an official facility representative.

DEPARTMENT CONTACT

Questions regarding this letter and all documents submitted as a response to the resident care deficiencies (those preceded by a "F" tag), and emergency preparedness deficiencies (those preceded by an "E" tag), i.e., the plan of correction should be directed to:

Susie Haben, Unit Supervisor
St. Cloud A Survey Team
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
3333 West Division Street, Suite 212
St. Cloud, Minnesota 56301
Email: susie.haben@state.mn.us

Phone: 320-223-7356

PRESUMPTION OF COMPLIANCE - CREDIBLE ALLEGATION OF COMPLIANCE

The facility's ePoC will serve as your allegation of compliance upon the Department's acceptance. In order for your allegation of compliance to be acceptable to the Department, the ePoC must meet the criteria listed in the plan of correction section above. You will be notified by the Minnesota Department of Health - Health Regulation Division staff and/or the Department of Public Safety, State Fire Marshal Division staff, if your ePoC for their respective deficiencies (if any) is acceptable.

VERIFICATION OF SUBSTANTIAL COMPLIANCE

Upon receipt of an acceptable ePoC, a Post Certification Revisit (PCR), of your facility will be conducted to validate that substantial compliance with the regulations has been attained in accordance with your verification.

If substantial compliance has been achieved, certification of your facility in the Medicare and/or Medicaid program(s) will be continued and remedies will not be imposed. Compliance is certified as of the latest correction date on the approved ePoC, unless it is determined that either correction actually occurred between the latest correction date on the ePoC and the date of the first revisit, or correction occurred sooner than the latest correction date on the ePoC.

FAILURE TO ACHIEVE SUBSTANTIAL COMPLIANCE BY THE SIXTH MONTH AFTER THE LAST DAY OF THE SURVEY

We will also recommend to the CMS Region V Office and/or the Minnesota Department of Human Services that your provider agreement be terminated by March 3, 2021 if your facility does not achieve substantial compliance. This action is mandated by the Social Security Act at § 1819(h)(2)(C) and 1919(h)(3)(D) and Federal regulations at 42 CFR § 488.412 and § 488.456.

Please note that this notice does not constitute formal notice of imposition of alternative remedies or termination of your provider agreement. Should the Centers for Medicare & Medicaid Services determine that termination or any other remedy is warranted, it will provide you with a separate formal notification of that determination.

APPEAL RIGHTS

If you disagree with this action imposed on your facility, you or your legal representative may request a hearing before an administrative law judge of the Department of Health and Human Services, Departmental Appeals Board (DAB). Procedures governing this process are set out in 42 C.F.R. 498.40, et seq. You must file your hearing request electronically by using the Departmental Appeals Board's Electronic Filing System (DAB E-File) at https://dab.efile.hhs.gov no later than sixty (60) days after receiving this letter. Specific instructions on how to file electronically are attached to this notice. A copy of the hearing request shall be submitted electronically to:

Tamika.Brown@cms.hhs.gov

Requests for a hearing submitted by U.S. mail or commercial carrier are no longer accepted as of October 1, 2014, unless you do not have access to a computer or internet service. In those circumstances you may call the Civil Remedies Division to request a waiver from e-filing and provide an explanation as to why you cannot file electronically or you may mail a written request for a waiver along with your written request for a hearing. A written request for a hearing must be filed no later than sixty (60) days after receiving this letter, by mailing to the following address:

Department of Health & Human Services
Departmental Appeals Board, MS 6132
Director, Civil Remedies Division
330 Independence Avenue, S.W.
Cohen Building – Room G-644
Washington, D.C. 20201
(202) 565-9462

A request for a hearing should identify the specific issues, findings of fact and conclusions of law with which you disagree. It should also specify the basis for contending that the findings and conclusions are incorrect. At an appeal hearing, you may be represented by counsel at your own expense. If you have any questions regarding this matter, please contact Tamika Brown, Principal Program Representative by phone at (312) 353-1502 or by e-mail at Tamika.Brown@cms.hhs.gov.

INFORMAL DISPUTE RESOLUTION (IDR) / INDEPENDENT INFORMAL DISPUTE RESOLUTION (IIDR)

In accordance with 42 CFR 488.331, you have one opportunity to question cited deficiencies through an informal dispute resolution process. You are required to send your written request, along with the specific deficiencies being disputed, and an explanation of why you are disputing those deficiencies, to:

> Nursing Home Informal Dispute Process Minnesota Department of Health Health Regulation Division P.O. Box 64900 St. Paul, Minnesota 55164-0900

This request must be sent within the same ten days you have for submitting an ePoC for the cited deficiencies. All requests for an IDR or IIDR of federal deficiencies must be submitted via the web at: https://mdhprovidercontent.web.health.state.mn.us/ltc_idr.cfm

You must notify MDH at this website of your request for an IDR or IIDR within the 10 calendar day period allotted for submitting an acceptable electronic plan of correction. A copy of the Department's informal dispute resolution policies are posted on the MDH Information Bulletin website at: https://www.health.state.mn.us/facilities/regulation/infobulletins/ib04 8.html

Please note that the failure to complete the informal dispute resolution process will not delay the dates specified for compliance or the imposition of remedies.

Questions regarding all documents submitted as a response to the Life Safety Code deficiencies (those preceded by a "K" tag) i.e., the plan of correction, request for waivers, should be directed to:

> Mr. Tom Linhoff, Fire Safety Supervisor **Health Care Fire Inspections** Minnesota Department of Public Safety State Fire Marshal Division 445 Minnesota Street, Suite 145 St. Paul, Minnesota 55101-5145 Email: tom.linhoff@state.mn.us Telephone: (651) 430-3012

Fax: (651) 215-0525

Feel free to contact me if you have questions.

Sincerely,

alison Helm

Alison Helm, Enforcement Specialist Licensing and Certification

Minnesota Department of Health

P.O. Box 64970 Saint Paul, Minnesota 55164-0970

Phone: 651-201-4206

Email: alison.helm@state.mn.us

PRINTED: 10/08/2020 FORM APPROVED OMB NO. 0938-0391

	FOF DEFICIENCIES OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MUL A. BUILD		LE CONSTRUCTION		E SURVEY PLETED
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		245336	B. WING			09/	03/2020
NAME OF F	PROVIDER OR SUPPLIER				STREET ADDRESS, CITY, STATE, ZIP CODE		
THE EST	ATES AT DELANO L	LC			33 COUNTY ROAD 30		
					DELANO, MN 55328		
(X4) ID		ATEMENT OF DEFICIENCIES	ID		PROVIDER'S PLAN OF CORRECTIO		(X5)
PREFIX TAG		Y MUST BE PRECEDED BY FULL SC IDENTIFYING INFORMATION)	PREFI TAG		(EACH CORRECTIVE ACTION SHOULD CROSS-REFERENCED TO THE APPROP		COMPLETION DATE
		,			DEFICIENCY)		
E 000	Initial Comments		ΕC	000			
	A survey for compl	liance with CMS Appendix Z					
		edness Requirements, was					
	conducted on 8/31/	/20-9/3/20, during a					
		ey. The facility is in compliance					
		Z Emergency Preparedness					
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		gh 9/3/20, a standard					
		ey was conducted at your t investigation was also					
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	Facilities.	-					
	TI 6 II	Literary Complete to					
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	H5336045C	ED.					
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	H5336048C						
	H5336049C						
	The facility's plan of	f correction (POC) will serve					
		of compliance upon the					
		ptance. Because you are					
		our signature is not required					
	at the bottom of the	e first page of the CMS-2567					
		ic submission of the POC will					
	be used as verificat	tion of compliance.					
	Unon receipt of an	acceptable electronic POC, an					
		ur facility may be conducted to					
		antial compliance with the					
		en attained in accordance with					
LABORATOR'	_	DER/SUPPLIER REPRESENTATIVE'S SIG	NATURE		TITLE		(X6) DATE

Electronically Signed 10/02/2020

Any deficiency statement ending with an asterisk (*) denotes a deficiency which the institution may be excused from correcting providing it is determined that other safeguards provide sufficient protection to the patients. (See instructions.) Except for nursing homes, the findings stated above are disclosable 90 days following the date of survey whether or not a plan of correction is provided. For nursing homes, the above findings and plans of correction are disclosable 14 days following the date these documents are made available to the facility. If deficiencies are cited, an approved plan of correction is requisite to continued program participation.

	FOF DEFICIENCIES OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	l ` ′	PLE CONSTRUCTION G	(X3) DATE S COMPLE	
		245336	B. WING _		C 09/03 /	/2020
	PROVIDER OR SUPPLIER	LC		STREET ADDRESS, CITY, STATE, ZIP CODE 433 COUNTY ROAD 30 DELANO, MN 55328		
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F 000		ge 1	F 00	0		
	your verification. Care Plan Timing a CFR(s): 483.21(b)(F 65	7	10)/9/20
	§483.21(b)(2) A colbe- (i) Developed withir the comprehensive (ii) Prepared by an includes but is not I (A) The attending p (B) A registered nu resident. (C) A nurse aide wi resident. (D) A member of fo (E) To the extent properties that the explanation must medical record if the and their resident resident resident's care pland (F) Other appropriate disciplines as deteror as requested by (iii)Reviewed and reteam after each assessments. This REQUIREMED by: Based on observative review, the facility following discharge Physical Therapy for the side of the comprehensive and assessments.	interdisciplinary team, that imited to inysician. rse with responsibility for the th responsibility for the th responsibility for the od and nutrition services staff. racticable, the participation of the resident's representative(s). It is included in a resident's representative is determined the development of the the development of the included by the resident's needs the resident. The staff or professionals in the mined by the resident's needs the resident. The staff or professionals in the staff or professionals in the staff or professionals in the resident. The staff or professionals in the staff or professionals in the resident.		F657: Care Plan Timing and Revis Immediate Corrective Action: R6 therapy recommendations were reviewed. R6 is currently being assiby therapy to determine appropriate	essed	

	OF DEFICIENCIES OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	l ` ′		E CONSTRUCTION		SURVEY PLETED
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F 657	Findings include: R6's annual Minima 7/10/20, indicated Ir right-side weaknes depression, anxiety MDS further indical cognitively impaired toileting, was not st with staff assistand standing position a transfers. A document titled If Summary, electron dated 8/14/20, indicat at all times while ou document further ir use of the AFO. R6's care plan, prir did not include ROI AFO. On 8/31/20, at 4:55 in their room, sitting ankle-foot orthosis The AFO was obse R6's nightstand. On 9/01/20, at 1:51 out of their room by laying on the floor r nightstandat 4:05 p.m. R6 wa facility via transport	um Data Set (MDS) dated R6 had diagnoses including s, end-stage renal disease, y, and speech difficulty. The ted R6 was severely d, needed extensive assist for teady and only able to stabilize the when moving from seated to a surface to surface. Physical Therapy PT Discharge ically signed by Staff-E and cated R6 should have AFO on at of bed and transferring. The indicated staff were trained in the Mexercises or application of the year of the wheelchair, with (AFO) splint not on R6's foot. The erved laying on the floor next to the residents as observed returning to the transit of the	F6	557	of AFO. The recommendations will added to the care plan and point of task list. Action as it is applied to others: Restorative nursing services policy reviewed and remains current Care Planning policy was reviewed remains current Therapy recommendations for all residents were reviewed from the pmonths to ensure that all recommendations were added to the plan and point of care task list. Communication process between the and clinical was reviewed and revise Reeducation has been initiated with therapy and clinical staff regarding communication, care planning, implementing and completing there are commendations. Date of completion: _10/9/2020 Recurrence will be prevented by: Audits of 3 therapy recommendation be completed weekly x 4 then mon months to ensure that the therapy recommendations are care planned are being followed. The results of the audits will be reviewed with the facion QAPI Committee for input on the new increase, decrease or discontinue the audits. The correction will be monitored by DON and/or designee	was and ast 3 ne care herapy ed. n upy ans will thly x2 d and these lity eed to the	

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F 657	On 9/02/20, at 8:0 -B with NA-C were incontinence care: NA-B and NA-C d or shoes. NA-B pl. NA-B then transfe gait belt and pivot -at 8:26 a.m., NA-R6 wears gripper is responsible for lower extremity. Neen an order for -at 9:10 a.m., NA-the NA's will do ra on the right arm at R6's lower extrem NA's do not do an -at 9:53 a.m., RN-there were no dev day, and R6 wore easier for R6 to drin place. RN-D ver ROM exercises at AFO in R6's chart On 9/03/20, at 10: AFO recommendation foot lift, and to mat On 9/2/20, at 12:4 and verified there AFO to R6's Right sheet nor in R6's con 9/2/20, at 8:40	2 a.m., Nursing Assistant (NA) e observed to perform and dressed R6 for the day. id not offer to put on R6's AFO acced blue gripper socks on R6. rred R6 to the wheelchair with a transfer. B was interviewed and stated socks all the time and therapy placing the AFO to R6's right NA-B further stated she had not putting on the AFO. B was interviewed and stated nge of motion (ROM) exercises a R6 allows. NA-B further stated ities were "pretty good" and y ROM on them. D was interviewed and stated ices R6 needed to wear every gripper socks because it was ag the right foot without a shoe rified there was no order for and no order for application of an 23 a.m., Staff-E stated R6's ations were in place to promote intain ROM. 7 p.m., RN-E was interviewed was no directive to apply an allower extremity on the NA care	F 6	357		

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would be receiventies accrecommendation on 9/2/20, at 8 Therapist, state maintenance properties to follow therapist of following the fol	expectation would be that R6 ving ROM to both upper and lower cording to previous therapy ons. 8:44 a.m., Staff-E, Physical ed R6 is on a functional program and he would expect staff by recommendations. 8:47 p.m., RN-E was interviewed rapy staff gives a copy of their ons to RN-E to add to the NA care at of care task list in the resident's RN-E stated the therapy would resident's care plan. RN-E stated being any therapy ons for R6 during the month of verified there are no ons from therapy for R6 in the staff a book for the month of August. Perified there was no directive to to R6's Right lower extremity on the nor in R6's care plan. RN-E stated expectation to receive a copy of commendations. RN-E verified here are at the bottom of the document T Functional Maintenance Program outlining therapy	f e d	7		

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F 657	The Monarch Healt titled Care Planning care plan is to be m	hcare Management policy g, revised 6/2019, indicated the nodified and updated as the needs of the resident	F 65	7	
	ADL Care Provided CFR(s): 483.24(a)(§483.24(a)(2) A resout activities of dail services to maintain personal and oral had This REQUIREMED by: Based on observareview, the facility for with grooming for 1 reviewed for activiting who were depended. R11's Minimum Daindicated severe codependent with translift and always inco Braden scale 14, work skin breakdown. R11's bladder asset identified, R11 was retraining. R11's Care Area Assidentified, R11 trigged during assessments.	sident who is unable to carry y living receives the necessary n good nutrition, grooming, and ygiene; NT is not met as evidenced tion, interview, and document ailed to provide assistance of 2 residents (R11), es of daily living (ADLs) and	F 67	F677: ADL Care Provided for Dependen Residents Immediate Corrective Action: A bowel and bladder assessment was completed for R11 and a toileting plan was implemented and care planned. R11 is being assisted with toileting per her plan. Action as it is applied to others: Urinary Continence and Incontinence-Assessment and Management Policy was reviewed and remains current. A bowel and bladder assessment were completed for residents to determine an appropriate toileting plan for each. These plans were implemented, and care planned. Education has been initiated with nursing staff regarding assessing bowel and bladder and initiating individualized toilet plans. Education was implemented for al	

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F 677	diagnosis Type 1 Dabuse, ataxic (invomuscle weakness, Other symptoms at functions and awar unspecified, abnored History of transient (mini-stroke), and owithout residual de During observation nursing assistant (lassistance in R11's laying R11 down at NA-B came in to as hooked up to mach moving resident frowas laid down, briewet. NA-A removed sheet, which was laid her wheelchair NA-A cleaned R11 skin while NA-B will NA-B will NA-B will NA-B will NA-B will NA-B will NA-A stated R11's coccyx appeared stool. R11's skin at wiped area. NA's opropped up her leg their hands and ex During interview or stated staff usually about every 2.5-3 in paper to remind her	ted 7/24/20, indicated pertinent Diabetes Mellitus, Alcohol fluntary movement) gait, Poly neuropathy (nerve pain), and signs involving cognitive eness, Disorientation malities of gait and mobility, eschemic attacks (TIA) cerebral infarction (stroke) ficits. If on 9/1/20, at 1:23 p.m. NA)-A requested transfer are not checking/changing brief. Easist with transfer. Hoyer was nine both aides assisted om wheelchair to bed. W R11 and pants were identified as a R11's soiled clothing. Hoyer and pants were identified as a R11's soiled clothing. Hoyer and pants was not red, but the lightly pink after removing ped down wheelchair cushion. skin was not red, but the lightly pink after removing opeared to blanched when NA ompleted R11's cares and as with a pillow before washing iting the room. In 9/1/20, at 1:30 p.m., NA-B check/change R11's brief nour, and tries to write down on erself of next changes. NA-B usually had large amount of	F 6	nursing staff regarding following providing toileting/incontinence Date of completion: 10/9/2020 Recurrence will be prevented Audits of 3 toileting plans will completed weekly x 4 then menths to ensure that resider assisted with toilet use/incont per their individualized plan. of these audits will be reviewed facility QAPI Committee for inneed to increase, decrease of the audits. The correction will be monitor or/and designee	by: be onthly x2 ots are being inent cares The results ed with the uput on the r discontinue	

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F 677	stated R11 usually and always a check R11 needed assist transfers. NA-A indineeds to urinate or NA-A further stated urine in brief when and the last brief of this morning at 8:00 time to get to her, the and reposition her and reposition her and reposition R11 event in the wheelchanget back in bed after stated if R11 did not 1:30 p.m. it would risk for skin break of During observation laying on left side, urine. Door was ophallway. During observation and NA-C were in Ingetting R11 in wheelstated R11 had a dR11 requested to get the stated to get the stated to get the stated to get the stated R11 had a dR11 requested R11 had	in 9/1/20, at 1:50 p.m. NA-A lays down after every meal is and change. NA-A stated of two and a Hoyer lift for all licated R11 cannot tell if she have a bowel movement. If R11 had large amount of R11 laid down this afternoon hange and repositioned was 0 am, as they did not have but they usually check, change, every 2-3 hours. In 9/1/20, at 2:53 p.m. RN)-C stated staff try to ry hour or so and checked for C further stated R11 likes to be in for meals and then likes to be in for meals and then likes to be exact meal. RN-C further of get toileted from 8:00 a.m. to not be acceptable, as R11 is at down. In 9/2/20, at 7:04 a.m. R11 room has strong smells of the can smell urine from In 9/2/20, at 7:25 a.m. NA-B R11's room to assist with elchair for breakfast. NA-B lay brief and offered a bedpan. Bet off bedpan and would try	F 6	77		
	again later. NA-B a cleaned peri area v skin observed on c Hoyer sheet put un	pet off bedpan and would try issisted R11 off bedpan, with wipe, no redness, or pink occyx, new brief applied. Ider R11 and R11 rolled herself llow clothing adjustment. R11				

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F 677	and NA-C. During observation noted to be laying off. R11's rooms since since before R11 wants to lay down. I foot of R11. R11 to bed since before R11 wanted help. I laying R11 in bed. I small wet brief with she has a history of practitioner. Period observed. During interview or stated R11 brief was check and change. R11 at 3:30 a.m. a before getting her work of the since was sitting in her work as in her wheelch	elchair via Hoyer lift by NA-B on 9/2/20, at 9:46 a.m. R11 on right side in bed with lights mells of urine. I and interview on 9/2/20, at and interview on 9/2/20, at and interview on 9/2/20, at and interview on the delchair, stated she Urine odor noted when within stated she had not been back a lunch. Surveyor notified RN-D RN-D called for assistance with bed. NA-C assisted RN-D with RN-D stated resident had a scant amount of blood, which of and will notify the nurse are performed, no redness on 9/2/20, at 2:38 p.m. NA-B as usually "soaked" with each as usually "soaked" with each and then again, at 5:30 a.m. up for breakfast this morning.		7		
	stated R11's asses	n 9/3/20, at 9:35 a.m., RN-B esments directed staff to check bout every 2-2.5 hours and it is				

STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA AND PLAN OF CORRECTION IDENTIFICATION NUMBER:		(X2) MULTIPLE CONSTRUCTION A. BUILDING			(X3) DATE SURVEY COMPLETED		
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F 677	down of her skin. R for repositioning an unable to do this ca During observation noted to be sitting u noted, call light in re watching television. During interview on stated R11 was rep this was the first tim NA-A stated no red R11's coccyx. NA-A large amount of urin sheet indicates repe every 2-3 hours. R1 wheelchair for lunch During document re AM observed NA-A R11 brief changed 7:15 a.m. Policy Urinary Cont Incontinence-Asses dated 9/2010 "As in remains incontinent causes of incontine toileting plan. As ap the category and ca will provide schedul or other intervention incontinence."	beyond that to prevent break N-B stated R11 relies on staff d bladder/bowel care as R11 re for herself. on 9/3/20, at 10:28 a.m. R11 re in wheelchair. No urine odor each. R11 appeared to be 9/3/20, at 10:46 a.m. NA-A ositioned last at 7:15 a.m. and re she could get back to her. ress or skin breakdown of a described R11's brief had re. NA-A stated R11's care ositioning should occur about 1 was transferred back into red. eview on 09/03/20, at 10:56 rescribed and got up for breakfast at sinence and sement and Management redicated, and if the individual at despite treating transient rece, the staff will initiate a propriate, based on assessing red toileting, prompted voiding,	F 6				10/9/20
	CFR(s): 483.25(b)(гО	00			10/9/20

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F 686	§483.25(b) Skin Int §483.25(b)(1) Pres Based on the compresident, the facility (i) A resident receive professional standar pressure ulcers and ulcers unless the indemonstrates that (ii) A resident with necessary treatme with professional signature promote healing, promote heali	regrity sure ulcers. orehensive assessment of a must ensure that- ves care, consistent with ards of practice, to prevent d does not develop pressure ndividual's clinical condition they were unavoidable; and pressure ulcers receives nt and services, consistent tandards of practice, to revent infection and prevent eveloping. NT is not met as evidenced tion, interview, and document failed to implement care plan event pressure ulcers for 2 of 3), who were dependent on staff mum Data Set (MDS) dated 1 had severe cognitive ed extensive assistance with ent on staff to turn and oed and in wheelchair. seed 4/7/20, identified R1 was ulcers related to impaired hrive, iron deficiency anemia,	F 680	F686 Treatment services to prever pressure ulcers Immediate Corrective Action: R1 air mattress was assessed by th Maintenance Director and is function Orders to assess mattress function been added. R1 was reviewed and assessed for risk of pressure ulcers to determine repositioning schedule plan has been updated to reflect proulcer interventions including reposit Resident is being turned and reposit Resident is being turned and reposit Resident is being turned and remains control of the residents with air mattresses reviewed to ensure air mattress is functioning and orders were entered regarding monitoring the air mattresses residents have been assessed for pressure ulcers and to determine	ne onal. have s and e. Care essure tioning. itioned ment were d ss.		

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F 686	upward towards ce During observation assistant (NA)-B tradip in the center of about one and a hainches high and 6 in stated the air mattrasank in and was co During observation 10:26 a.m. NA-B aridentified R1's air mosetting for her weig would be contacted R1 should not sink cocooned into the rincrease R1's risk to the word of	at 9/2/20, at 8:46 a.m. nursing ansferred R1 out of bed, noted the bed. NA-B stated it was all feet across the bed, 10 nches deep. NA-B further ess had always done that, R1 cooned in the air mattress. and interview on 9/2/20 at not registered nurse (RN)-B nattress was set at the correct ht. RN-B stated maintenance to check the air mattress as into a pocket and be nattress, because that would o develop a pressure sore. on 9/2/20, at 1:12 p.m. or stated the air mattress was r loss light on but the alarm did arm had been set to silent so that there was a problem with	Fé	386	repositioning schedule when applic Residents care plans have been up to reflect pressure ulcer prevention interventions including repositioning where applicable. Reeducation has been initiated with nursing staff regarding air mattress following and providing pressure ulprevention interventions, including repositioning. Date of completion:10/9/2020 Recurrence will be prevented by: Audits of 3 pressure ulcer intervent will be completed weekly x 4 then rx2 months to ensure that residents being assisted with their interventions/repositioning per their individualized plan and that air mattare functional. The results of these will be reviewed with the facility QA Committee for input on the need to increase, decrease or discontinue taudits. The correction will be monitored by and/or designee	ions nonthly are tresses audits PI	
	indicated, R11 has impairment, total de has limited range o R11's Braden scale R11's Braden score	ta Set (MDS) dated 7/24/20, moderate cognition ependent with transfers. R11 f motion to bilateral lower legs. ed dated 7/24/20, indicated, e of 14, which indicates a kin breakdown. R11 reposition					

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F 686	schedule indicates needed. R11's Care plan da foot drop, abnorma weakness, and alchistory of falls. Hoy transfers. During observation transferred to bed nursing assistance change nursing assightly pink coccypalpating after hav denies any observabreakdown. During interview or stated R11 care sh should occur every care sheets indicate and Hoyer lift. During interview or indicated R11 usua with assist of two a indicated they report this morning and dher. NA-A stated R reposition every 2-During interview or registered nurse (F reposition R11 abor R11's dependence was unsure what F	every 2.5-3 hours or as ated 7/24/20, diagnosis of right al gait, ataxic gait, muscle ohol encephalopathy with ver with two assists for all a on 9/1/20, at 1:23 p.m. R11 with the assitance of two and a Hoyer lift. During brief sistant (NA)-A indicated a k, but skin blanches when ing a bowel movement. NA-A ation of red skin or skin a 9/1/20, at 1:30 p.m. NA-B seets indicated repositioning a 2.5-3 hours. NA-B stated R11 te transfer with assist of two a 9/1/20, at 1:50 p.m. NA-A ally lays down after every meal and the Hoyer lift. NA-A sitioned R11 last at 8:00 a.m. id not have time get back to care sheets indication to	F 68	36		

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION (X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:		l ` ′	PLE CONSTRUCTION G	(X3) DATE SURVEY COMPLETED		
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F 686	was sitting in her w room having break! During observation was in her wheelch in reach. No observation buring interview on indicated on admission tolerance, bowel, a then identifies how repositioned and to resident's cares by plans. RN-E stated indicated every 2-2 could not move her staff for repositioning for R11 indicated a repositioned within confirmed the imporprevent pressure sedown from forming. During observation continues to be sittle odor noted, call light to be watching televalue buring interview on stated R11 was reposition was the first tim NA-A stated no red R11's coccyx. NA-A	on 9/3/20, at 8:30 a.m. R11 heelchair in the the dining fast. on 9/3/20, at 9:03 a.m. R11 air sitting in her room, call light ration of position change. 9/3/20, at 9:35 a.m. RN-E sion all residents get a tissue and bladder evaluation, which often residents should be ileted. Staff can identify their care sheets and care R11 repositioning schedule 5 hours. RN-E indicated R11 self and was dependent on ag. RN-E stated assessments risk of skin break down if not her timeframe. RN-E rance to reposition R11 to ores and other skin break on 9/3/20, at 10:28 a.m. R11 ag in wheelchair. No urine at in reach. Resident appears	F 68			

STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA AND PLAN OF CORRECTION IDENTIFICATION NUMBER:		(X2) MULTIPLE CONSTRUCTION A. BUILDING				(X3) DATE SURVEY COMPLETED	
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F 686		ge 14 are sheet dated 9/3/20, at d at 7:15 a.m. R11 transferred	F 6	86			
F 688 SS=D	Policy Pressure Uld dated 7/17, indicate resident-centered of based on the risk far assessments, the of resident's overall of resident's stated wi interventions must be recognized standar the interventions mi plan must be modified changes, or if curre inadequate."	are plan and interventions actors identified in the ondition of the skin, the inical condition, and the shes and goals. a) The be based on current, ds of care. b) The effects of ust be evaluated. c) The Care ied as the resident's condition nt intervention are deemed	F 6	88			10/9/20
	resident who enters range of motion door range of motion unl	acility must ensure that a the facility without limited es not experience reduction in ess the resident's clinical ates that a reduction in range					
	motion receives appearvices to increase	ident with limited range of propriate treatment and erange of motion and/or to rease in range of motion.					
	receives appropriat assistance to maint the maximum pract	ident with limited mobility e services, equipment, and ain or improve mobility with icable independence unless a y is demonstrably unavoidable.					

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F 688	This REQUIREMENT by: Based on observatoreview, the facility faservices were provimotion (ROM) and 1 of 1 resident (R6) services. Findings include: R6's annual Minimum 7/10/20, indicated Fright-side weakness depression, anxiety MDS further indicate cognitively impaired toileting, was not st with staff assistance standing position and transfers. A document titled Foundated 8/14/20, indicated 8/14/20, indicate	ge 15 NT is not met as evidenced cion, interview and record ailed to ensure therapy ded to maintain range of prevent decline in mobility for reviewed for ROM/Therapy Im Data Set (MDS) dated R6 had diagnoses including s, end-stage renal disease, r, and speech difficulty. The ed R6 was severely d, needed extensive assist for eady and only able to stabilize e when moving from seated to end surface to surface Physical Therapy PT Discharge cally signed by Staff-E and cated R6 should have AFO on at of bed and transferring, the dicated staff were trained in ted 9/2/20, was reviewed, and of exercises or application of p.m., R6 was observed to be g in the wheelchair, with (AFO) splint not on R6's foot, rved laying on the floor next to	F	F 688 Increase/Prevent decreas ROM/Mobility Immediate Corrective Action: R6 therapy recommendations are assessed. R6 will have AFO app therapy recommendations. The recommendations were added to plan and point of care task list. Action as it is applied to others: Restorative nursing services poli reviewed and remains current Care Planning policy was revieweremains current Therapy recommendations for all residents were reviewed from the months to ensure that all recommendations were added to plan and point of care task list. Communication process between and clinical was reviewed and re Reeducation has been implement therapy and clinical staff regarding communication, care planning, implementing and providing there recommendations. Date of completion:10/09/2020 Recurrence will be prevented by: Audits of 3 therapy recommendations are care planting recommendations are care planting and providing there are being followed. The results of audits will be reviewed with the faction of the increase, decrease or discontinual recommendations.	e being ied per the care by was ed and e past 3 the care in therapy vised. Ited with ig apy tions will onthly x2 y ed and of these acility need to	

STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA AND PLAN OF CORRECTION IDENTIFICATION NUMBER:		A. BUILDING			COMPLETED	
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	PROVIDER OR SUPPLIER	LC		STREET ADDRESS, CITY, STATE, ZIP CODE 433 COUNTY ROAD 30 DELANO, MN 55328	1 00/	00/2020
(X4) ID PREFIX TAG	(EACH DEFICIENC)	ATEMENT OF DEFICIENCIES Y MUST BE PRECEDED BY FULL SC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECT (EACH CORRECTIVE ACTION SHOU CROSS-REFERENCED TO THE APPRO DEFICIENCY)	LD BE	(X5) COMPLETION DATE
F 688	On 9/01/20, at 1:51 out of their room by laying on the floor rightstandat 4:05 p.m. R6 was facility via transport gripper socks on he shoes nor the AFO On 9/02/20, at 8:02 -B with NA-C were incontinence cares NA-B and NA-C did or shoes. NA-B plat NA-B then transfer gait belt and pivot transfer gait belt gait gait gait gait gait gait gait gai	I p.m., R6 was observed to be ut R6's AFO was in the room next to the residents as observed returning to the t van in a wheelchair with blue er feet. R6 was not wearing a. a.m., Nursing Assistant (NA) observed to perform and dressed R6 for the day. If not offer to put on R6's AFO ced blue gripper socks on R6. red R6 to the wheelchair with a transfer. B was interviewed and stated tocks all the time and therapy blacing the AFO to R6's right A-B further stated she had not butting on the AFO. B was interviewed and stated age of motion (ROM) exercises R6 allows. NA-B further stated ties were "pretty good" and	F 68	audits. The correction will be monitored DON or/and designee	by: The	
		tions were in place to promote				

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245336 B. WING 09/0	3/2020
NAME OF PROVIDER OR SUPPLIER THE ESTATES AT DELANO LLC STREET ADDRESS, CITY, STATE, ZIP CODE 433 COUNTY ROAD 30 DELANO, MN 55328	70723
(X4) ID SUMMARY STATEMENT OF DEFICIENCIES ID PROVIDER'S PLAN OF CORRECTION PREFIX (EACH DEFICIENCY MUST BE PRECEDED BY FULL TAG REGULATORY OR LSC IDENTIFYING INFORMATION) TAG REGULATORY OR LSC IDENTIFYING INFORMATION) TAG CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)	(X5) COMPLETION DATE
On 9/2/20, at 12:47 p.m., RN-E was interviewed and verified there was no directive to apply an AFO to R6's Right lower extremity on the NA care sheet nor in R6's care plan. On 9/2/20, at 8:40 a.m., Staff-D, Certified Occupational Therapy Assistant, was interviewed and stated her expectation would be that R6 would be receiving ROM to both upper and lower extremities according to previous therapy recommendations. On 9/2/20, at 8:44 a.m., Staff-E, Physical Therapist, stated R6 is on a functional maintenance program and he would expect staff to follow therapy recommendations. On 9/2/20, at 12:47 p.m., RN-E was interviewed and stated therapy staff gives a copy of their recommendations to RN-E to add to the NA care sheet and point of care task list in the resident's medical chart. RN-E stated the therapy would also be in the resident's care plan. RN-E stated she did not receive any therapy recommendations from therapy for R6 in the staff communication book for the month of August. RN-E further verified there was no directive to apply an AFO to R6's Right lower extremity on the NA care sheet nor in R6's care plan. RN-E stated it would be her expectation to receive a copy of any therapy recommendations. RN-E verified her signature appears at the bottom of the document tittled PTi/OT/ST Functional Maintenance Program dated 8/13/20, outlining therapy recommendations.	

	OF DEFICIENCIES OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:				(X3) DATE SURVEY COMPLETED	
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F 761 SS=E	On 9/2/20, at 8:40 a Occupational Thera and stated her exposured would be receiving extremities according recommendations. On 9/2/20, at 8:44 a Therapist, stated Remaintenance prograte follow therapy retrieval to follow therapy retrieval July, 2017, receive restorative help promote optimathe policy further in objectives are indiversident-centered, resident's plan of calcabel/Store Drugs and biological CFR(s): 483.45(g) Labeling Drugs and biological labeled in accordant professional principappropriate access instructions, and the applicable. §483.45(h) Storage §483.45(h)(1) In acceptable acceptabl	a.m., Staff-D, Certified apy Assistant, was interviewed ectation would be that R6 ROM to both upper and lowering to previous therapy a.m., Staff-E, Physical 6 is on a functional am and he would expect staff commendations. storative Nursing Services, indicated Residents will nursing care as needed to hal safety and independence. Indicated restorative goals and idualized and and are outlined in the fare. and Biologicals h)(1)(2) g of Drugs and Biologicals als used in the facility must be not with currently accepted bles, and include the ory and cautionary expiration date when a compartment of the facility must store all drugs and discompartments under proper ls, and permit only authorized	F 6			10/9/20	

STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:		` ′	PLE CONSTRUCTION 3	COMF	TE SURVEY MPLETED	
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locked, permanent storage of controlled the Comprehensive Control Act of 1976 abuse, except when package drug distributed quantity stored is in the readily detected. This REQUIREME by: Based on observations (stored for anybody in the medications (stored for anybody in the medication administialled to properly resof 7 residents (R23 services. This had residents residing is survey. During observations review with RN-A con North medication of bottles of stock medication of the control of the contr	facility must provide separately ly affixed compartments for ed drugs listed in Schedule II of e Drug Abuse Prevention and 5 and other drugs subject to in the facility uses single unit libution systems in which the minimal and a missing dose can l. NT is not met as evidenced tion, interview, and document failed to ensure stock filled different medications that can used facility) were not expired at estration. In addition, the facility elable resident medication for 1 and resident medication for 1 and medication storage on 8/31/20, at 3:00 p.m. of eart, RN- verified the following edications were expired:), each expired 5/20 comp (milligrams) (acid	F 76	F761: Label/Store drugs & biologous limmediate Corrective Action: All expired medications were dis R23 medication was relabeled with change of order sticker. Action as it is applied to others: Medication Ordering and Receive Pharmacy Policy was reviewed remains current. Medication carts were reviewed change of order stickers were pall medication cards that had chorders. Reeducation was implemented licensed nursing staff regarding policy and procedure as it relates storage of medication and utilization cards that had chorders. Reeducation was implemented licensed nursing staff regarding policy and procedure as it relates to the storage of medication and utilization cards that had chorders. Recurrence will be prevented by Audits of all of the medications residents and of the stock medications and of the stock medications and of the stock medications and the stock medications are sidents and of the stock medications and the stock medications are sidents.	sposed of. with ving from and , and laced on lange of with facility es to ation of		

STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:		` ′	MULTIPLE CONSTRUCTION (X3) DATE SUR UILDING C			
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F 761	checked medication discontinue medicates tasks. RN-C further residents expired medications, if they therapeutic past ex residents could posimedications, if they buring observation review with licenses 8/31/20, at 3:35 p.m. LPN-A verified the medications were expired 3/202 Vitamin C 500 mg (expired 5/20. Aspirin 325 mg (no each expired 5/20. Aspirin 325 mg (no each expired 5/202. Oyster shell calcium supplement), each 24 hour all day alles each expired 6/20. Calcium antacid ex reducer), each expired 6/20. Calcium antacid ex reducer), each expired 6/20. Milk of magnesia (lieach expired 3/20. During interview on stated the night shir carts for expired an LPN-A further states	cN)-C stated night shift in carts for expired or itions every 1-2 weeks for their is stated staff should not give nedications, as they are not as piration date. RN-C stated all issibly use the stock is had an order for them. and medication storage d practical nurse LPN-A on in. of South medication cart, following bottles of stock expired: rgy 10 mg (antihistamine), 0. (Vitamin C supplement), each insteroidal anti-inflammatory), 0. in 500 mg (calcium expired 7/2020. rgy 10 mg (antihistamine), itra strength open (acid ired 2/16/20. rgy 10 mg (antihistamine), rgy 10 mg (antihistamine),	F 76	months to ensure the medicatic expired and to ensure proper la The results of these audits will reviewed with the facility QAPI for input on the need to increas decrease or discontinue the au The correction will be monitore or/and designee	abeling. be Committee se, dits.	

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION A. BUILDING			(X3) DATE SURVEY COMPLETED C	
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NAME OF PROVIDER OR SUPPLIER THE ESTATES AT DELANO LLC				43	REET ADDRESS, CITY, STATE, ZIP CODE 3 COUNTY ROAD 30 ELANO, MN 55328	1 001	00,2020
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F 761	During observation registered nurse (R mg One Capsule (2 per label on the me hard chart dated 9/ omeprazole to 20 n week then disconting interview on stated the omeprazyesterday and sher R23's bubble pack, she follows include information on the locorrect order. State order is the one who bubble card is correct medications medication errors for During interview on stated best practice changes would inclubible pack that identify the more consistent of the worthen wrote in pen "of During phone interview on the more pharmacist-B state not have a negative	on 9/2/20, at 8:40 a.m. (N)-A gave R23 omeprazole 20 (20) mg by mouth once a day edication bubble pack. Order in 1/20 states decrease ag every other day for one nue. 9/2/20, at 10:16 a.m. RN-A cole order came in late forgot to change the label on RN-A stated the process that is crossing off the incorrect bubble pack and entering the ed the nurse who writes the order. RN-A further stated is labels help prevent from occurring. 9/3/20, at 9:35 a.m. RN-E is for labeling medication ude placing a sticker on the entifies the medication. on 9/2/20, at 10:17 a.m. RN-A ds once daily with a single line.	F 7	61			
	Pharmacist-C state	view on 9/3/30, at 11:14 a.m. Indicate the consulting pharmacist acility in the past and has					

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION A. BUILDING			(X3) DATE SURVEY COMPLETED C	
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NAME OF PROVIDER OR SUPPLIER THE ESTATES AT DELANO LLC				STREET ADDRESS, CITY, STATE, ZIP COD 433 COUNTY ROAD 30 DELANO, MN 55328	•		
(X4) ID PREFIX TAG	(EACH DEFICIENC)	ATEMENT OF DEFICIENCIES Y MUST BE PRECEDED BY FULL SC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRE (EACH CORRECTIVE ACTION SH CROSS-REFERENCED TO THE APP DEFICIENCY)	OULD BE	(X5) COMPLETION DATE	
F 761	During documental Pharmacist-C identicart audit was on 3 expired OTC medic the physician's directions for use, 1 important label information or indicating directions for use, 1 important label informacy dated 4/labels are not alterway by nursing per transferred from or no circumstances a requested or accept the pharmacy/regis label on the medical physician's direction is inaccurate, the norder check chart indicating there is a taking care not to cinformation". Policy Labeling of Market in the container or package.	tion review dated 9/3/20, tified, "Our last Delano med /9/20. There were some cations identified at that time. If ections for use change or the the nurse may place a check chart" label on the g there is a change in taking care not to cover ormation." Ordering and Receiving from 2018, identified, "Medication ed, modified, or marked in any sonnel. Contents are not ne container to another. Under are unattached labels oted from the pharmacy. Only stered pharmacist may place a action container. 1) If the ns for use change or the label turse may place a "change of t" label on the container a change in directions for use, cover important label Medication Containers dated only the dispensing pharmacy ne label on a medication	F 76	51			
		ose in containers that are without secure closures are					

F 761 Continued From page 23 immediately removed from inventory, disposed of according to procedures for medication disposal (Section IE: Disposal of Medications and medication-related supplies), and reordered from the pharmacy. (See IC3: Ordering and receiving non-controlled medication from the dispensing pharmacy), if a current order exists. F 803 Menus Meet Resident Nds/Prep in Adv/Followed CFR(s): 483.60(c)(1)-(7) §483.60(c) Menus and nutritional adequacy. Menus must- §483.60(c)(1) Meet the nutritional needs of residents in accordance with established national guidelines.; §483.60(c)(2) Be prepared in advance; §483.60(c)(3) Be followed; §483.60(c)(4) Reflect, based on a facility's reasonable efforts, the religious, cultural and ethnic needs of the resident population, as well as input received from residents and resident groups; §483.60(c)(5) Be updated periodically; §483.60(c)(6) Be reviewed by the facility's dietitian or other clinically qualified nutrition professional for nutritional adequacy; and §483.60(c)(7) Nothing in this paragraph should be construed to limit the residents right to make	STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION A. BUILDING		COM	(X3) DATE SURVEY COMPLETED	
THE ESTATES AT DELANO LLC THE ESTATES AT DELANO LLC (X4) ID			245336	B. WING		1		
FREFIX TAG REGULATORY OR LSC IDENTIFYING INFORMATION) F 761 Continued From page 23 immediately removed from inventory, disposed of according to procedures for medication disposal (Section IE: Disposal of Medications and medication-related supplies), and reordered from the pharmacy, (See IG3: Ordering and receiving non-controlled medication from the dispensing pharmacy), if a current order exists. F 803 SS=D CFR(s): 483.60(c)(1)-(7) §483.60(c)(1) Meet the nutritional adequacy. Menus must- §483.60(c)(2) Be prepared in advance; §483.60(c)(3) Be followed; §483.60(c)(3) Be followed; §483.60(c)(4) Reflect, based on a facility's reasonable efforts, the religious, cultural and ethnic needs of the residents and resident groups; §483.60(c)(5) Be updated periodically; §483.60(c)(6) Be reviewed by the facility's dietitian or other clinically qualified nutrition professional for nutritional adequacy; and §483.60(c)(7) Nothing in this paragraph should be construed to limit the resident's right to make		NAME OF PROVIDER OR SUPPLIER			433 COUNTY ROAD 30			
immediately removed from inventory, disposed of according to procedures for medication disposal (Section IE: Disposal of Medications and medication-related supplies), and reordered from the pharmacy. (See IC3: Ordering and receiving non-controlled medication from the dispensing pharmacy), if a current order exists. F 803 Menus Meet Resident Nds/Prep in Adv/Followed CFR(s): 483.60(c)(1)-(7) §483.60(c) Menus and nutritional adequacy. Menus must- §483.60(c) Menus and nutritional needs of residents in accordance with established national guidelines.; §483.60(c)(2) Be prepared in advance; §483.60(c)(3) Be followed; §483.60(c)(4) Reflect, based on a facility's reasonable efforts, the religious, cultural and ethnic needs of the resident population, as well as input received from residents and resident groups; §483.60(c)(5) Be updated periodically; §483.60(c)(6) Be reviewed by the facility's dietitian or other clinically qualified nutrition professional for nutritional adequacy; and §483.60(c)(7) Nothing in this paragraph should be construed to limit the resident's right to make	PRÉFIX	(EACH DEFICIENCY MUST BE PRECEDED BY FULL		PREFIX (EACH CORRECTIVE ACTION SHOUTS TAG CROSS-REFERENCED TO THE APPR		JLD BE	COMPLETION	
personal dietary choices. This REQUIREMENT is not met as evidenced by:	F 803	immediately remove according to proceed (Section IE: Disposs medication-related the pharmacy. (See non-controlled med pharmacy), if a curr Menus Meet Reside CFR(s): 483.60(c)(1) Meet residents in accordaguidelines.; §483.60(c)(1) Meet residents in accordaguidelines.; §483.60(c)(2) Be professionable efforts, ethnic needs of the input received from groups; §483.60(c)(5) Be up §483.60(c)(6) Be redicitian or other clir professional for nut §483.60(c)(7) Nothic construed to limit the personal dietary characterists.	ed from inventory, disposed of dures for medication disposal al of Medications and supplies), and reordered from a IC3: Ordering and receiving ication from the dispensing rent order exists. ent Nds/Prep in Adv/Followed 1)-(7) and nutritional adequacy. the nutritional needs of ance with established national repared in advance; allowed; act, based on a facility's the religious, cultural and resident population, as well as residents and resident addated periodically; eviewed by the facility's nically qualified nutrition ritional adequacy; and and ing in this paragraph should be the resident's right to make poices.				10/9/20	

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	l ` ′	PLE CONSTRUCTION G	COM	(X3) DATE SURVEY COMPLETED C 09/03/2020	
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	PROVIDER OR SUPPLIER	LC		STREET ADDRESS, CITY, STATE, ZIP C 433 COUNTY ROAD 30 DELANO, MN 55328			
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F 803	Based on observareview, the facility for preferences for 2 or reviewed for resident Findings include: R2's quarterly Minit 6/3/20, indicated R received a regular During an interview stated he preferred with breakfast but wadded that breakfast protein over any ottand it was important because he has prostated it would be a eggs every once in When interviewed assistant (NA)-A st bacon and sausage When interviewed registered nurse (F mentioned several breakfast. RN-F fur served at breakfast. RN-F fur served at breakfast.	tion, interview, and document failed to accommodate dietary of 3 residents (R2, R20) ent choices. The mum Data Set (MDS) dated 2 was cognitively intact, diet and had a pressure sore. To on 8/31/20, at 5:01 p.m. R2 It to have bacon or sausage was not receiving regularly. R2 st meats was his preferred her meat throughout the day not that he ate enough protein essure ulcers. Lastly, R2 also nice to have scrambled a while. To 9/3/20, at 8:54 a.m. nursing ated R2 had requested more	F 80	F803: Food preferences R2 food preferences have be completed. R20 discharged facility on 9/26/2020. Food preferences complete residents. Re-education initiated to cult completing food preferences resident food preferences to capabilities. Audits of 3 residents food preferences with the resident interview completed weekly. Audits with the completed weekly in a very completed weekly and the residents preferences are being met. These audits will be reviewed facility QAPI Committee for need to increase, decrease the audits. Correction will be monitored services direct and/or designates.	from the d on current linary staff on s and follow o facilities references, vs will be will be monthly x2 food The results of d with the input on the or discontinue		

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION (X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:		(X2) MUL A. BUILD		(X3) DATE SURVEY COMPLETED C			
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	PROVIDER OR SUPPLIER TATES AT DELANO L	LC		4	TREET ADDRESS, CITY, STATE, ZIP CODE 33 COUNTY ROAD 30 ELANO, MN 55328	,	
(X4) ID PREFIX TAG	(EACH DEFICIENCY	TEMENT OF DEFICIENCIES MUST BE PRECEDED BY FULL SC IDENTIFYING INFORMATION)	ID PREFI TAG		PROVIDER'S PLAN OF CORRECTIO (EACH CORRECTIVE ACTION SHOULD CROSS-REFERENCED TO THE APPROP DEFICIENCY)) BE	(X5) COMPLETION DATE
F 803	7/31/20, did not ide R20's care plan dat food preferences. R20's Food Prefere requested but not p R20's clinical nutriti does not identify for On 9/1/20, at 8:18 at to have pancakes, eggs. R20 stated h preference for brea to facility. On 9/2/30, at 2:35 p -1 stated breakfast alternative with the pandemic restrictio place to make staff stated she had cho restructured the me for breakfast, add t sausage three time on the menu. DM-1 preference an asse however, if a prefer resident is informed meet the resident's The Estates at Dela dated 8/30/20 iden served 3 days out of The Estates at Dela	sessment (CAA) dated ntify food preferences. ded 7/24/20, did not identify ence Assessment was provided. on evaluation dated 7/27/20, a.m. R20 stated he would like bacon and sunny side up he had not been ask for his kfast proteins since admission on the dietary manager (DM) meal did not have an menu and due to the COVID has to menu items were put in ing accommodations. DM-1 sen to keep staff hours and enu which included less meat he menu now only included as a week and bacon was not stated when a resident had a resment was completed ence cannot be met, that did but no future attempts to preferences are made.	F 8	803			

STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA AND PLAN OF CORRECTION IDENTIFICATION NUMBER:		(X2) MUL [*] A. BUILDI		(X3) DATE SURVEY COMPLETED			
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F 803	Continued From pa	ge 26	F8	03			
	The Estates at Dela policy was requeste Infection Preventior CFR(s): 483.80(a)(n & Control	F 8	80			10/2/20
	infection prevention designed to provide comfortable enviror	tablish and maintain an and control program a safe, sanitary and ament and to help prevent the ansmission of communicable					
	program. The facility must es	tablish an infection prevention (IPCP) that must include, at owing elements:					
	infections and commersidents, staff, voluindividuals providing arrangement based	g, investigating, and controlling municable diseases for all unteers, visitors, and other g services under a contractual I upon the facility assessment ig to §483.70(e) and following					
	procedures for the put are not limited to (i) A system of surve possible communic infections before the persons in the facility	eillance designed to identify able diseases or ey can spread to other					

		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	` ′	IPLE CONSTRUCTION IG	CON	(X3) DATE SURVEY COMPLETED		
		245336	B. WING _			C / 03/2020		
	NAME OF PROVIDER OR SUPPLIER THE ESTATES AT DELANO LLC			STREET ADDRESS, CITY, STATE, ZIP 433 COUNTY ROAD 30 DELANO, MN 55328				
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F 880	communicable discreported; (iii) Standard and to be followed to provide to be followed to provide the followed the foll	ransmission-based precautions revent spread of infections; isolation should be used for a but not limited to: uration of the isolation, e infectious agent or organism that the isolation should be the sible for the resident under the ces under which the facility oyees with a communicable skin lesions from direct ints or their food, if direct it the disease; and ne procedures to be followed direct resident contact. Stem for recording incidents e facility's IPCP and the taken by the facility. Indle, store, process, and as to prevent the spread of review. duct an annual review of its heir program, as necessary. NT is not met as evidenced	F 88	F880 Infection Prevention Immediate Corrective Actionursing staff member was educated regarding appropand doffing procedure.	on: Identified immediately			

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION A. BUILDING			(X3) DATE SURVEY COMPLETED	
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NAME OF	PROVIDER OR SUPPLIER				TREET ADDRESS, CITY, STATE, ZIP CODE		
THE EST	ATES AT DELANO L	LC			33 COUNTY ROAD 30 ELANO, MN 55328		
(X4) ID PREFIX TAG	(EACH DEFICIENCY	ATEMENT OF DEFICIENCIES / MUST BE PRECEDED BY FULL SC IDENTIFYING INFORMATION)	ID PREFI TAG		PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD CROSS-REFERENCED TO THE APPROPI DEFICIENCY)	BE	(X5) COMPLETION DATE
F 880	practice had the poresiding at the facil to disinfect the Hoy for 1 of 2 residents control. This had the residents in the fact the Hoyer lift. Findings include: On 8/31/20, at 4:22 therapist assistant exiting a room desi isolation precaution be wearing a gown asked what the proprotective equipmeshe removes her gwashes her hands, places items in recishe is exiting. CO on doffing PPE in Manager did the transplacement of the weather that was designate precautions. AD no mask, or gloves. We required to be done a resident on isolat that she removes her mouse her mouse her mouse her mouse her moves her	tential to affect all 27 residents ity. In addition, the facility failed er lift between residents use (R11) reviewed for infection in potential to affect all illity who required the use of P.p.m., certified occupational (COTA)-A was observed gnated for a resident with its. COTA-A was noted to not play gloves, or mask. When cedure for doffing person int (PPE), COTA-A stated that own, then removes gloves and then removes her mask, and eptacle in resident room as TA-A stated she was instructed March or April and that a nurse	F	380	Infections from the previous 2 monwere re-reviewed including tracking analyzing for trends and actions ne All mechanical lifts were disinfected. Action as it is applied to others: Policy Cleaning and Disinfection of Resident-Care items and Equipmer Policy was reviewed and remains of Coronavirus policy was reviewed at remains current. Reeducation was initiated for staff regarding cleaning and disinfection resident-care items, specifically mechanical lifts. Reeducation was initiated with staff regarding donnin doffing PPE per CDC guidelines. Date of completion: 10/2/20 Recurrence will be prevented by: Audits of PPE use and lift use will be completed weekly x 4 then monthly months to ensure PPE is donned a doffed per CDC guidance and that are properly disinfected after each The results of these audits will be reviewed with the facility QAPI Confor input on the need to increase, decrease or discontinue the audits. The correction will be monitored by and/or designee	g and eded. d. nt urrent nd of g and ee x2 nd the lifts use. nmittee	

AND PLAN OF CORRECTION IDENTIFIC		L IDENTIFICATION NUMBER. L '		FIPLE CONSTRUCTION NG) COM	(X3) DATE SURVEY COMPLETED	
		245336	B. WING		C 09/03/2020		
	NAME OF PROVIDER OR SUPPLIER THE ESTATES AT DELANO LLC			STREET ADDRESS, CITY, STATE, ZIP C 433 COUNTY ROAD 30 DELANO, MN 55328			
(X4) ID PREFIX TAG	(EACH DEFICIENC)	TEMENT OF DEFICIENCIES MUST BE PRECEDED BY FULL SC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF COF (EACH CORRECTIVE ACTION CROSS-REFERENCED TO THE . DEFICIENCY)	SHOULD BE	(X5) COMPLETION DATE	
F 880	-A was observed to entering a resident precaution. Upon e observed to have d facemask, and was shieldat 10:28 a.m., NArevealed staff have control procedures instructions to doff facemask, before e On 9/2/2020, at 9:2 nursing (ADON) was resident room that required airborne is noted to be wearing shield as she exited ADON about doffin stated that before sprecautions she religarbage can, remogarbage can, and wis out of the room sin garbage, then reface shield with dishands. ADON state with staff in April 20 were instructed on PPE. ADON stated remove their mask designated an airborals also stated the direstaff on the door to rooms. ADON stated	4 a.m. Nursing Assistant (NA) don appropriate PPE before room identified with droplet xiting the room, NA-A was offed the gown, gloves and conly wearing a plastic face. A was interviewed and been trained on infection stating the training included all PPE, including the exiting an isolation room. 3 a.m., assistant director of as exiting a COVID-19 positive was designated as a room that colation precautions. ADON a face mask and a face of the room. Writer asked g PPE procedure. ADON the leaves a room that has moves her gloves, places in wes her gown, places them in washes her hands. When she he removes her mask, puts it moves her face shield, cleans infectant and then washes her ed that she did PPE education 120. ADON stated all staff the proper way to don and doff at it is her expectation staff will after leaving a room that is orne isolation room. ADON ctions are clearly defined for any of the airborne isolation es they have been completing y to ensure staff are donning	F 8	80			

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION (X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:		` ′	TIPLE CONSTRUCTION NG	COM	(X3) DATE SURVEY COMPLETED		
245336			B. WING_			C 09/03/2020	
	PROVIDER OR SUPPLIER	LC		STREET ADDRESS, CITY, STATE, ZIP COI 433 COUNTY ROAD 30 DELANO, MN 55328			
(X4) ID PREFIX TAG	(EACH DEFICIENCY	TEMENT OF DEFICIENCIES MUST BE PRECEDED BY FULL SC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORR (EACH CORRECTIVE ACTION SI CROSS-REFERENCED TO THE AF DEFICIENCY)	HOULD BE	(X5) COMPLETION DATE	
F 880	Facilities In-Service objectives for drople positive COVID ide PPE before leaving wipe goggles/face signed the in-service service training she trainer was. Facility Coronavirus states appropriate It protection) per isola and removed when the resident's room Control (CDC) Don Facility provided CI guidance to writer. staff are to remove may exit the room. hand hygiene, remote then remove and diffuse instead of resident room. No of Hoyer. During observation nursing assistant (Non R11 then pushe resident room. No of Hoyer. During interview on stated she forgot to between residents should have cleane R11's room to previruses, especially states.	et raining record dated 4/5/20, et and contact precautions for ntified that staff must remove the room and use red top to shield. 10 staff members the training record sheet. In the tet does not designate who the est (COVID-19) Policy, undated, PPE (gloves, gown, mask, eye ation guidelines will be applied entering and before exiting, per Centers for Disease ning and Doffing Guidance. DC donning and doffing Per CDC doffing guidance gloves, remove gown, and Staff are then to perform ove face shield or goggles, scard respirator (or facemask)	F 88	30			

STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA AND PLAN OF CORRECTION IDENTIFICATION NUMBER:		(X2) MULTIPLE CONSTRUCTION A. BUILDING				(X3) DATE SURVEY COMPLETED	
		245336	B. WING			C 09/03/2020	
	PROVIDER OR SUPPLIER	LC		STREET ADDRESS, CITY, ST 433 COUNTY ROAD 30 DELANO, MN 55328	FATE, ZIP CODE	, 007	30,2020
(X4) ID PREFIX TAG	(EACH DEFICIENCY	TEMENT OF DEFICIENCIES MUST BE PRECEDED BY FULL SC IDENTIFYING INFORMATION)	ID PREFIX TAG	X (EACH CORRECTI' CROSS-REFERENCE	LAN OF CORRECTION IVE ACTION SHOULD ED TO THE APPROPE FICIENCY)	BE	(X5) COMPLETION DATE
F 880	registered nurse (R equipment must be residents use to pre Policy Cleaning and items and Equipme "Reusable items are sterilized between redurable medical equipment will	9/1/20, at 8:30 a.m. N)-A stated all shared disinfected after each event spreading germs. d Disinfection of Resident-Care nt dated 10/2018, identified, e cleaned and disinfected or esident (e.g., stethoscopes, uipment)." "Reusable resident be decontaminated and/or esidents according to	F8	.80			



Protecting, Maintaining and Improving the Health of All Minnesotans

DIRECTED PLAN OF CORRECTION

A Directed Plan of Correction (DPOC) is imposed in accordance with 42 CFR § 488.424. Your facility must include the following in their POC for the deficient practice cited at F880:

PERSONAL PROTECTIVE EQUIPMENT (PPE)

- Address how corrective action will be accomplished for those residents found to have been affected by the deficient practice.
 - Address how the facility will identify other residents having the potential to be affected by the same deficient practice.

POLICIES/PROCEDURES/SYSTEM CHANGES:

• The facility's Quality Assurance and Performance Improvement Committee must conduct a root cause analysis (RCA) to identify the problem(s) that resulted in this deficiency and develop intervention or corrective action plan to prevent recurrence.

The Infection Preventionist and Director of Nursing, shall complete the following:

- Review policies and procedures for donning/doffing PPE during COVID-19 with current guidelines to include crisis standard of care, contingency standard of care and standard care.
- Develop and implement a policy and procedure for source control masks.
- Review policies regarding standard and transmission based precautions and revise as needed.

TRAINING/EDUCATION:

As a part of corrective action plan, the facility must provide training for the Infection Preventionist, the Director of Nursing, all staff providing direct care to residents, and all staff entering resident's rooms, whether it be for residents' dietary needs or cleaning and maintenance services. The training must cover standard infection control practices, including but not limited to, transmission-based precautions, appropriate PPE use, and donning and doffing of PPE.

- The training may be provided by the Director of Nursing, Infection Preventionist, or Medical Director with an attestation statement of completion.
 - The training must include competency testing of staff and this must be documented.
- Residents and their representatives should receive education on the facility's Infection Prevention Control Program as it related to them and to the degree possible/consistent with resident's capacity.
- Online infection prevention training courses may be utilized. The CDC and MDH websites have several infection control training modules and materials.

CDC RESOURCES:

Infection Control Guidance: https://www.cdc.gov/coronavirus/2019-ncov/hcp/infection-control.html

CDC: Isolation Precautions Guideline:

https://www.cdc.gov/infectioncontrol/guidelines/isolation/index.html

CDC: Guideline for Isolation Precautions: Preventing Transmission of Infectious Agents in Healthcare

Settings (2007): https://www.cdc.gov/infectioncontrol/guidelines/isolation/index.html

CDC: Personal Protective Equipment: https://www.cdc.gov/niosh/ppe/

Healthcare Infection Prevention and Control FAQs for COVID-19:

https://www.cdc.gov/coronavirus/2019-ncov/hcp/faq.html?CDC AA refVal=https%3A%2F%2Fwww.cd

c.gov%2Fcoronavirus%2F2019-ncov%2Fhcp%2Finfection-control-faq.html

MDH RESOURCES:

Personal Protective Equipment (PPE) for Infection Control:

https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/ppe/index.html

MDH Contingency Standards of Care for COVID-19: Personal Protective Equipment for Congregate Care

Settings (PDF): https://www.health.state.mn.us/communities/ep/surge/crisis/ppegrid.pdf

Interim Guidance on Facemasks as a Source Control Measure (PDF):

https://www.health.state.mn.us/diseases/coronavirus/hcp/maskssource.pdf

Interim Guidance on Alternative Facemasks (PDF):

https://www.health.state.mn.us/diseases/coronavirus/hcp/masksalt.pdf

Aerosol-Generating Procedures and Patients with Suspected or Confirmed COVID-19 (PDF):

https://www.health.state.mn.us/diseases/coronavirus/hcp/aerosol.pdf

Droplet Precautions:

https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/pre/droplet.html

Airborne Precautions:

https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/pre/droplet.html

MONITORING/AUDITING:

- The Director of Nursing, the Infection Preventionist, and other facility leadership will conduct audits of donning/doffing PPE with Transmission Based Precautions i.e. Droplet precautions.
- The Director of Nursing, Infection Preventionist, and other facility leadership will conduct routine audits on all shifts four times a week for one week, then twice weekly for one week once compliance is met. Audits should continue until 100% compliance is met on source control masking for staff, visitors and residents.
 - The Director of Nursing, Infection Preventionist, and other facility leadership will conduct real time audits on all aerosolized generating procedures to ensure PPE is in us.
 - The Director of Nursing, Infection Preventionist, or designee will review the results of audits and monitoring with the Quality Assurance Program Improvement (QAPI) program.

EQUIPMENT/ENVIRONMENT

- Address how corrective action will be accomplished for those residents found to have been affected by the deficient practice.
- Address how the facility will identify other residents having the potential to be affected by the same deficient practice.

POLICIES/PROCEDURES/SYSTEM CHANGES:

- The facility's Quality Assurance and Performance Improvement Committee must conduct a root cause analysis (RCA) to identify the problem(s) that resulted in this deficiency and develop intervention or corrective action plan to prevent recurrence.
- The director of housekeeping, director of maintenance, and director of nursing must review
 policies and procedures regarding disinfecting multiuse/shared equipment/items and/or
 environmental disinfection to ensure they meet the CDC guidance for disinfection in health
 care facilities and follow disinfectant product manufacturer directions for use including contact
 time.

TRAINING/EDUCATION:

• The Director of Housekeeping/Maintenance, and/or Director of Nursing, or Infection Preventionist must train all staff responsible for resident care equipment and environment on the facility policies/practices for proper disinfection, including following manufacturer direction for use. Each staff person must demonstrate competency at the conclusion of the training.

Training and competency testing must be documented. The Minnesota Department of Health (MDH), Center for Disease Control (CDC), and Environmental Protection Agency have education materials that may be used for training.

- CDC: Infection Control Guidelines and Guidance Library. https://www.cdc.gov/infectioncontrol/guidelines/index.html/eic_in_HCF_03.pdf
- MDH COVID-19 Toolkit. https://www.health.state.mn.us/diseases/coronavirus/hcp/ltctoolkit.pdf
- EPA: List N: Disinfectants for Use Against SARS-CoV-2 (COVID-19) https://www.epa.gov/pesticide-registration/list-n-disinfectants-use-against-sars-cov-2-covid-19

CDC RESOURCES:

Infection Control Guidance: https://www.cdc.gov/coronavirus/2019-ncov/hcp/infection-control.html CDC: Isolation Precautions Guideline:

https://www.cdc.gov/infectioncontrol/guidelines/isolation/index.html

CDC: Guideline for Isolation Precautions: Preventing Transmission of Infectious Agents in Healthcare Settings (2007): https://www.cdc.gov/infectioncontrol/guidelines/isolation/index.html

CDC: Personal Protective Equipment: https://www.cdc.gov/niosh/ppe/

Healthcare Infection Prevention and Control FAQs for COVID-19:

https://www.cdc.gov/coronavirus/2019-ncov/hcp/faq.html?CDC AA refVal=https%3A%2F%2Fwww.cdc.gov%2Fcoronavirus%2F2019-ncov%2Fhcp%2Finfection-control-faq.html

MDH RESOURCES:

Personal Protective Equipment (PPE) for Infection Control:

https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/ppe/index.html
MDH Contingency Standards of Care for COVID-19: Personal Protective Equipment for Congregate Care
Settings (PDF): https://www.health.state.mn.us/communities/ep/surge/crisis/ppegrid.pdf
Interim Guidance on Facemasks as a Source Control Measure (PDF):

https://www.health.state.mn.us/diseases/coronavirus/hcp/maskssource.pdf

Interim Guidance on Alternative Facemasks (PDF):

https://www.health.state.mn.us/diseases/coronavirus/hcp/masksalt.pdf

Aerosol-Generating Procedures and Patients with Suspected or Confirmed COVID-19 (PDF):

https://www.health.state.mn.us/diseases/coronavirus/hcp/aerosol.pdf

Droplet Precautions:

https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/pre/droplet.html Airborne Precautions:

https://www.health.state.mn.us/facilities/patientsafety/infectioncontrol/pre/droplet.html

MONITORING/AUDITING:

• The Director of Nursing, the Infection Preventionist, and/or other facility leadership will conduct audits for proper cleaning and disinfection of resident use equipment/environmental cleaning, on all shifts every day for one week, then may decrease frequency as determined by compliance.

In accordance with 42 CFR § 488.402(f), this remedy is effective 15 calendar days from the date of the enforcement letter. The DPOC may be completed on or after that date. The effective date is not a deadline for completion of the DPOC. However, a revisit will not be approved prior to receipt of documentation confirming the DPOC was completed. To demonstrate that the facility successfully completed the DPOC, the facility must provide all of the following documentation. Documentation should be uploaded as attachments through ePOC.

Imposition of this DPOC does not replace the requirement that the facility must submit a complete POC for all cited deficiencies (including F880) within 10 days after receipt of the Form CMS 2567.

Item	Checklist: Documents Required
	for Successful Completion of the Directed Plan
1	Documentation of the RCA and intervention or corrective action plan based on the results with
	signatures of the QAPI Committee members.
2	Documentation that the interventions or corrective action plan that resulted from the RCA was
	fully implemented
3	Content of the training provided to staff, including a syllabus, outline, or agenda, as well as any
	other materials used or provided to staff for the training
4	Names and positions of all staff that attended and took the trainings
5	Staff training sign-in sheets
6	Summary of staff training post-test results, to include facility actions in response to any failed
	post-tests
7	Documentation of efforts to monitor and track progress of the interventions or corrective action
	plan



In order to speed up our review, identify all submitted documents with the number in the "Item" column.

Attach all items into ePOC.

F5336029

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STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA AND PLAN OF CORRECTION IDENTIFICATION NUMBER:				LE CONSTRUCTION 01 - MAIN BUILDING 01	(X3) DATE SURVEY COMPLETED			
	245336		B. WING	B. WING			09/01/2020	
	PROVIDER OR SUPPLIER	-C		4	STREET ADDRESS, CITY, STATE, ZIP CODE 133 COUNTY ROAD 30 DELANO, MN 55328			
(X4) ID PREFIX TAG	(EACH DEFICIENCY	ATEMENT OF DEFICIENCIES Y MUST BE PRECEDED BY FULL SC IDENTIFYING INFORMATION)	ID PREF TAG		PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOUL CROSS-REFERENCED TO THE APPROPERTION OF T	D BE	(X5) COMPLETION DATE	
K 000	INITIAL COMMEN		K	000				
	ALLEGATION OF ODEPARTMENT'S A SIGNATURE AT THE PAGE OF THE CM	POC WILL SERVE AS YOUR COMPLIANCE UPON THE CCEPTANCE. YOUR HE BOTTOM OF THE FIRST S-2567 FORM WILL BE CATION OF COMPLIANCE.						
	ONSITE REVISIT (CONDUCTED TO SUBSTANTIAL CO REGULATIONS HA	OF AN ACCEPTABLE POC, AN OF YOUR FACILITY MAY BE VALIDATE THAT OF WITH THE AS BEEN ATTAINED IN ITH YOUR VERIFICATION.						
	Minnesota Departm Fire Marshal Division The Estates at Dela compliance with the in Medicare/Medica 483.70(a), Life Safe edition of National I	Survey was conducted by the nent of Public Safety, State on. At the time of this survey, ano was found not in a requirements for participation aid at 42 CFR, Subpart ety from Fire, and the 2012 Fire Protection Association 01, Life Safety Code (LSC), g Health Care.						
		E AN EPOC, A PAPER COPY CORRECTION IS NOT						
	PLEASE RETURN CORRECTION FO DEFICIENCIES (K-TAGS) TO:	THE PLAN OF R THE FIRE SAFETY						
LABORATORY	Health Care Fire In State Fire Marshal		MATURE		TITLE		(X6) DATE	

Electronically Signed 10/02/2020

Any deficiency statement ending with an asterisk (*) denotes a deficiency which the institution may be excused from correcting providing it is determined that other safeguards provide sufficient protection to the patients. (See instructions.) Except for nursing homes, the findings stated above are disclosable 90 days following the date of survey whether or not a plan of correction is provided. For nursing homes, the above findings and plans of correction are disclosable 14 days following the date these documents are made available to the facility. If deficiencies are cited, an approved plan of correction is requisite to continued program participation.

AND PLAN OF CORRECTION (X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:		· '	TIPLE CONSTRUCTION NG 01 - MAIN BUILDING 01		(X3) DATE SURVEY COMPLETED			
		245336	B. WING		09/	09/01/2020		
	PROVIDER OR SUPPLIER	.c		STREET ADDRESS, CITY, STATE, ZIP CODE 433 COUNTY ROAD 30 DELANO, MN 55328	,			
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K 000	THE PLAN OF CO DEFICIENCY MUS FOLLOWING INFO	Suite 145 -5145, or Inspections@state.mn.us RRECTION FOR EACH IT INCLUDE ALL OF THE DRMATION: what has been, or will be, done	K 0	00				
	2. The actual, or processions and addition was condetermined to be Tyne facility has a find department notifical.	oposed, completion date. In title of the person rection and monitoring to ence of the deficiency. Surveyed as one building. In ano is a 1-story building with building was constructed at 3 eroriginal building was and was determined to be of cruction. In 1988 a single story ructed to the South Wing and frype II (000) construction. In structed in 2008 and was type II (000) to the East Wing. In a larm system with smoke ridors and spaces open to the enitored for automatic fire tion.						
		time of the survey. 42 CFR. Subpart 483.70(a) is						

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STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA (X2) MULTIPLE CONSTRUCTION (X3) DATE SURVEY AND PLAN OF CORRECTION **IDENTIFICATION NUMBER:** COMPLETED A. BUILDING 01 - MAIN BUILDING 01 245336 B. WING 09/01/2020 NAME OF PROVIDER OR SUPPLIER STREET ADDRESS, CITY, STATE, ZIP CODE **433 COUNTY ROAD 30** THE ESTATES AT DELANO LLC **DELANO, MN 55328** SUMMARY STATEMENT OF DEFICIENCIES PROVIDER'S PLAN OF CORRECTION (X5) COMPLETION ID (X4) ID (EACH DEFICIENCY MUST BE PRECEDED BY FULL PRÉFIX PREFIX (EACH CORRECTIVE ACTION SHOULD BE DATE REGULATORY OR LSC IDENTIFYING INFORMATION) TAG CROSS-REFERENCED TO THE APPROPRIATE TAG **DEFICIENCY**) K 000 | Continued From page 2 K 000 NOT MET as evidenced by: K 345 Fire Alarm System - Testing and Maintenance K 345 10/9/20 CFR(s): NFPA 101 SS=F Fire Alarm System - Testing and Maintenance A fire alarm system is tested and maintained in accordance with an approved program complying with the requirements of NFPA 70, National Electric Code, and NFPA 72, National Fire Alarm and Signaling Code. Records of system acceptance, maintenance and testing are readily available. 9.6.1.3, 9.6.1.5, NFPA 70, NFPA 72 This REQUIREMENT is not met as evidenced Based on staff interview and a review of the K345: available documentation, the facility has not Facility received proper documentation to maintained the fire alarm system testing and ensure compliance with NFPA72. maintenance documentation in accordance with Facility will ensure compliance on a NFPA 72 National Fire Alarm Code 2010 edition. quarterly basis of NFPA72. This deficient practice could affect 54 of 54 Re-education was initiated with residents, as well as an undetermined number of maintenance staff specific to NFPA72 staff, and visitors to the facility. requirements. NFPA72 policy Audits of documentation specific to NFPA72 compliance will be complete Findings include: quarterly X 1 year to ensure appropriate life safety documentation. The results of On facility tour between 8 AM -12 PM on September 1st, 2020, during a review of all these audits will be reviewed with the available fire alarm maintenance and testing facility QAPI Committee for input on the documentation for the last 12 months, and an need to increase, decrease or discontinue interview with the Maintenance Director revealed the audits. that at the time of the inspection the facility's fire Monitored by: Maintenance and/or alarm test documentation did not contain a designee detailed list of all the devices that had been tested and the results of the testing completed on the devices.

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STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA (X2) MULTIPLE CONSTRUCTION (X3) DATE SURVEY AND PLAN OF CORRECTION IDENTIFICATION NUMBER: COMPLETED A. BUILDING 01 - MAIN BUILDING 01 245336 B. WING 09/01/2020 NAME OF PROVIDER OR SUPPLIER STREET ADDRESS, CITY, STATE, ZIP CODE **433 COUNTY ROAD 30** THE ESTATES AT DELANO LLC **DELANO, MN 55328** SUMMARY STATEMENT OF DEFICIENCIES PROVIDER'S PLAN OF CORRECTION (X5) COMPLETION (X4) ID (EACH DEFICIENCY MUST BE PRECEDED BY FULL PRÉFIX PREFIX (EACH CORRECTIVE ACTION SHOULD BE REGULATORY OR LSC IDENTIFYING INFORMATION) TAG CROSS-REFERENCED TO THE APPROPRIATE TAG **DEFICIENCY**) K 345 Continued From page 3 K 345 This deficient practice was confirmed by the Maintenance Director. HVAC K 521 10/9/20 K 521 SS=F | CFR(s): NFPA 101 **HVAC** Heating, ventilation, and air conditioning shall comply with 9.2 and shall be installed in accordance with the manufacturer's specifications. 18.5.2.1, 19.5.2.1, 9.2 This REQUIREMENT is not met as evidenced Based on observations and an interview, it is Waiver request for September 2020 Life Safety Code Inspection. Waiver request revealed that the facility is using the corridors as part of the air distribution system to provide submitted on September 9/3/2020 Currently, The Estates at Delano is using make-up air for the sleeping rooms' bathroom exhaust, throughout the building which is not in the corridors for both North and South accordance with NFPA 90A. This deficient wings as part of the heating, ventilation, and air conditioning air distribution system practice could allow the products of combustion to travel far from the fire origin and negatively to provide make-up air for both resident affect all residents, staff and visitors by restricting rooms and bathrooms. This waiver is their means of egress in a fire situation. being requested for the following reasons: 1. There will be no adverse effect on the Findings include: health and safety of the facility's residents. family members, and staff because the building is equipped with an approved full On the facility tour between 8:00 AM to 12:00 PM on 09/01/2020, observations revealed that the smoke detector system, along with an heating, ventilation, and air conditioning systems automated full shutdown for the ventilation for the building is using the corridor system as system and fans upon detection of smoke part of the air distribution system for make-up air or activation of the building fire alarm or for the bathrooms exhaust. sprinkler system. 2. The facility is protected by a 24-hour

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION (X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:		, ,	IPLE CONSTRUCTION NG 01 - MAIN BUILDING 01	(X3) DATE SURVEY COMPLETED		
		245336	B. WING _		09/0	1/2020
	PROVIDER OR SUPPLIER	.c		STREET ADDRESS, CITY, STATE, ZIP CODE 433 COUNTY ROAD 30 DELANO, MN 55328		
(X4) ID PREFIX TAG	(EACH DEFICIENCY	TEMENT OF DEFICIENCIES / MUST BE PRECEDED BY FULL SC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD CROSS-REFERENCED TO THE APPROPROFICIENCY)	O BE	(X5) COMPLETION DATE
K 521	Continued From pa	tion was confirmed by the	K 52	supervised automatic sprinkler systems are prominently posted at all entrances/exits. There is a design exterior smoking area on the fare the patio in the back of the building only be a few residents. The area equipped with approved metal self containers for used cigarettes. 4. Annual service and maintenanc contracts exist to service all the far fire protection system including fire sprinkler—system, and portable extinguishers. 5. The building fire alarm system monitored to provide automatic fire department notification. 6. Fire safety training is provided employees on an annual basis and orientation for all new hires. 7. Fire drills are conducted quarte each shift. 8. Compliance with this provision impose an unreasonable hardship facility due to the disruption during weeks of construction to the corrid leading to all the resident rooms. Additionally, the electrical system is building would need to be upgrade handle the power load requirement air handling system. The initial bid proposed the installation of duct would negatively affect the structurintegrity of the building. The Estates at Delano was not abuilding a more cost effective solution making the ventilation system upg meet the current codes NFPA 90 Americans.	ee and major nated and of g, used is f-closing ce cility's e alarm, e is e for all during would on the 6 dors in the ed to ork that ral le to for rades to	

PRINTED: 10/23/2020 FORM APPROVED OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA (X2) MULTIPLE CONSTRUCTION (X3) DATE SURVEY AND PLAN OF CORRECTION **IDENTIFICATION NUMBER:** COMPLETED A. BUILDING 01 - MAIN BUILDING 01 245336 B. WING 09/01/2020 NAME OF PROVIDER OR SUPPLIER STREET ADDRESS, CITY, STATE, ZIP CODE **433 COUNTY ROAD 30** THE ESTATES AT DELANO LLC **DELANO, MN 55328** SUMMARY STATEMENT OF DEFICIENCIES PROVIDER'S PLAN OF CORRECTION (X5) COMPLETION (X4) ID (EACH DEFICIENCY MUST BE PRECEDED BY FULL PRÉFIX PREFIX (EACH CORRECTIVE ACTION SHOULD BE REGULATORY OR LSC IDENTIFYING INFORMATION) TAG CROSS-REFERENCED TO THE APPROPRIATE TAG **DEFICIENCY**) K 918 | Continued From page 5 K 918 Electrical Systems - Essential Electric Syste K 918 K 918 10/9/20 SS=F CFR(s): NFPA 101 Electrical Systems - Essential Electric System Maintenance and Testing The generator or other alternate power source and associated equipment is capable of supplying service within 10 seconds. If the 10-second criterion is not met during the monthly test, a process shall be provided to annually confirm this capability for the life safety and critical branches. Maintenance and testing of the generator and transfer switches are performed in accordance with NFPA 110. Generator sets are inspected weekly, exercised under load 30 minutes 12 times a year in 20-40 day intervals, and exercised once every 36 months for 4 continuous hours. Scheduled test under load conditions include a complete simulated cold start and automatic or manual transfer of all EES loads, and are conducted by competent personnel. Maintenance and testing of stored energy power sources (Type 3 EES) are in accordance with NFPA 111. Main and feeder circuit breakers are inspected annually, and a program for periodically exercising the components is established according to manufacturer requirements. Written records of maintenance and testing are maintained and readily available. EES electrical panels and circuits are marked, readily identifiable, and separate from normal power circuits. Minimizing the possibility of damage of the emergency power source is a design consideration for new installations. 6.4.4, 6.5.4, 6.6.4 (NFPA 99), NFPA 110, NFPA 111, 700.10 (NFPA 70) This REQUIREMENT is not met as evidenced by:

PRINTED: 10/23/2020 FORM APPROVED OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES (X1) PROVIDER/SUPPLIER/CLIA (X2) MULTIPLE CONSTRUCTION (X3) DATE SURVEY AND PLAN OF CORRECTION IDENTIFICATION NUMBER: COMPLETED A. BUILDING 01 - MAIN BUILDING 01 245336 B. WING 09/01/2020 NAME OF PROVIDER OR SUPPLIER STREET ADDRESS, CITY, STATE, ZIP CODE **433 COUNTY ROAD 30** THE ESTATES AT DELANO LLC **DELANO, MN 55328** SUMMARY STATEMENT OF DEFICIENCIES PROVIDER'S PLAN OF CORRECTION (X5) COMPLETION (X4) ID (EACH DEFICIENCY MUST BE PRECEDED BY FULL PRÉFIX PREFIX (EACH CORRECTIVE ACTION SHOULD BE DATE REGULATORY OR LSC IDENTIFYING INFORMATION) TAG CROSS-REFERENCED TO THE APPROPRIATE TAG DEFICIENCY) K 918 | Continued From page 6 K 918 Based on record review and staff interview the K918 Load bank facility failed to provide test documentation in Generator load bank tested and accordance with the 2012 edition of the Life completed 9/3/2020. Safety Code (NFPA 101) section 9.1.3.1 and the Facility will ensure compliance on a 2010 edition of NFPA 110 the Standard for annual basis to ensure load bank testing Emergency and Standby Power Systems. This is completed per life safety code. deficient practice could affect the safety of all of the 54 residents if the generator failed to operate Re-education was initiate with during a power outage. maintenance staff regarding life safety code for load bank testing. Findings include: Audits of load bank will be completed annually X1 year. During the facility tour between 8:30 AM to 9:30 AM on 09/01/2020 during documentation review The results of these audits will be revealed the load bank testing as required for reviewed with the facility QAPI Committee facilities that cannot meet the required 30% has for input on the need to increase, not been conducted since 2/15/2019. decrease or discontinue the audits. Correction will be monitored by This deficient condition was confirmed by the maintenance or designee. Maintenance Director.

Name of Facility

Estates at Delano Waiver for K521 Survey Date 9/3/2020

2012 LIFE SAFETY CODE

PART III - RECOMMENDATION FOR WAIVER OF SPECIFIC LIFE SAFETY CODE PROVISIONS

For each item of the Life Safety Code recommended for waiver, list the survey report form item number and state the reason for the conclusion that:
(a) the specific provisions of the code, if rigidly applied, would result in unreasonable hardship on the facility, and (b) the waiver of such unmet provisions will not adversely affect the health and safety of the patients. If additional space is required, attach additional sheet(s).

PROVISION NUMBER(S)

JUSTIFICATION

K400

K521

Waiver request for September 2020 Life Safety Code Inspection. Waiver request submitted on September 9/3/2020 Currently, The Estates at Delano is using the corridors for both North and South wings as part of the heating, ventilation, and air conditioning air distribution system to provide make-up air for both resident rooms and bathrooms. This waiver is being requested for the following reasons:

1. There will be no adverse effect on the health and safety of the facility's residents, family members, and staff because the building is

equipped with an approved full smoke detector system, along with an automated full shutdown for the ventilation system and fans upon detection of smoke or activation of the building fire alarm or sprinkler system.

2. The facility is protected by a 24-hour supervised automatic sprinkler system.

3. The internal facility is smoke-free and signs are prominently posted at all major entrances/exits. There is a designated exterior smoking area on the far end of the patio in the back of the building, used only be a few residents. The area is equipped with approved metal self-closing containers for used cigarettes.

4. Annual service and maintenance contracts exist to service all the facility's fire protection system including fire alarm, sprinkler

system, and portable extinguishers.

5. The building fire alarm system is monitored to provide automatic fire department notification.

6. Fire safety training is provided for all employees on an annual basis and during orientation for all new hires.

7. Fire drills are conducted quarterly on each shift.

8. Compliance with this provision would impose an unreasonable hardship on the facility due to the disruption during 6 weeks of construction to the corridors leading to all the resident rooms. Additionally, the electrical system in the building would need to be upgraded to handle the power load requirements of the air handling system. The initial bid also proposed the installation of duct work that would negatively affect the structural integrity of the building.

The Estates at Delano was not able to find a more cost effective solution for making the ventilation system upgrades to meet the current

codes NFPA 90 A.

Surveyor (Signature)	Title	Office	Date
Fire Authority Official (Signature)	Title FINE SAFETY SUPERVISOR	Office MN STATE FIRE MARSHAL	Date 10-22-20

Sent to Tom Linkoff 10/13/20

Ben Scherer Plumbing & Heating Inc.

4520 85th Street SE Delano, MN 55328 63 972 8137 schererplumbing@gmail.com

Name / Address	
The Estates at Delano	
433 County Road 30 SE	
Delano, MN 55328	

Estimate

Date	Estimate #	
9/10/2020	2705	Mary Control of the Control

				Project
		T T	4	
Description	Qty	Cost		Total
Price for changing the heating system so that it is not using the corridor system as part of the air distribution system and adding multiple-ducted make up air systems This system will include adding 3 new Renewaire CA2XRT Roof top heat and humidity air exchangers. This also includes running all of the existing roof top heating and cooling units to be piped in with sheet-metal instead of using the central return air system that they currently have. This price includes all sheet-metal work, new equipment, and controls		495,000.00		495,000.00
		Total		\$495,000.00
			_	w.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,

Customer Signature

Name of Facility

Estates at Delano Waiver for K521 Survey Date 9/3/2020

2012 LIFE SAFETY CODE

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Name / Address	
The Estates at Delano	
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Estimate

Date	Estimate #	
9/10/2020	2705	Mary Control of the Control

				Project
		T T	4	
Description	Qty	Cost		Total
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		Total		\$495,000.00
			_	w.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,

Customer Signature

TATEMENT OF ISOLATED DEFICIENCIES WHICH CAUSE TO HARM WITH ONLY A POTENTIAL FOR MINIMAL HARM OR SNFs AND NFs	PROVIDER # 245336	MULTIPLE CONSTRUCTION A. BUILDING: B. WING	DATE SURVEY COMPLETE: 9/3/2020		
IAME OF PROVIDER OR SUPPLIER THE ESTATES AT DELANO LLC	STREET ADDRESS, CITY, STATE, ZIP CODE 433 COUNTY ROAD 30 DELANO, MN				
D REFIX AG SUMMARY STATEMENT OF DEFICIEN	NCIES				
Notice of Bed Hold Policy Before/Upon CFR(s): 483.15(d)(1)(2) §483.15(d) Notice of bed-hold policy a §483.15(d)(1) Notice before transfer. B goes on therapeutic leave, the nursing for representative that specifies— (i) The duration of the state bed-hold poresume residence in the nursing facility (ii) The reserve bed payment policy in t (iii) The nursing facility's policies regar (1) of this section, permitting a resident (iv) The information specified in paragrams §483.15(d)(2) Bed-hold notice upon tratherapeutic leave, a nursing facility must which specifies the duration of the bed-This REQUIREMENT is not met as even Based on interview and document reviet of hospital transfer for 1 of 1 residents (Findings include: R8's Significant Change Minimum Data R8's progress note(s) identified: - 7/3/20, at 6:47 p.m. R8 was admitted the lungs). - 8/4/20, at 2:09 a.m. R8 was transferred verbal consent for bed hold. R8's medical record lacked evidence a variansfer. During interview on 9/3/20, at 10:51 a.m. verbal acceptance or denial to a bed hold (BOM) followed up with the bed hold a	nd return- defore a nursing face acility must provide acility must provide oblicy, if any, during the state plan, under the return; and raph (e)(1) of this state provide to the rehold policy described policy described policy described with a Set (MDS) dated a Set (MDS) dated to the hospital for a date to the hospital for a date of the hospital	de written information to the resident of g which the resident is permitted to return a which the resident is permitted to return a which must be consistent with particular, which must be consistent with particular and the resident for hospitalizar sident and the resident representative wheel in paragraph (d)(1) of this section. The provides the written bed hold police weed for hospitalization. 7/17/20, identified R8 had intact cognitions and pleural effusion (fluid builder high blood pressure, headache and fearement/policy was provided at the time of the provided the nurse should trying to the hospital. The business office in	resident urn and ragraph (e) tion or vritten notice ry at the time ition. d up in ver, R8 gave ne of hospital r and get a nanager		

Any deficiency statement ending with an asterisk (*) denotes a deficiency which the institution may be excused from correcting providing it is determined that other safeguards provide sufficient protection to the patients. (See instructions.) Except for nursing homes, the findings stated above are disclosable 90 days following the date of survey whether or not a plan of correction is provided. For nursing homes, the above findings and plans of correction are disclosable 14 days following the date these documents are made available to the facility. If deficiencies are cited, an approved plan of

Event ID: MTA311

The above isolated deficiencies pose no actual harm to the residents

031099

	F ISOLATED DEFICIENCIES WHICH CAUSE	PROVIDER #	MULTIPLE CONSTRUCTION	DATE SURVEY		
NO HARM WIT: FOR SNFs AND	H ONLY A POTENTIAL FOR MINIMAL HARM NFs		A. BUILDING:	COMPLETE:		
		245336	B. WING	9/3/2020		
	VIDER OR SUPPLIER SES AT DELANO LLC	STREET ADDRESS, CITY, STATE, ZIP CODE 433 COUNTY ROAD 30 DELANO, MN				
ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIE	NCIES				
F 625	Continued From Page 1 When interviewed on 9/3/20, at 12:42 president and/or the family wanted to hot to hold the bed she sent a written copy of BOM identified there was no notes indi 7/3/20 or 8/4/20. BOM stated a signed wanted to hold the bed for readmission The facility bed hold policy was reques	ld the bed while hos of the bed hold agre icating a signed bed bed hold was obtain and obtain consent	spitalized. If the resident/responsible party freement/policy to the responsible party freement/policy to the responsible party freed was obtained for R8's hospital added to identify that the resident/responsito charge to hold the bed.	arty wanted for signature. Imissions on		

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SURVEY TEAM COMPOSITION AND WORKLOAD REPORT

Public reporting burden for this collection of information is estimated to average 10 minutes per response, including time for reviewing instructions, searching existing data sources, gathering and maintaining data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing the burden, to Office of Financial Management, HCFA, P.O. Box 26684, Baltimore, MD 21207; or to the Office of Management and Budget, Paperwork Reduction Project(0838-0583), Washington, D.C. 20503.

245336	Number		vider/Supplie E ESTATES AT I					
pe of Survey (sele		pply):		Investigation vestigation	F Inspec G Valida	tion of Car	re J Sand	certification ction/Hearing te License w
A A	lect all that	app1y):	A Routine/St B Extended S C Partial Ex D Other Surv	urvey (HHA o	r long term		ity)	
and onton the wor	land informa		SURVEY TEAM A			armatian nu	mbox	
lease enter the wor Surveyor Id Number (A)	First Date Arrived (B)	Last Date Departed (C)	Pre-Survey Preparation Hours (D)	On-Site Hours 12am-8am (E)	veyor's info On-Site Hours 8am-6pm (F)	On-Site Hours 6pm-12am (G)		Off-Site Report Preparation Hours (I)
Team Leader 1. 40938	09-01-2020	09-03-2020	3.00	1.00	18.00	0.00	2.00	10.00
2. 42581	08-31-2020	09-09-2020	0.00	1.00	34.00	2.00	2.00	6.50
42583	08-31-2020	09-03-2020	0.00	1.00	24.50	2.00	4.00	9.00
43081	08-31-2020	09-03-2020	0.00	1.00	27.00	2.00	2.00	7.00
43082	08-31-2020	09-03-2020	0.00	0.00	16.00	16.00	4.50	10.00
5.								
7.								
3.								
9.								

SURVEY TEAM COMPOSITION AND WORKLOAD REPORT

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	J Sand K Stat L Chov	ction/Hearing te License
B Extended Survey (HHA or long term care facility C Partial Extended Survey (HHA) D Other Survey SURVEY TEAM AND WORKLOAD DATA Please enter the workload information for each surveyor. Use the surveyor's information numb Surveyor Id Number Date Arrived (B) (A) Date Arrived (B) (B) Departed (C) Departed (C) Departed (C) Departed (D) Departed (D) Departed (D) Departed (E) Departed (D) Departed (E) Departed	ty)	
Please enter the workload information for each surveyor. Use the surveyor's information number Surveyor Id Number Car		
Surveyor Id Number		
Surveyor Id Number (A) Date Arrived (B) Departed (C) Departed (D) Departed (D) Departed (D) Departed (D) Departed (E) Depa		+
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otal Supervisory Review Hours		0.00
Cotal Clerical/Data Entry Hours		

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Paperwork Reduction	Project(0838-	0583), Washir	ngton, D.C. 2	0503.				
Provider/Supplier	Number	Pro	vider/Supplie	er Name				
245336	245336 THE ESTATES AT DELANO LLC							
Type of Survey (sele	ct all that a	pply):	B Dumping In C Federal Mc	vestigation nitoring	F Inspec G Valida	tion of Car tion	ce J Sand K Stat	certification ction/Hearing ce License
Extent of Survey (Se	lect all that	apply):	D Follow-up A Routine/St	andard (all	providers/s		L Chov	W
			B Extended S C Partial Ex D Other Surv	tended Surve	_	care racri	y)	
			SURVEY TEAM A					
Please enter the wor Surveyor Id Number (A)	First Date Arrived (B)	Last Date Departed (C)	Pre-Survey Preparation Hours (D)	Use the sur On-Site Hours 12am-8am (E)	veyor's info On-Site Hours 8am-6pm (F)	On-Site Hours 6pm-12am (G)		Off-Site Report Preparation Hours (I)
Team Leader 1. 40938	08-31-2020	09-01-2020	0.00	0.00	3.00	0.00	0.00	0.00
2.								
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10.								
								'
Total Supervisory Re	view Hours							0.25
Total Clerical/Data	Entry Hours							2

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Arrived (B) Departed (C) Hours (E) 8am-6pm (G) 6pm-12am (H) Hours (I) Team Leader 1. 40938 08-31-2020 09-01-2020 0.00 0.00 1.00 2.00 0.00 0.00 2. 3. 4. 5. 6. 6. 7. 6. 7. 6. 7. 6. 7. 6. 7. 6. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7. 7.	Paperwork Reduction		1						
ype of Survey (select all that apply): A Complaint Investigation E Dumping Investigation C Pederal Monitoring D Follow-up Visit B Life safety Code L Chow E Survey (Select all that apply): A Routine/Standard (all providers/suppliers) B Extended Survey (IHHA) D Other Survey SURVEY TEAM AND NORKLOAD DATA Surveyor Id Number Arrived (A) Pirst Date Pate Arrived (A) (C) (C) (C) Partial Extended Survey (D) (E) (C) (C) (C) (C) (C) (C) (C) (C) (C) (C		Number							
C Federal Monitoring G Validation K State License D Follow-up Visit H Life safety Code L Chow Extent of Survey (Select all that apply): A Routine/Standard (all providers/suppliers) B Extended Survey (HHA or long term care facility) C Partial Extended Survey (HHA) D Other Survey SURVEY TEAM AND WORKLOAD DATA Clease enter the workload information for each surveyor. Use the surveyor's information number. Surveyor Id Number (A) Pirst Last Date Arrived (B) C(C) Charles (Bours Hours Sum-Spm (G)) Team Leader 1, 40938 08-31-2020 09-01-2020 0.00 0.00 1.00 2.00 0.00 0.00 0.00 2. 3. 4. 5. 6. 6. 7. 9. 9. 9. 10. 10. 10. 10. 10. 10. 10. 10. 10. 10		ect all that a				n E Initia	l Certifica	ation I Red	certification
A Routine/Standard (all providers/suppliers) B Rxtended Survey (HHA) or long term care facility) C Partial Extended Survey (HHA) D Other Survey SURVEY TEAM AND WORKLOAD DATA Clease enter the workload information for each surveyor. Use the surveyor's information number.	A			C Federal Monitoring G Validation K			K Stat	te License	
B Extended Survey (HHA or long term care facility) C Partial Extended Survey (HHA) D Other Survey	Extent of Survey (Se	lect all that	apply):						
Surveyor Id Number First Date Date Date Arrived (B) On-1-2020 On-0 O	А			B Extended Survey (HHA or long term care facility) C Partial Extended Survey (HHA)					
Surveyor Id Number (A) First Date Arrived (B) Compared (C) Date Arrived (B) Date Arrived (B) Date Arrived (C) Date	Please enter the wor	kload informa					ormation nu	mber.	
1. 40938	Surveyor Id Number	First Date Arrived	Last Date Departed	Pre-Survey Preparation Hours	On-Site Hours 12am-8am	On-Site Hours 8am-6pm	On-Site Hours 6pm-12am	Travel (Preparation Hours
3. 4. 5. 6. 7. 8. 9. 9. 9. 9. 9. 9. 9. 9. 9. 9. 9. 9. 9.		08-31-2020	09-01-2020	0.00	0.00	1.00	2.00	0.00	0.00
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7. 8. 9. 10. otal Supervisory Review Hours 0.25	5.								
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9. 10. otal Supervisory Review Hours 0.25	7.								
otal Supervisory Review Hours	8.								
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otal Clerical/Data Entry Hours 2	Total Supervisory Re	view Hours							0.25
	Fotal Clerical/Data	Entry Hours							2

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Paperwork Reduction	Project(0838-	0583), Washir	ngton, D.C. 2	0503.				
Provider/Supplier	Number	Pro	vider/Supplie	er Name				
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Type of Survey (sele	ct all that a	pply):	B Dumping In C Federal Mc	vestigation nitoring	F Inspec G Valida	tion of Car tion	e J Sand K Stat	certification ction/Hearing ce License
Extent of Survey (Se	lect all that	apply):	D Follow-up	Visit	H Life s	afety Code	L Chov	∛
A			A Routine/St B Extended S C Partial Ex D Other Surv	urvey (HHA o	r long term		ity)	
		S	SURVEY TEAM A	ND WORKLOAD	DATA			
Please enter the wor	kload informa	tion for each	n surveyor.	Use the sur	veyor's info	ormation nu	mber.	1
Surveyor Id Number (A)	First Date Arrived (B)	Last Date Departed (C)	Pre-Survey Preparation Hours (D)	On-Site Hours 12am-8am (E)	On-Site Hours 8am-6pm (F)	On-Site Hours 6pm-12am (G)	Travel (H)	Off-Site Report Preparation Hours (I)
Team Leader 1. 40938	08-31-2020	09-01-2020	0.00	0.00	2.50	0.00	0.00	0.00
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10. Total Supervisory Rev Total Clerical/Data 1								0.25

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Paperwork Reduction	Project(0838-	0583), Washir	ngton, D.C. 2	0503.				
Provider/Supplier	Number	Pro	vider/Supplie	er Name				
245336	245336 THE ESTATES AT DELANO LLC							
Type of Survey (sele	ct all that a	pply):	B Dumping In C Federal Mc	vestigation nitoring	F Inspec G Valida	tion of Car tion	e J Sand K Stat	certification ction/Hearing ce License
Extent of Survey (Se	lect all that	apply):	D Follow-up	Visit	H Life s	afety Code	L Chow	∛
A			A Routine/St B Extended S C Partial Ex D Other Surv	urvey (HHA o	r long term		ity)	
		S	SURVEY TEAM A	ND WORKLOAD	DATA			
Please enter the wor	kload informa	tion for each	n surveyor.	Use the sur	veyor's info	ormation nu	mber.	1
Surveyor Id Number (A)	First Date Arrived (B)	Last Date Departed (C)	Pre-Survey Preparation Hours (D)	On-Site Hours 12am-8am (E)	On-Site Hours 8am-6pm (F)	On-Site Hours 6pm-12am (G)	Travel (H)	Off-Site Report Preparation Hours (I)
Team Leader 1. 40938	08-31-2020	09-01-2020	0.00	0.00	2.50	0.00	0.00	0.00
2.								
3.								
4.								
5.								
6.								
7.								
8.								
9.								
10.								
10. Total Supervisory Rev Total Clerical/Data 1								0.25

U

SURVEY TEAM COMPOSITION AND WORKLOAD REPORT

Public reporting burden for this collection of information is estimated to average 10 minutes per response, including time for reviewing instructions, searching existing data sources, gathering and maintaining data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing the burden, to Office of Financial Management, HCFA, P.O. Box 26684, Baltimore, MD 21207; or to the Office of Management and Budget, Paperwork Reduction Project(0838-0583), Washington, D.C. 20503.

Paperwork Reduction			ingcon, b.c. z						
Provider/Supplier 245336	Number		ovider/Supplie E ESTATES AT I						
			E ESTATES AT 1	DELIANO LILO					
Type of Survey (sele	ct all that a	appiy):	C Federal Monitoring G Validation K S				e J Sand	anction/Hearing tate License	
Extent of Survey (Se	lect all that	apply):							
A			A Routine/Standard (all providers/suppliers) B Extended Survey (HHA or long term care facility) C Partial Extended Survey (HHA) D Other Survey						
			SURVEY TEAM A						
Please enter the wor Surveyor Id Number (A)	First Date Arrived (B)	Last Date Departed (C)	Pre-Survey Preparation Hours (D)	Use the sur On-Site Hours 12am-8am (E)	veyor's info On-Site Hours 8am-6pm (F)	On-Site Hours 6pm-12am (G)		Off-Site Report Preparation Hours (I)	
1. 43081			0.00	0.00	0.00	0.00	0.00	0.50	
2.									
3.									
4.									
5.									
6.									
7.									
8.									
9.									
10.									
		1	,					!	
Total Supervisory Re	view Hours							0.25	
Total Clerical/Data									
.ocal Cierical/Data	Enery Hours							2	

SURVEY TEAM COMPOSITION AND WORKLOAD REPORT

Public reporting burden for this collection of information is estimated to average 10 minutes per response, including time for reviewing instructions, searching existing data sources, gathering and maintaining data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing the burden, to Office of Financial Management, HCFA, P.O. Box 26684, Baltimore, MD 21207; or to the Office of Management and Budget, Paperwork Reduction Project(0838-0583), Washington, D.C. 20503.

	_		ngton, D.C. 2					
Provider/Supplier	Number	Pro	vider/Supplie	er Name				
245336		THE	THE ESTATES AT DELANO LLC					
Type of Survey (sele	ct all that a	.pply):	A Complaint B Dumping In C Federal Mo D Follow-up	vestigation onitoring	F Inspec G Valida	tion of Car	e J Sand	certification ction/Hearing ce License
Extent of Survey (Se	lect all that	apply):						
A			B Extended S	andard (all garvey (HHA o	r long term		ity)	
			SURVEY TEAM A	ND WORKLOAD I	DATA			
Please enter the wor	kload informa	tion for eac	h surveyor.	Use the sur	veyor's info	ormation nu	mber.	1
Surveyor Id Number	First Date Arrived (B)	Last Date Departed (C)	Pre-Survey Preparation Hours (D)	On-Site Hours 12am-8am (E)	On-Site Hours 8am-6pm (F)	On-Site Hours 6pm-12am (G)	Travel (H)	Off-Site Report Preparation Hours (I)
Team Leader 1. 34764	09-01-2020	09-02-2020	1.00	0.00	7.00	0.00	3.00	0.00
2.								
3.								
4.								
5.								
6.								
7.								
8.								
9.								
10.								6.50

CMS-671 Page 1 of 3





 $\label{thm:protecting} \mbox{Minnesota Department of Health: Protecting, maintaining improving the health of all Minnesotans.}$



Confirmation page! Thank you for using the data entry system. If you have comments please send to:

monica.larson@state.mn.us

Please print this page and give it to your state survey team. A page for both the CMS-671 and CMS-672 will be required to complete the process.	Print this Page
Would you like to go to the CMS-672 form for data entry?	Go to CMS-672
I'm finished and would like to exit the application.	Exit

Standard Survey Date Format: mm/dd/yy From F1: 08/31/20 To F2: 09/03/20							
Name of Facility: THE ESTATES AT DELANO LLC	Provider Number: 245336	Fiscal Year ending:					
Address: 433 COUNTY ROAD 30, DELANO, WRIG	GHT, MN 55328						
Telephone Number: F6 7639722987 State/County Code: MN / WRIGHT State/Region Code: MN / 05							
A. F9 03 - SNF/NF - Medicare/Medicaid B. Is this facility hospital based? F10 No If yes, indicate Hopsital Provider Number: F11							
Ownership: F12 03 - For Profit - Corporation	on						
Owned or leased by Multi-Facility Organization: F13 Yes Name of Multi-Facility Organization: F14 Monarch Healthcare Management							
Dedicated Special Care Units (show number of beds for all that apply)							
AIDS F15 0 Al	AIDS F15 0 Alzheimer's Disease F16 0						
Dialysis F17 0 Di	sabled Child Young Adu	lt F18 0					
Head Trama F19 0 Hospice F20 0							

CMS-671 Page 2 of 3

Huntington's Disease F21 0 Ver Other Spec Rehab. F23 4	ntilator/Respiratory Care	e F22 0					
Does the facility currently have an organized r	resident group? F24	Yes					
Does the facility currently have an organized gmembers of residents? F25	group of family	Yes					
Does the facility conduct experimental research	ch? F26	No					
Is the facility part of a continuing care retirem (CCRC)? F27	ent community	No					
the date(s) of the last approval. Indicate the nu granted. If the facility does not have a waiver, Waiver of seven day RN requirement. Waiver of 24 hr licensed nursing requirement	write NA in the blanks. Date: mm/dd/yy F28	* *					
Does the facility currently have an approved n competency program? F32	urse aide training and	No					
The following three questions are to be con	The following three questions are to be completed by the survey team.						
1) Was this a staggered Survey?	No - Not S	00					
2) If staggered, day of the week starting?	•	to Complete					
3) If staggered, starting time?	Surveyor t	to complete AM					

Name of Person Completing Form:	Date:
DEREK RIVARD	09/16/20

• Share This

Spotlight

Minnesota eLicensing

Questions?

Please contact our Health Regulation Division: <u>health.fpc-web@state.mn.us</u> or 651-201-4101.

See also > <u>Health Regulation</u>

CMS-672 Page 1 of 4





Minnesota Department of Health: Protecting, maintaining improving the health of all Minnesotans.



Confirmation page! Thank you for using the data entry system. If you have comments please send to:

monica.larson@health.state.mn.us

Please print this page and give it to your state survey team. A page for both the CMS-671 and CMS-672 will be required to complete the process.	Print this Page
Would you like to go to the CMS-671 form for data entry?	Go to CMS-671
I'm finished and would like to exit the application.	<u>Exit</u>

THE ESTATES AT DELANO LLC					
Provider No. 245336	Medicare F75	Medicaid F76	Other F77	Total Residents F78 27	

ADL	Independent	Assist of One Two Staff	Dependent
Bathing	F79 1	F80 26	F81 0
Dressing	F82 4	F83 23	F84 0
Transferring	F85 3	F86 19	F87 5
Toilet Use	F88 4	F89 22	F90 1
Eating	F91 23	F92 4	F93 0

	-	1/10		~
Δ	Rowe	4/K	ladder	Status

F94 2 With indwelling or external catheter.

F95 Of total number of residents with catheters, 2 were present on admission.

B. Mobility

F100 0 Bedfast all or most of time..

F101 17 In chair all or most of time.

F102 1 Independently ambulatory.

CMS-672 Page 2 of 4

F96 9 Occasionally or frequently incontinent of bladder.

F97 8 Occasionally or frequently incontinent of bowel.

F98 **0** On individually written bladder training program.

F99 **0** On individually written bowel training program.

F103 9 Ambulation with assistance or assistive device.

F104 0 Physically restrained.

F105 Of total number of residents with restrained, **0** were admitted with orders for restraints.

F106 4 With contractures.

F107 Of total number of residents with contractures, 4 had contractures on admission.

C. Mental Status

F108 0 With mental retardation.

F109 **18** With documentation signs and symptoms of depression.

F110 13 With documentation psychiatric diagnosis (excluding dementias and depression).

F111 10 Dementia: multi-infarct, senile, Alzheimer's type, or other than Alzheimer's type.

F112 5 With behavioral symptoms.

F113 5 Of the total number of residents with behavioral symptoms, the total number receiving a behavior management prpgram.

F114 0 Receiving health rehabilitative services for MI/MR.

D. Skin Integrity

F115 1 With pressure sores (exclude stage I).

F116 1 Of the total number of residents with pressure sores excluding stage I, how many residents had pressure sores on admission?

F117 **22** Receiving preventive skin care.

F118 0 With rashes.

E. Special Care

F119 2 Receiving hospice care benefit.

F120 0 Receiving radiation therapy.

F121 **0** Receiving chemotherapy.

F127 0 Receiving suction.

F128 6 Receiving injections (exclude vitamin B12 injections)

F129 0 Receiving tube feedings.

CMS-672 Page 3 of 4

F122 4 Receiving dialysis.

F123 0 Receiving intravenous therapy, parenteral nutrition, and/or blood transfusion.

F124 0 Receiving respiratory treatment.

F125 0 Receiving tracheostomy care.

F126 2 Receiving ostomy care.

F. Medication	G. Other
F133 17 Receiving any psychoactive medication.	F140 3 With unplanned significant weight loss/gain.
F134 5 Receiving antipsychotic medications.	F141 0 Who do not communicate in the dominant language of the facility (includes those who use sign language).
F135 6 Receiving antianxiety medications.	F142 1 Who use non-oral communication devices.
F136 14 Receiving antidepressant medications.	F143 27 With advance directives.
F137 0 Receiving hypnotic medication.	F144 20 Received influenza immunization.
F138 4 Receiving antibiotics.	F145 24 Received pneumococcal vaccine.
F139 19 On pain management program.	

I certify that this Information is accurate to the best of my knowledge.					
Name of Person Completing Title Date					
Suzie Koosman DON 09/16/2020					

To be completed by MDH survey team.
F146 Was ombudsman office notified prior to survey? Yes
F147 Was ombudsman present during any portion of the survey? No
F148 Medication error rate 4%

• Share This

MINNESOTA DEPARTMENT OF HEALTH Health Regulation Division 85 East Seventh Place, Suite 300, P.O. Box 64900 St. Paul, Minnesota 55164-0900

Email for Administ	rator: Lhogendorn @monachmn.com
Administrator:	rator: Lhogendorn @monachmn.com
National Provider I One facility may have type for this survey, i	dentifier (NPI) Number: 1720039001 e multiple NPI Numbers. Please verify the NPI number associated with the provider i.e. for a nursing home survey, the NPI Number will be associated with the Nursing
OWNERSHIP INFOR	RMATION AT THE TIME OF SURVEY
Name of Facility: <u>T</u>	HE ESTATES AT DELANO LLC City: DELANO
Name of Legal Entit	ty Operating Provider: THE ESTATES AT DELANO LLC
Name and Address	of Governing Board President:
_	JOSH LEGUM
Address: _	638 SOUTHBEND AVENUE
- - City/State/Zip:	MANKATO, MN 56001
If legal entity or pre provide the inform	esident of the governing board is different than what is noted above, please ation below.
Name of Facility:	City:
Name of Legal En	tity Operating Provider:
Name and Addre	ss of Governing Board President:
Name:	· · · · · · · · · · · · · · · · · · ·
Address:	
City/State/2	Zip:
SIGNATURE	2
Completed by:	Tym Eligenden
Title: <u> </u>	NIA
Date: $\frac{2}{2}$	3(31/2030

FIRE SAFETY SURVEY REPORT CRUCIAL DATA EXTRACT (TO BE USED WITH CMS-2786 FORMS)

PROVIDER NUMBER		Y NAME ESTATES AT DELANO LLC		SURVEY DATE *K4 09/01/2020	
K6 DATE OF PL APPROVAL		K3: MULTIPLE CONSTRUCTION TOTAL NUMBER OF BUILDINGS NUMBER OF THIS BUILDING	1A	A BUILDING B WING C FLOOR D APARTMENT UNIT	
13 2	Health Care 786 R 786 R ASC For	2012 EXISTING 2012 NEW	COMPLETE IF ICF/MR IS SURVEYED U SMALL (16 BEDS 0 1 PROMP 2 SLOW 3 IMPRAO LARGE	OR LESS) T CTICAL	
16 2	786 U ICF/MR F 786 V, W, X	2012 EXISTING	K8: 4 PROMP 5 SLOW 6 IMPRAC		
*K7 12 SELECT NUMBER OF FORM USED FROM ABOVE (Check if K321 or K351 are marked as not applicable in the		APARTMENT HOUSE 7 PROMPT 8 SLOW 9 IMPRACTICAL			
,	U, V, W, X, Y and Z.,		ENTER E-SCORE HERE K5: e.g 2.5		
*K9 : FACILITY MATERIAL *K9 :	TH (AC	ON: (Check all that apply) A2 X A3 CEPTABLE POC) (WA	AIVERS) (FSES)	A5 [PERFORMANCE BASED DESIGN)	
FACILITY DOES I	NOT MEET LSC:	K180: A. X FULLY SPRINKLI (All required areas are s			
*MANDATORY					

2012 LIFE SAFETY CODE

Form Approved OMB Exempt

				1	
	ORT - 2012 LIFE SAFETY COD _THCARE	1. (A) PI	ROVIDER NUMBER	1. (B) MEDICAID I.D. NO.	
OPTIONAL — CI		Facilities Code, Ne commendation for V Crucial Data Extra	w and Existing Waiver ct	ncies – CMS-2786T	
Identifying information as shown in applic	able records. Enter changes, if any, alor	ngside each item, (giving date of change.		
2. NAME OF FACILITY	2. (A) MULTIPLE CONSTRUCTION (BLDGS) A. BUILDING B. WING C. FLOOR	2. (B) ADDRESS OF I	FACILITY (STREET, CITY,	STATE, ZIP CODE) A. Fully Sprinklered (All required areas are sprinklered) B. Partially Sprinklered (Not all required areas are sprinklered) C. None (No sprinkler system) K0180	
3. SURVEY FOR	4. DATE OF SURVEY	DATE OF PLAN APP	ROVAL SURVEY I	<u> </u>	
☐ MEDICARE ☐ MEDICAID	К4	5 K7		2012 EXISTING 6. 2012 NEW	
5. SURVEY FOR CERTIFICATION OF	· · ·				
1. HOSPITAL 2. SKILLED/NU	RSING FACILITY 4. ICF/IID UN	DER HEALTH CARE	5. HOSPICE	Ξ	
IF "2" OR "5" ABOVE IS MARKED, CHECK APPRO	· <i>,</i>		3. IF DISTINCT PART	OF HOSPITAL, IS HOSPITAL ACCREDITED?	
	HOSPITAL BEDS c. NUMBER OF SKILLED CERTIFIED FOR MED	-	JMBER OF SKILLED BED ERTIFIED FOR MEDICAID		
	IONS 2. ACCEPTANCE OF A PLAN OF CO		COMMENDED WAIVERS	4. FSES 5. PERFORMANCE BASED DESIGN	
B. THE FACILITY DOES NOT MEET THE S	STANDARD				
SURVEYOR (EKimberly Swense	TITLE	OFFICE		DATE	
SURVEYOR ID					
K10		055105			
FIRE AUTHORITY OFFICIAL (Signature)	TITLE	OFFICE		DATE	
CMS FORMS SHALL BE COMPLETED AND RETAIL	AINED AS PART OF THE SURVEY RECORD.	·			

ID PREFIX		MET	NOT MET	N/A	REMARKS
	PART I – NFPA 101 LSC REQUIREMENTS (Items in italics relate to the FSES)				
	SECTION 1 – GENERAL REQUIREMENTS				
K100	General Requirements – Other				
	List in the REMARKS section any LSC Section 18.1 and 19.1 General Requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567.				
K111	Building Rehabilitation				
	Repair, Renovation, Modification, or Reconstruction				
	Any building undergoing repair, renovation, modification, or reconstruction complies with both of the following:				
	Requirements of Chapter 18 and 19.				
	Requirements of the applicable Sections 43.3, 43.4, 43.5, and 43.6.				
	18.1.1.4.3, 19.1.1.4.3, 43.1.2.1				
	Change of Use or Change of Occupancy				
	Any building undergoing change of use or change of occupancy classification complies with the requirements of Section 43.7, unless permitted by 18.1.1.4.2 or 19.1.1.4.2.				
	18.1.1.4.2 (4.6.7 and 4.6.11), 19.1.1.4.2 (4.6.7 and 4.6.11), 43.1.2.2 (43.7)				
	Additions				
	Any building undergoing an addition shall comply with the requirements of Section 43.8. If the building has a common wall with a nonconforming building, the common wall is a fire barrier having at least a two hour fire resistance rating constructed of materials as required for the addition.				
	Communicating openings occur only in corridors and are protected by approved self-closing fire doors with at least a 1-1/2 hour fire resistance rating. Additions comply with the requirements of Section 43.8.				
	18.1.1.4.1 (4.6.7 and 4.6.11), 18.1.1.4.1.1 (8.3), 18.1.1.4.1.2, 18.1.1.4.1.3, 19.1.1.4.1 (4.6.7 and 4.6.11), 19.1.1.4.1.1 (8.3), 19.1.1.4.1.2, 19.1.1.4.1.3, 43.1.2.3(43.8)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K112	Sprinkler Requirements for Major Rehabilitation If a nonsprinklered smoke compartment has undergone major rehabilitation the automatic sprinkler requirements of 18.3.5 have been applied to the smoke compartment. In cases where the building is not protected throughout by a sprinkler system, the requirements of 18.4.3.2, 18.4.3.3, and 18.4.3.8 are also met. Note: Major rehabilitation involves the modification of more than 50 percent, or more than 4500 ft² of the area of the smoke compartment. 18.1.1.4.3.3, 19.1.1.4.3.3				
K131	 Multiple Occupancies – Sections of Health Care Facilities Sections of health care facilities classified as other occupancies meet all of the following: They are not intended to serve four or more inpatients for purposes of housing, treatment, or customary access. They are separated from areas of health care occupancies by construction having a minimum two hour fire resistance rating in accordance with Chapter 8. The entire building is protected throughout by an approved, supervised automatic sprinkler system in accordance with Section 9.7. Hospital outpatient surgical departments are required to be classified as an Ambulatory Health Care Occupancy regardless of the number of patients served. 18.1.3.3, 19.1.3.3, 42 CFR 482.41, 42 CFR 485.623 				
K132	Multiple Occupancies – Contiguous Non-Health Care Occupancies Non-health care occupancies that are located immediately next to a Health Care Occupancy, but are primarily intended to provide outpatient services are permitted to be classified as Business or Ambulatory Health Care Occupancies, provided the facilities are separated by construction having not less than two hour fire resistance-rated construction, and are not intended to provide services simultaneously for four or more inpatients. Outpatient surgical departments must be classified as Ambulatory Health Care Occupancy regardless of the number of patients served. 18.1.3.4.1, 19.1.3.4.1				

ID PREFIX					MET	NOT MET	N/A	REMARKS
K133	Mι	ıltiple	Occupancies - Constructi	on Type				
	Where separated occupancies are in accordance with 18/19.1.3.2 or 18/19.1.3.4, the most stringent construction type is provided throughout the building, unless a two hour separation is provided in accordance with 8.2.1.3, in which case the construction type is determined as follows:							
	The construction type and supporting construction of the health care occupancy is based on the story in which it is located in the building in accordance with 18/19.1.6 and Tables 18/19.1.6.1.							
	The construction type of the areas of the building enclosing the other occupancies shall be based on the applicable occupancy chapters.							
16101			, 19.1.3.5, 8.2.1.3	• • •				
K161		_	g Construction Type and He ISTING	eight				
				meets Table 19.1.6.1, unless				
			e permitted by 19.1.6.2 throu					
	19.1.6.4, 19.1.6.5							
			Construction Type					
	/	1	l (442), l (332), ll (222)	Any number of stories non-sprinklered or sprinklered	ı			
	2	2	II (111)	One story non-sprinklered Maximum 3 stories sprinklered				
	3	3	II (000)					
	2	4	III (211)	Not allowed non-sprinklered				
	5	5	IV (2HH)	Maximum 2 stories sprinklered				
	6	3	V (111)					
		7	III (200)	Not allowed non-sprinklered				
	8	8	V (000)	Maximum 1 story sprinklered				
				ed throughout by an approved, rdance with section 9.7. (See 19.3.5)				
	inc fire	luding barrie	basements, floors on which pa	f the construction, the number of stories, atients are located, location of smoke or amplete sketch or attach small floor				

ID PREFIX				MET	NOT MET	N/A	REMARKS
K161	2012 NEW Building construction type and stories meets Table 18.1.6.1, unless otherwise permitted by 18.1.6.2 through 18.1.6.7 18.1.6.4, 18.1.6.5						
		Construction Type					
	1	I (442), I (332), II (222)	Not allowed non-sprinklered Any number of stories sprinklered				
	2	II (111)	Not allowed non-sprinklered Maximum 3 stories sprinklered				
	3	II (000)	Not allowed non-sprinklered Maximum 1 story sprinklered				
	4	III (211)					
	5	IV (2HH)					
	6	V (111)					
	7	III (200)	Not allowed non-sprinklered				
	8	V (000)	·				
	Sprinklered stories must be sprinklered throughout by an approved, supervised automatic system in accordance with section 9.7. (See 18.3.5) Give a brief description, in REMARKS, of the construction, the number of stories, including basements, floors on which patients are located, location of smoke or fire barriers and dates of approval. Complete sketch or attach small floor plan of the building as appropriate.						
K162		n <mark>g Systems Involving Comb</mark> u EXISTING	stibles				
	Buildings of Type I (442), Type I (332), Type II (222), or Type II (111) having roof systems employing combustible roofing supports, decking or roofing meet the following:						
		of covering meets Class C requ					
	no	2. roof is separated from occupied building portions with a noncombustible floor assembly using not less than 2½ inches concrete or gypsum fill.					
	attic or other space is either unoccupied or protected throughout by an approved automatic sprinkler system.						
	19.1.6.2*, ASTM E108, ANSI/UL 790						

ID PREFIX		MET	NOT MET	N/A	REMARKS
K162	2012 NEW				
	Buildings of Type I (442), Type I (332), Type II (222), Type II (111) having roof systems employing combustible roofing supports, decking or roofing meet the following:				
	roof covering meets Class A requirements.				
	2. roof is separated from occupied building portions with 2 hour fire resistive noncombustible floor assembly using not less than 2½ inches concrete or gypsum fill.				
	 the structural elements supporting the rated floor assembly meet the required fire resistance rating of the building. 				
	18.1.6.2, ASTM E108, ANSI/UL 790				
K163	Interior Nonbearing Wall Construction				
	Interior nonbearing walls in Type I or II construction are constructed of noncombustible or limited-combustible materials.				
	Interior nonbearing walls required to have a minimum 2 hour fire resistance rating are permitted to be fire-retardant-treated wood enclosed within noncombustible or limited-combustible materials, provided they are not used as shaft enclosures.				
	18.1.6.4, 18.1.6.5, 19.1.6.4, 19.1.6.5				
	SECTION 2 – MEANS OF EGRESS REQUIREMENTS				
K200	Means of Egress Requirements – Other				
	List in the REMARKS section any LSC Section 18.2 and 19.2 Means of Egress requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567.				
	18.2, 19.2				
K211	Means of Egress – General				
	Aisles, passageways, corridors, exit discharges, exit locations, and accesses are in accordance with Chapter 7, and the means of egress is continuously maintained free of all obstructions to full use in case of emergency, unless modified by 18/19.2.2 through 18/19.2.11.				
	18.2.1, 19.2.1, 7.1.10.1				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K221	Patient Sleeping Room Doors Locks on patient sleeping room doors are not permitted unless the keylocking device that restricts access from the corridor does not restrict egress from the patient room, or the locking arrangement is permitted for patient clinical, security or safety needs in accordance with 18.2.2.2.5 or 19.2.2.2.5. 18.2.2.2, 19.2.2.2, TIA 12-4				
K222	Egress Doors Doors in a required means of egress shall not be equipped with a latch or a lock that requires the use of a tool or key from the egress side unless using one of the following special locking arrangements:				
	□ CLINICAL NEEDS OR SECURITY THREAT LOCKING Where special locking arrangements for the clinical security needs of the patient are used, only one locking device shall be permitted on each door and provisions shall be made for the rapid removal of occupants by: remote control of locks; keying of all locks or keys carried by staff at all times; or other such reliable means available to the staff at all times. 18.2.2.2.5.1, 18.2.2.2.6, 19.2.2.2.5.1, 19.2.2.2.6				
	□ SPECIAL NEEDS LOCKING ARRANGEMENTS Where special locking arrangements for the safety needs of the patient are used, all of the Clinical or Security Locking requirements are being met. In addition, the locks must be electrical locks that fail safely so as to release upon loss of power to the device; the building is protected by a supervised automatic sprinkler system and the locked space is protected by a complete smoke detection system (or is constantly monitored at an attended location within the locked space); and both the sprinkler and detection systems are arranged to unlock the doors upon activation. 18.2.2.2.5.2, 19.2.2.2.5.2, TIA 12-4				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K222	□ DELAYED-EGRESS LOCKING ARRANGEMENTS Approved, listed delayed-egress locking systems installed in accordance with 7.2.1.6.1 shall be permitted on door assemblies serving low and ordinary hazard contents in buildings protected throughout by an approved, supervised automatic fire detection system or an approved, supervised automatic sprinkler system. 18.2.2.2.4, 19.2.2.2.4 □ ACCESS-CONTROLLED EGRESS LOCKING ARRANGEMENTS Access-Controlled Egress Door assemblies installed in accordance with 7.2.1.6.2 shall be permitted. 18.2.2.2.4, 19.2.2.2.4 □ ELEVATOR LOBBY EXIT ACCESS LOCKING ARRANGEMENTS Elevator lobby exit access door locking in accordance with 7.2.1.6.3 shall be permitted on door assemblies in buildings protected throughout by an approved, supervised automatic fire detection system and an approved, supervised automatic sprinkler system. 18.2.2.2.4, 19.2.2.2.4				
K223	Doors with Self-Closing Devices Doors in an exit passageway, stairway enclosure, or horizontal exit, smoke barrier, or hazardous area enclosure are self-closing and kept in the closed position, unless held open by a release device complying with 7.2.1.8.2 that automatically closes all such doors throughout the smoke compartment or entire facility upon activation of: Required manual fire alarm system; and Local smoke detectors designed to detect smoke passing through the opening or a required smoke detection system; and Automatic sprinkler system, if installed; and Loss of power. 18.2.2.2.7, 18.2.2.2.8, 19.2.2.2.7, 19.2.2.2.8				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K224	Horizontal-Sliding Doors				
	Horizontal-sliding doors permitted by 7.2.1.14 that are not automatic-closing are limited to a single leaf and shall have a latch or other mechanism to ensure the door will not rebound.				
	Horizontal-sliding doors serving an occupant load fewer than 10 shall be permitted, providing all of the following criteria are met:				
	Area served by the door has no high hazard contents.				
	Door is operable from either side without special knowledge or effort.				
	 Force required to operate the door in the direction of travel is ≤ 30 lbf to set the door in motion and ≤ 15 lbf to close or open to the required width. 				
	Assembly is appropriately fire rated, and where rated, is self-or automatic-closing by smoke detection per 7.2.1.8, and installed per NFPA 80.				
	Where required to latch, the door has a latch or other mechanism to ensure the door will not rebound.				
	18.2.2.2.10, 19.2.2.2.10				
K225	Stairways and Smokeproof Enclosures				
	Stairways and Smokeproof enclosures used as exits are in accordance with 7.2.				
	18.2.2.3, 18.2.2.4, 19.2.2.3, 19.2.2.4, 7.2				
K226	Horizontal Exits				
	Horizontal exits, if used, are in accordance with 7.2.4 and the provisions of 18.2.2.5.1 through 18.2.2.5.7, or 19.2.2.5.1 through 19.2.2.5.4.				
	18.2.2.5, 19.2.2.5				
K227	Ramps and Other Exits				
	Ramps, exit passageways, fire and slide escapes, alternating tread devices, and areas of refuge are in accordance with the provisions 7.2.5 through 7.2.12.				
	18.2.2.6 to 18.2.2.10 or 19.2.2.6 to 19.2.2.10				
K231	Means of Egress Capacity				
	The capacity of required means of egress is in accordance with 7.3. 18.2.3.1, 19.2.3.1				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K232	Aisle, Corridor or Ramp Width				
	2012 EXISTING				
	The width of aisles or corridors (clear or unobstructed) serving as exit access shall be at least 4 feet and maintained to provide the convenient removal of nonambulatory patients on stretchers, except as modified by 19.2.3.4, exceptions 1-5.				
	19.2.3.4, 19.2.3.5				
	2012 NEW				
	The width of aisles or corridors (clear and unobstructed) serving as exit access in hospitals and nursing homes shall be at least 8 feet. In limited care facility and psychiatric hospitals, width of aisles or corridors shall be at least 6 feet, except as modified by the 18.2.3.4 or 18.2.3.5 exceptions.				
	18.2.3.4, 18.2.3.5				
K233	Clear Width of Exit and Exit Access Doors				
	2012 EXISTING				
	Exit access doors and exit doors are of the swinging type and are at least 32 inches in clear width. Exceptions are provided for existing 34-inch doors and for existing 28-inch doors where the fire plan does not require evacuation by bed, gurney, or wheelchair. 19.2.3.6, 19.2.3.7				
	2012 NEW				
	Exit access doors and exit doors are of the swinging type and are at least 41.5 inches in clear width. In psychiatric hospitals or limited care facilities, doors are at least 32 inches wide. Doors not subject to patient use, in exit stairway enclosures, or serving newborn nurseries shall be no less than 32 inches in clear width. If using a pair of doors, the doors shall be provided with a rabbet, bevel, or astragal at the meeting edge, at least one of the doors shall provide 32 inches in clear width, and the inactive leaf of the pair shall be secured with automatic flush bolts.				
1/044	18.2.3.6, 18.2.3.7				
K241	Number of Exits – Story and Compartment Not less than two exits, remote from each other, and accessible from every				
	part of every story are provided for each story. Each smoke compartment shall likewise be provided with two distinct egress paths to exits that do not require the entry into the same adjacent smoke compartment.				
	18.2.4.1-18.2.4.4, 19.2.4.1-19.2.4.4				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K251	Dead-End Corridors and Common Path of Travel 2012 EXISTING				
	Dead-end corridors shall not exceed 30 feet. Existing dead-end corridors greater than 30 feet shall be permitted to be continued to be used if it is impractical and unfeasible to alter them.				
	19.2.5.2				
K251	2012 NEW				
	Dead-end corridors shall not exceed 30 feet. Common path of travel shall not exceed 100 feet.				
	18.2.5.2, 18.2.5.3				
K252	Number of Exits – Corridors				
	Every corridor shall provide access to not less than two approved exits in accordance with Sections 7.4 and 7.5 without passing through any intervening rooms or spaces other than corridors or lobbies.				
	18.2.5.4, 19.2.5.4				
K253	Number of Exits – Patient Sleeping and Non-Sleeping Rooms				
	Patient sleeping rooms of more than 1,000 square feet or nonsleeping rooms of more than 2,500 square feet have at least two exit access doors remotely located from each other.				
	18.2.5.5.1, 18.2.5.5.2, 19.2.5.5.1, 19.2.5.5.2				
K254	Corridor Access				
	All habitable rooms not within suites have a door leading directly outside to grade or have a door leading to an exit access corridor. Patient sleeping rooms with less than eight patient beds may have one room intervening to reach an exit access corridor provided the intervening room is equipped with an approved automatic smoke detection system.				
	18.2.5.6.1 through 18.2.5.6.4, 19.2.5.6.1 through 19.2.5.6.4				
K255	Suite Separation, Hazardous Content, and Subdivision				
	All suites are separated from the remainder of the building (including from other suites) by construction meeting the separation provisions for corridor construction (18.3.6.2-18.3.6.5 or 19.3.6.2-19.3.6.5). Existing approved barriers shall be allowed to continue to be used provided they limit the transfer of smoke. Intervening rooms have no hazardous areas and hazardous areas within suites comply with 18/19.2.5.7.1.3. Subdivision of suites shall be by noncombustible or limited-combustible construction. 18.2.5.7.1.2 through 18.2.5.7.1.4, 19.2.5.7.1.2, 19.2.5.7.1.3, 19.2.5.7.1.4				

ID PREFIX		MET	NOT MET	N/A	REMARKS
	Sleeping Suites Occupants shall have exit access to a corridor or direct access to a horizontal exit. Where ≥ 2 exits are required, one exit access door may be to a stairway, passageway or to the exterior. Suites shall be provided with constant staff supervision. Staff shall have direct visual supervision of patient sleeping rooms, from a constantly attended location or the room shall be provided with an automatic smoke detection system. Suites more than 1,000 ft² shall have 2 or more remote exits. One means of egress from the suite shall be to a corridor and one may be into an adjacent suite separated in accordance with corridor requirements. Suites shall not exceed the following size limitations: • 5,000 square feet if the suite is not fully smoke detected or fully sprinklered. • 7,500 square feet if the suite is either fully smoke detected and fully sprinklered and the sleeping rooms have direct supervision from a constantly attended location. Travel distance between any point in a suite to exit access shall not exceed 100 feet and distance to an exit shall not exceed 150 feet (200 feet if	MET		N/A	REMARKS
	building is fully sprinklered). 18.2.5.7.2, 19.2.5.7.2				
K257	Non-Sleeping Suites Occupants shall have exit access to a corridor or direct access to a horizontal exit. Where ≥ 2 exits are required, one exit access door may be to a stairway, passageway or to the exterior. Suites more than 2,500 ft² shall have 2 or more remote exits. One means of egress from the suite shall be to a corridor and one may be into an adjacent suite separated in accordance with corridor requirements. Suites shall not exceed 10,000 ft². Travel distance between any point in a suite to exit access shall not exceed 100 feet and distance to an exit shall not exceed 150 feet (200 feet if building is fully sprinklered). 18.2.5.7.3, 19.2.5.7.3				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K261	Travel Distance to Exits				
	Travel distance (excluding suites) to exits are measured in accordance with 7.6.				
	 From any point in the room or suite to exit less than or equal to 150 feet (less than or equal to 200 feet if the building is fully sprinklered). 				
	 Point in a room to room door less than or equal to 50 feet. 				
	18.2.6, 19.2.6				
K271	Discharge from Exits				
	Exit discharge is arranged in accordance with 7.7, provides a level walking surface meeting the provisions of 7.1.7 with respect to changes in elevation and shall be maintained free of obstructions. Additionally, the exit discharge shall be a hard packed all-weather travel surface. 18.2.7, 19.2.7				
K281	Illumination of Means of Egress				
	Illumination of means of egress, including exit discharge, is arranged in accordance with 7.8 and shall be either continuously in operation or capable of automatic operation without manual intervention. 18.2.8, 19.2.8				
K291	Emergency Lighting				
	Emergency lighting of at least 1-1/2 hour duration is provided automatically in accordance with 7.9.				
	18.2.9.1, 19.2.9.1				
K292	Life Support Means of Egress 2012 NEW (INDICATE N/A FOR EXISTING)				
	Buildings equipped with or requiring the use of life support systems (electro- mechanical or inhalation anesthetics) have illumination of means of egress, emergency lighting equipment, exit, and directional signs supplied by the life safety branch of the electrical system described in NFPA 99.				
	(Indicate N/A if life support equipment is for emergency purposes only.) 18.2.9.2, 18.2.10.5				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K293	Exit Signage		IVIEI		
N293	2012 EXISTING				
	Exit and directional signs are displayed in accordance with 7.10 with continuous illumination also served by the emergency lighting system. 19.2.10.1				
	(Indicate N/A in one-story existing occupancies with less than 30 occupants where the line of exit travel is obvious.)				
	2012 NEW				
	Exit and directional signs are displayed in accordance with 7.10 with continuous illumination also served by the emergency lighting system. 18.2.10.1				
	SECTION 3 – PROTECTION				
K300	Protection – Other				
	List in the REMARKS section any LSC Section 18.3 and 19.3 Protection requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567.				
K311	Vertical Openings – Enclosure 2012 EXISTING				
	Stairways, elevator shafts, light and ventilation shafts, chutes, and other vertical openings between floors are enclosed with construction having a fire resistance rating of at least 1-hour. An atrium may be used in accordance with 8.6.				
	19.3.1.1 through 19.3.1.6				
	If all vertical openings are properly enclosed with construction providing at least a 2 hour fire resistance rating, also check this box. □				
	2012 NEW				
	Stairways, elevator shafts, light and ventilation shafts, chutes, and other vertical openings between floors are enclosed with construction having a fire resistance rating of at least 2 hours connecting four or more stories. (1-hour for single story building and buildings up to three stories in height.) An atrium may be used in accordance with 8.6.7. 18.3.1 through 18.3.1.5				

ID PREFIX						MET	NOT MET	N/A	REMARKS
K321	Hazardous Areas – Enclosure 2012 EXISTING Hazardous areas are protected by resistance rating (with ¾ hour fire rextinguishing system in accordance approved automatic fire extinguishing shall be separated from other space doors in accordance with 8.4. Door closing and permitted to have nonrethat do not exceed 48 inches from Describe the floor and zone location in REMARKS. 19.3.2.1, 19.3.5.9	rated doors) or an a e with 8.7.1 or 19.3 ing system option i es by smoke resist rs shall be self-clos rated or field-applie the bottom of the d	automatic fir 3.5.9. When s used, the ting partition sing or autor d protective door.	the the areas is and matic- plates	S				
	Area	Automatic Sprinkler	Separation	N/A	1				
	a. Boiler and Fuel-Fired Heater Rooms								
	b. Laundries (larger than 100 sq. ft.)				-				
	c. Repair, Maintenance, and Paint Shops								
	d. Soiled Linen Rooms (exceeding 64 gal.) e. Trash Collection Rooms (exceeding 64 gal.)				-				
	f. Combustible Storage Rooms/Spaces (over 50 sq. ft.)				-				
	g. Laboratories (if classified as Severe Hazard - see K322)								

ID PREFIX						MET	NOT MET	N/A	REMARKS
K321	2012 NEW								
	Hazardous areas are protected in a shall be enclosed with a 1-hour fire door without windows (in accordant closing or automatic-closing in accordant protected by a sprinkler system 8.4.	e-rated barrier, with ice with 8.7.1.1). Do ordance with 7.2.1	a ¾ hour fi oors shall b .8. Hazardo	re-rate e self- us are	ed eas				
	Describe the floor and zone locations of hazardous areas that are deficient in REMARKS.								
	18.3.2.1, 7.2.1.8, 8.4, 8.7, 9.7								
	Area	Automatic Sprinkler	Separation	N/A					
	a. Boiler and Fuel-Fired Heater Rooms								
	b. Laundries (larger than 100 sq. ft.)								
	c. Repair, Maintenance, and Paint Shops								
	d. Soiled Linen Rooms (exceeding 64 gal.)								
	e. Trash Collection Rooms (exceeding 64 gal.)								
	f. Combustible Storage Rooms/Spaces (over 50 and less than 100 sq. ft.)								
	g. Combustible Storage Rooms/Spaces (over 100 sq. ft.)								
	h. Laboratories (if classified as Severe Hazard - see K322)								

ID PREFIX		MET	NOT MET	N/A	REMARKS
ID PREFIX	Laboratories Laboratories employing quantities of flammable, combustible, or hazardous materials that are considered a severe hazard are protected by 1-hour fire resistance-rated separation, automatic sprinkler system, and are in accordance with 8.7 and with NFPA 99. Laboratories not considered a severe hazard are protected as hazardous areas (see K321). Laboratories using chemicals are in accordance with NFPA 45, Standard on Fire Protection for Laboratories Using Chemicals. Gas appliances are of appropriate design and installed in accordance with NFPA 54. Shutoff valves are marked to identify material they control. Devices requiring medical grade oxygen from the piped distribution system meet the requirements under 11.4.2.2 (NFPA 99).	MET	NOT MET	N/A	REMARKS
	18.3.2.2, 19.3.2.2, 8.7, 8.7.4.1 (LSC) 9.3.1.2, 11.4.3.2, 15.4 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K323	Anesthetizing Locations				
	Areas designated for administration of general anesthesia (i.e., inhalation anesthetics) are in accordance with 8.7 and NFPA 99.				
	Zone valves are: located immediately outside each life-support, critical care, and anesthetizing location of moderate sedation, deep sedation, or general anesthesia for medical gas or vacuum; readily accessible in an emergency; and arranged so shutting off any one anesthetizing location will not affect others.				
	Area alarm panels are provided to monitor all medical gas, medical-surgical vacuum, and piped WAGD systems. Panels are at locations that provide for surveillance, indicate medical gas pressure decreases of 20 percent and vacuum decreases of 12 inch gauge HgV, and provide visual and audible indication. Alarm sensors are installed either on the source side of individual room zone valve box assemblies or on the patient/use side of each of the individual zone box valve assemblies.				
	The EES critical branch supplies power for task illumination, fixed equipment, select receptacles, and select power circuits, and EES equipment system supplies power to ventilation system.				
	Heating, cooling, and ventilation are in accordance with ASHRAE 170. Medical supply and equipment manufacturer's instructions for use are considered before reducing humidity levels to those allowed by ASHRAE, per S&C 13-58.				
	18.3.2.3, 19.3.2.3 (LSC) 5.1.4.8.7, 5.1.4.8.7.2, 5.1.9.3, 5.1.9.3.4, 6.4.2.2.4.2 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K324	Cooking Facilities				
	Cooking equipment is protected in accordance with NFPA 96, Standard for Ventilation Control and Fire Protection of Commercial Cooking Operations, unless:				
	 residential cooking equipment (i.e., small appliances such as microwaves, hot plates, toasters) are used for food warming or limited cooking in accordance with 18.3.2.5.2, 19.3.2.5.2. 				
	 cooking facilities open to the corridor in smoke compartments with 30 or fewer patients comply with the conditions under 18.3.2.5.3, 19.3.2.5.3, or 				
	 cooking facilities in smoke compartments with 30 or fewer patients comply with conditions under 18.3.2.5.4, 19.3.2.5.4. 				
	Cooking facilities protected according to NFPA 96 per 9.2.3 are not required to be enclosed as hazardous areas, but shall not be open to the corridor.				
	18.3.2.5.1 through 18.3.2.5.4, 19.3.2.5.1 through 19.3.2.5.5, 9.2.3, TIA 12-2				
K325	Alcohol Based Hand Rub Dispenser (ABHR)				
	ABHRs are protected in accordance with 8.7.3.1, unless all conditions are met:				
	Corridor is at least 6 feet wide.				
	 Maximum individual dispenser capacity is 0.32 gallons (0.53 gallons in suites) of fluid and 18 ounces of Level 1 aerosols. 				
	Dispensers shall have a minimum of four foot horizontal spacing.				
	Not more than an aggregate of 10 gallons of fluid or 1135 ounces of aerosol are used in a single smoke compartment outside a storage cabinet, excluding one individual dispenser per room.				
	Storage in a single smoke compartment greater than 5 gallons complies with NFPA 30.				
	Dispensers are not installed within 1 inch of an ignition source.				
	Dispensers over carpeted floors are in sprinklered smoke compartments.				
	ABHR does not exceed 95 percent alcohol.				
	Operation of the dispenser shall comply with Section 18.3.2.6(11) or 19.3.2.6(11).				
	ABHR is protected against inappropriate access.				
	18.3.2.6, 19.3.2.6, 42 CFR Parts 403, 418, 460, 482, 483, and 485				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K331	Interior Wall and Ceiling Finish 2012 EXISTING Interior wall and ceiling finishes, including exposed interior surfaces of buildings such as fixed or movable walls, partitions, columns, and have a flame spread rating of Class A or Class B. The reduction in class of interior finish for a sprinkler system as prescribed in 10.2.8.1 is permitted. 10.2, 19.3.3.1, 19.3.3.2 Indicate flame spread rating(s).				
	Interior wall and ceiling finishes, including exposed interior surfaces of buildings such as fixed or movable walls, partitions and columns have a flame spread rating of Class A. The reduction in class of interior finish for a sprinkler system as prescribed in 10.2.8.1 is permitted. Individual rooms not exceeding four persons may have a Class A or B finish. Lower half of corridor walls, not exceeding 4 feet in height, may have a Class A or B flame spread rating. 10.2, 18.3.3.1, 18.3.3.2 Indicate flame spread rating(s).				
K332	Interior Floor Finish 2012 NEW (Indicate N/A for 2012 EXISTING) Interior finishes shall comply with 10.2. Floor finishes in exit enclosures and exit access corridors and spaces not separated by walls that resist the passage of smoke shall be Class I or II. 18.3.3.3.1, 18.3.3.3.2, 18.3.3.3.3, 10.2, 10.2.7.1, 10.2.7.2				
K341	Fire Alarm System – Installation A fire alarm system is installed with systems and components approved for the purpose in accordance with NFPA 70, National Electric Code, and NFPA 72, National Fire Alarm Code to provide effective warning of fire in any part of the building. In areas not continuously occupied, detection is installed at each fire alarm control unit. In new occupancy, detection is also installed at notification appliance circuit power extenders, and supervising station transmitting equipment. Fire alarm system wiring or other transmission paths are monitored for integrity. 18.3.4.1, 19.3.4.1, 9.6, 9.6.1.8				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K342	Fire Alarm System – Initiation				
	Initiation of the fire alarm system is by manual means and by any required sprinkler system alarm, detection device, or detection system. Manual alarm boxes are provided in the path of egress near each required exit. Manual alarm boxes in patient sleeping areas shall not be required at exits if manual alarm boxes are located at all nurse's stations or other continuously attended staff location, provided alarm boxes are visible, continuously accessible, and 200' travel distance is not exceeded. 18.3.4.2.1, 18.3.4.2.2, 19.3.4.2.1, 19.3.4.2.2, 9.6.2.5				
K343	Fire Alarm – Notification				
	2012 EXISTING				
	Positive alarm sequence in accordance with 9.6.3.4 are permitted in buildings protected throughout by a sprinkler system. Occupant notification is provided automatically in accordance with 9.6.3 by audible and visual signals.				
	In critical care areas, visual alarms are sufficient. The fire alarm system transmits the alarm automatically to notify emergency forces in the event of a fire.				
	19.3.4.3, 19.3.4.3.1, 19.3.4.3.2, 9.6.4, 9.7.1.1(1)				
	2012 NEW				
	Positive alarm sequence in accordance with 9.6.3.4 are permitted. Occupant notification is provided automatically in accordance with 9.6.3 by audible and visual signals.				
	In critical care areas, visual alarms are sufficient. The fire alarm system transmits the alarm automatically to notify emergency forces in the event of a fire.				
	Annunciation and annunciation zoning for fire alarm and sprinklers shall be provided by audible and visual indicators and zones shall not be larger than 22,500 square feet per zone.				
	18.3.4.3 through 18.3.4.3.3, 9.6.4				
K344	Fire Alarm – Control Functions				
	The fire alarm automatically activates required control functions and is provided with an alternative power supply in accordance with NFPA 72.				
	18.3.4.4, 19.3.4.4, 9.6.1, 9.6.5, NFPA 72				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K345	Fire Alarm System – Testing and Maintenance A fire alarm system is tested and maintained in accordance with an approved program complying with the requirements of NFPA 70, National Electric Code, and NFPA 72, National Fire Alarm and Signaling Code. Records of system acceptance, maintenance and testing are readily available.				
K346	9.6.1.3, 9.6.1.5, NFPA 70, NFPA 72 Fire Alarm – Out of Service Where required fire alarm system is out of services for more than 4 hours in a 24 hour period, the authority having jurisdiction shall be notified, and the building shall be evacuated or an approved fire watch shall be provided for all parties left unprotected by the shutdown until the fire alarm system has been returned to service. 9.6.1.6				
K347	Smoke Detection 2012 EXISTING Smoke detection systems are provided in spaces open to corridors as required by 19.3.6.1. 19.3.4.5.2				
	2012 NEW Smoke detection systems are provided in spaces open to corridors as required by 18.3.6.1 In nursing homes, an automatic smoke detection system is installed in the corridors of all smoke compartments containing resident sleeping rooms, unless the resident sleeping rooms have: • smoke detection, or • automatic door closing devices with integral smoke detectors on the room side that provide occupant notification. Such detectors are electrically interconnected to the fire alarm system. 18.3.4.5.2, 18.3.4.5.3				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K351	Sprinkler System – Installation 2012 EXISTING				
	Nursing homes, and hospitals where required by construction type, are protected throughout by an approved automatic sprinkler system in accordance with NFPA 13, Standard for the Installation of Sprinkler Systems.				
	In Type I and II construction, alternative protection measures are permitted to be substituted for sprinkler protection in specific areas where state or local regulations prohibit sprinklers.				
	In hospitals, sprinklers are not required in clothes closets of patient sleeping rooms where the area of the closet does not exceed 6 ft² and sprinkler coverage covers the closet footprint as required by NFPA 13, Standard for Installation of Sprinkler Systems.				
	19.3.5.1, 19.3.5.2, 19.3.5.3, 19.3.5.4, 19.3.5.5, 19.4.2, 19.3.5.10, 9.7, 9.7.1.1(1)				
	2012 NEW				
	Buildings are to be protected throughout by an approved automatic sprinkler system in accordance with NFPA 13, Standard for the Installation of Sprinkler Systems.				
	In Type I and II construction, alternative protection measures are permitted to be substituted for sprinkler protection in specific areas where State and local regulations prohibit sprinklers.				
	Listed quick-response or listed residential sprinklers are used throughout smoke compartments with patient sleeping rooms.				
	In hospitals, sprinklers are not required in clothes closets of patient sleeping rooms where the area of the closet does not exceed 6 ft² and sprinkler coverage covers the closet footprint as required by NFPA 13, Standard for Installation of Sprinkler Systems.				
	18.3.5.1, 18.3.5.4, 18.3.5.5, 18.3.5.6, 9.7, 9.7.1.1(1), 18.3.5.10				
K352	Sprinkler System – Supervisory Signals				
	Automatic sprinkler system supervisory attachments are installed and monitored for integrity in accordance with NFPA 72, <i>National Fire Alarm and Signaling Code</i> , and provide a signal that sounds and is displayed at a continuously attended location or approved remote facility when sprinkler operation is impaired.				
l	9.7.2.1, NFPA 72				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K353	Sprinkler System – Maintenance and Testing Automatic sprinkler and standpipe systems are inspected, tested, and maintained in accordance with NFPA 25, Standard for the Inspection, Testing, and Maintaining of Water-based Fire Protection Systems. Records of system design, maintenance, inspection and testing are maintained in a secure location and readily available. a) Date sprinkler system last checked. b) Who provided system test. c) Water system supply source. Provide in REMARKS information on coverage for any non-required or partial automatic sprinkler system. 9.7.5, 9.7.7, 9.7.8, and NFPA 25				
K354	Sprinkler System – Out of Service Where the sprinkler system is impaired, the extent and duration of the impairment has been determined, areas or buildings involved are inspected and risks are determined, recommendations are submitted to management or designated representative, and the fire department and other authorities having jurisdiction have been notified. Where the sprinkler system is out of service for more than 10 hours in a 24 hour period, the building or portion of the building affected are evacuated or an approved fire watch is provided until the sprinkler system has been returned to service. 18.3.5.1, 19.3.5.1, 9.7.5, 15.5.2 (NFPA 25) Portable Fire Extinguishers Portable fire extinguishers are selected, installed, inspected, and				
K361	maintained in accordance with NFPA 10, <i>Standard for Portable Fire Extinguishers</i> . 18.3.5.12, 19.3.5.12, NFPA 10 Corridors – Areas Open to Corridor Spaces (other than patient sleeping rooms, treatment rooms and hazardous areas), waiting areas, nurse's stations, gift shops, and cooking facilities, open to the corridor are in accordance with the criteria under 18.3.6.1 and 19.3.6.1. 18.3.6.1, 19.3.6.1				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K362	Corridors – Construction of Walls 2012 EXISTING Corridors are separated from use areas by walls constructed with at least ½ hour fire resistance rating. In fully sprinklered smoke compartments, partitions are only required to resist the transfer of smoke. In nonsprinklered buildings, walls extend to the underside of the floor or roof deck above the ceiling. Corridor walls may terminate at the underside of ceilings where specifically permitted by Code. Fixed fire window assemblies in corridor walls are in accordance with Section 8.3, but in sprinklered compartments there are no restrictions in area or fire resistance of glass or frames. If the walls have a fire resistance rating, give the rating if the walls terminate at the underside of the ceiling, give brief description in	=	MET		
	REMARKS, describing the ceiling throughout the floor area. 19.3.6.2, 19.3.6.2.7 2012 NEW Corridor walls shall form a barrier to limit the transfer of smoke. Such walls shall be permitted to terminate at the ceiling where the ceiling is constructed to limit the transfer of smoke. No fire resistance rating is required for the corridor walls. 18.3.6.2				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K363	Corridor – Doors 2012 EXISTING Doors protecting corridor openings in other than required enclosures of vertical openings, exits, or hazardous areas resist the passage of smoke and are made of 1¾ inch solid-bonded core wood or other material capable of resisting fire for at least 20 minutes. Doors in fully sprinklered smoke compartments are only required to resist the passage of smoke. Corridor doors and doors to rooms containing flammable or combustible materials have positive latching hardware. Roller latches are prohibited by CMS regulation. These requirements do not apply to auxiliary spaces that do not contain flammable or combustible material. Powered doors complying with 7.2.1.9 are permissible if provided with a device capable of keeping the door closed when a force of 5lbf is applied, whether or not power is applied. Clearance between bottom of door and floor covering is not exceeding 1 inch. There is no impediment to the closing of the doors. Hold open devices that release when the door is pushed or pulled are permitted. Nonrated protective plates of unlimited height are permitted. Dutch doors meeting 19.3.6.3.6 are permitted. Door frames shall be labeled and made of steel or other materials in compliance with 8.3, unless the smoke compartment is sprinklered. Fixed fire window assemblies are allowed per 8.3. In sprinklered compartments there are no restrictions in area or fire resistance of glass or frames in window assemblies. 19.3.6.3, 42 CFR Parts 403, 418, 460, 482, 483, and 485 Show in REMARKS details of doors such as fire protection ratings, automatics closing devices, etc.				
	Doors protecting corridor openings shall be constructed to resist the passage of smoke. Corridor doors and doors to rooms containing flammable or combustible materials have self-latching and positive latching hardware. Roller latches are prohibited by CMS regulation. These requirements do not apply to auxiliary spaces that do not contain flammable or combustible material. Powered doors complying with 7.2.1.9 are permissible if provided with a device capable of keeping the door closed when a force of 5lbf is applied, whether or not power is applied. Clearance between bottom of door and floor covering is not exceeding 1 inch. There is no impediment to the closing of the doors. Hold open devices that release when the door is pushed or pulled are permitted. Nonrated protective plates of unlimited height are permitted. Dutch doors meeting 18.3.6.3.6 are permitted. 18.3.6.3, 42 CFR Parts 403, 418, 460, 482, 483, and 485 Show in REMARKS details of doors such as fire protection ratings, automatic closing devices, etc.				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K364	Corridor – Openings				
	Transfer grilles are not used in corridor walls or doors. Auxiliary spaces that do not contain flammable or combustible materials are permitted to have louvers or be undercut.				
	In other than smoke compartments containing patient sleeping rooms, miscellaneous openings are permitted in vision panels or doors, provided the openings per room do not exceed 20 in² and are at or below half the distance from floor to ceiling. In sprinklered rooms, the openings per room do not exceed 80 in².				
	Vision panels in corridor walls or doors shall be fixed window assemblies in approved frames. (In fully sprinklered smoke compartments, there are no restrictions in the area and fire resistance of glass and frames.) 18.3.6.5.1, 19.3.6.5.2, 8.3				
K371	Subdivision of Building Spaces – Smoke Compartments				
	2012 EXISTING				
	Smoke barriers shall be provided to form at least two smoke compartments on every sleeping floor with a 30 or more patient bed capacity. Size of compartments cannot exceed 22,500 square feet or a 200-foot travel distance from any point in the compartment to a door in the smoke barrier.				
	19.3.7.1, 19.3.7.2				
	Detail in REMARKS zone dimensions including length of zones and deadend corridors.				
	2012 NEW				
	Smoke barriers shall be provided to form at least two smoke compartments on every floor used by inpatients for sleeping or treatment, and on every floor with an occupant load of 50 or more persons, regardless of use.				
	Size of compartments cannot exceed 22,500 square feet or a 200-foot travel distance from any point in the compartment to a door in the smoke barrier.				
	Smoke subdivision requirements do not apply to any of the stories or areas described in 18.3.7.2.				
	18.3.7.1, 18.3.7.2				
	Detail in REMARKS zone dimensions including length of zones and deadend corridors.				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K372	Subdivision of Building Spaces – Smoke Barrier Construction		IVILI		
	2012 EXISTING				
	Smoke barriers shall be constructed to a ½ hour fire resistance rating per 8.5. Smoke barriers shall be permitted to terminate at an atrium wall. Smoke dampers are not required in duct penetrations in fully ducted HVAC systems where an approved sprinkler system is installed for smoke compartments adjacent to the smoke barrier.				
	19.3.7.3, 8.6.7.1(1)				
	Describe any mechanical smoke control system in REMARKS.				
	2012 NEW				
	Smoke barriers shall be constructed to provide at least a 1-hour fire resistance rating and constructed in accordance with 8.5. Smoke barriers shall be permitted to terminate at an atrium wall. Smoke dampers are not required in duct penetrations of fully ducted HVAC systems.				
	18.3.7.3, 18.3.7.4, 18.3.7.5, 8.3				
	Describe any mechanical smoke control system in REMARKS.				
K373	Subdivision of Building Spaces – Accumulation Space Space shall be provided on each side of smoke barriers to adequately accommodate the total number of occupants in adjoining compartments. 18.3.7.5.1, 18.3.7.5.2, 19.3.7.5.1, 19.3.7.5.2				
K374	Subdivision of Building Spaces – Smoke Barrier Doors 2012 EXISTING Doors in smoke barriers are 1¾-inch thick solid bonded wood-core doors or of construction that resists fire for 20 minutes. Nonrated protective plates of unlimited height are permitted. Doors are permitted to have fixed fire window assemblies per 8.5. Doors are self-closing or automatic-closing, do not require latching, and are not required to swing in the direction of egress travel. Door opening provides a minimum clear width of 32 in for swinging or horizontal doors. 19.3.7.6, 19.3.7.8, 19.3.7.9				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K374	2012 NEW				
	Doors in smoke barriers have at least a 20-minute fire protection rating or are at least 1¾-inch thick solid bonded core wood.				
	Required clear widths are provided per 18.3.7.6(4) and (5).				
	Nonrated protective plates of unlimited height are permitted. Horizontal- sliding doors comply with 7.2.1.14. Swinging doors shall be arranged so that each door swings in an opposite direction.				
	Doors shall be self-closing and rabbets, bevels, or astragals are required at the meeting edges. Positive latching is not required.				
	18.3.7.6, 18.3.7.7, 18.3.7.8				
K379	Smoke Barrier Door Glazing				
	2012 EXISTING				
	Openings in smoke barrier doors shall be fire-rated glazing or wired glass panels in steel frames.				
	19.3.7.6, 19.3.7.6.2, 8.5				
	2012 NEW				
	Windows in smoke barrier doors shall be installed in each cross corridor swinging or horizontal-sliding door protected by fire-rated glazing or by wired glass panels in approved frames.				
	18.3.7.9				
K381	Sleeping Room Outside Windows and Doors				
	Every patient sleeping room has an outside window or outside door. In new occupancies, sill height does not exceed 36 inches above the floor. Windows in atrium walls are considered outside windows. Newborn nurseries and rooms intended for occupancy less than 24 hours have no outside window or door requirements. Window sills in special nursing care areas (e.g., ICU, CCU, hemodialysis, neonatal) do not exceed 60 inches above the floor.				
	42 CFR 403, 418, 460, 482, 483, and 485				
1446	SECTION 4 – SPECIAL PROVISIONS				
K400	Special Provisions – Other				
	List in the REMARKS section any LSC Section 18.4 and 19.4 Special Provisions requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or				
	NFPA standard citation, should be included on Form CMS-2567.				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K421	High-Rise Buildings 2012 EXISTING High-rise buildings are protected throughout by an approved, supervised automatic sprinkler system in accordance with Section 9.7 within 12 years of LSC final rule effective date. 19.4.2				
	2012 NEW High-rise buildings comply with section 11.8. 18.4.2				
	SECTION 5 – BUILDING SERVICES				
K500	Building Services – Other List in the REMARKS section any LSC Section 18.5 and 19.5 Building Services requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567.				
K511	Utilities – Gas and Electric				
	Equipment using gas or related gas piping complies with NFPA 54, <i>National Fuel Gas Code</i> , electrical wiring and equipment complies with NFPA 70, <i>National Electric Code</i> . Existing installations can continue in service provided no hazard to life.				
	18.5.1.1, 19.5.1.1, 9.1.1, 9.1.2				
K521	HVAC Heating, ventilation, and air conditioning shall comply with 9.2 and shall be installed in accordance with the manufacturer's specifications. 18.5.2.1, 19.5.2.1, 9.2				
K522	HVAC – Any Heating Device				
	Any heating device, other than a central heating plant, is designed and installed so combustible materials cannot be ignited by device, and has a safety feature to stop fuel and shut down equipment if there is excessive temperature or ignition failure. If fuel fired, the device also:				
	is chimney or vent connected.				
	takes air for combustion from outside.				
	 provides for a combustion system separate from occupied area atmosphere. 18.5.2.2, 19.5.2.2 				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K523	HVAC – Suspended Unit Heaters				
	Suspended unit heaters are permitted provided the following are met:				
	Not located in means of egress or in patient rooms.				
	Located high enough to be out of reach of people in the area.				
	Has a safety feature to stop fuel and shut down equipment if there is excessive temperature or ignition failure.				
	18.5.2.3(1), 19.5.2.3(1)				
K524	HVAC - Direct-Vent Gas Fireplaces				
	Direct-vent gas fireplaces, as defined in NFPA 54, inside of all smoke compartments containing patient sleeping areas comply with the requirements of 18.5.2.3(2), 19.5.2.3(2). 18.5.2.3(2), 19.5.2.3(2), NFPA 54				
K525	HVAC - Solid Fuel-Burning Fireplaces				
	Solid fuel-burning fireplaces are permitted in other than patient sleeping areas provided:				
	Areas are separated by 1-hour fire resistance construction.				
	Fireplace complies with 9.2.2.				
	Fireplace enclosure resists breakage up to 650°F and has heat- tempered glass.				
	Room has supervised CO detection per 9.8.				
	18.5.2.3(3) and 19.5.2.3(3)				
K531	Elevators				
	2012 EXISTING				
	Elevators comply with the provision of 9.4. Elevators are inspected and tested as specified in ASME A17.1, Safety Code for Elevators and Escalators. Firefighter's Service is operated monthly with a written record.				
	Existing elevators conform to ASME/ANSI A17.3, Safety Code for Existing Elevators and Escalators. All existing elevators, having a travel distance of 25 feet or more above or below the level that best serves the needs of emergency personnel for firefighting purposes, conform with Firefighter's Service Requirements of ASME/ANSI A17.3. (Includes firefighter's service Phase I key recall and smoke detector automatic recall, firefighter's service Phase II emergency in-car key operation, machine room smoke detectors, and elevator lobby smoke detectors.)				
	19.5.3, 9.4.2, 9.4.3				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K531	2012 NEW Elevators comply with the provision of 9.4. Elevators are inspected and tested as specified in ASME A17.1, <i>Safety Code for Elevators and Escalators</i> . Firefighter's Service is operated monthly with a written record. New elevators conform to ASME/ANSI A17.1, <i>Safety Code for Elevators and Escalators</i> , including Firefighter's Service Requirements. (Includes firefighter's Phase I key recall and smoke detector automatic recall, firefighter's service Phase II emergency in-car key operation, machine room smoke detectors, and elevator lobby smoke detectors.) 18.5.3, 9.4.2, 9.4.3				
K532	Escalators, Dumbwaiters, and Moving Walks 2012 EXISTING Escalators, dumbwaiters, and moving walks comply with the provisions of 9.4. All existing escalators, dumbwaiters, and moving walks conform to the requirements of ASME/ANSI A17.3, Safety Code for Existing Elevators and Escalators. (Includes escalator emergency stop buttons and automatic skirt obstruction stop. For power dumbwaiters, includes hoistway door locking to keep doors closed except for floor where car is being loaded or unloaded.) 19.5.3, 9.4.2.2				
	2012 NEW Escalators, dumbwaiters, and moving walks comply with the provisions of 9.4. 18.5.3, 9.4.2.2				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K541	Rubbish Chutes, Incinerators, and Laundry Chutes				
	2012 EXISTING				
	(1) Any existing linen and trash chute, including pneumatic rubbish and linen systems, that opens directly onto any corridor shall be sealed by fire resistive construction to prevent further use or shall be provided with a fire door assembly having a fire protection rating of 1-hour. All new chutes shall comply with 9.5.				
	(2) Any rubbish chute or linen chute, including pneumatic rubbish and linen systems, shall be provided with automatic extinguishing protection in accordance with 9.7.				
	(3) Any trash chute shall discharge into a trash collection room used for no other purpose and protected in accordance with 8.4. (Existing laundry chutes permitted to discharge into same room are protected by automatic sprinklers in accordance with 19.3.5.9 or 19.3.5.7.)				
	(4) Existing fuel-fed incinerators shall be sealed by fire resistive construction to prevent further use.				
	19.5.4, 9.5, 8.4, NFPA 82				
	2012 NEW				
	Rubbish chutes, incinerators, and laundry chutes shall comply with the provisions of Section 9.5, unless otherwise specified in 18.5.4.2.				
	The fire resistance rating of chute charging room shall not be required to exceed 1-hour.				
	 Any rubbish chute or linen chute shall be provided with automatic extinguishing protection in accordance with Section 9.7. 				
	 Chutes shall discharge into a trash collection room used for no other purpose and shall be protected in accordance with 8.7. 18.5.4.2, 8.7, 9.5, 9.7, NFPA 82 				
	SECTION 6 – RESERVED				
	SECTION 6 - RESERVED SECTION 7 - OPERATING FEATURES				
1/700					
K700	Operating Features – Other List in the REMARKS section any LSC Section 18.7 and 19.7 Operating				
	Features requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or				
	NFPA standard citation, should be included in Form CMS-2567.				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K711	Evacuation and Relocation Plan				
	There is a written plan for the protection of all patients and for their evacuation in the event of an emergency.				
	Employees are periodically instructed and kept informed with their duties under the plan, and a copy of the plan is readily available with telephone operator or with security. The plan addresses the basic response required of staff per 18/19.7.2.1.2 and provides for all of the fire safety plan components per 18/19.7.2.2.				
	18.7.1.1 through 18.7.1.3, 18.7.2.1.2, 18.7.2.2, 18.7.2.3, 19.7.1.1 through 19.7.1.3, 19.7.2.1.2, 19.7.2.2, 19.7.2.3				
K712	Fire Drills				
K712	Fire Drills Fire drills include the transmission of a fire alarm signal and simulation of emergency fire conditions. Fire drills are held at expected and unexpected times under varying conditions, at least quarterly on each shift. The staff is familiar with procedures and is aware that drills are part of established routine. Where drills are conducted between 9:00 PM and 6:00 AM, a coded announcement may be used instead of audible alarms. 18.7.1.4 through 18.7.1.7, 19.7.1.4 through 19.7.1.7				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K741	 Smoking Regulations Smoking regulations shall be adopted and shall include not less than the following provisions: Smoking shall be prohibited in any room, ward, or compartment where flammable liquids, combustible gases, or oxygen is used or stored and in any other hazardous location, and such area shall be posted with signs that read NO SMOKING or shall be posted with the international symbol for no smoking. In health care occupancies where smoking is prohibited and signs are prominently placed at all major entrances, secondary signs with language that prohibits smoking shall not be required. Smoking by patients classified as not responsible shall be prohibited. The requirement of 18.7.4(3) shall not apply where the patient is under direct supervision. Ashtrays of noncombustible material and safe design shall be provided in all areas where smoking is permitted. Metal containers with self-closing cover devices into which ashtrays can be emptied shall be readily available to all areas where smoking is permitted. 		MEI		
K751	Draperies, Curtains, and Loosely Hanging Fabrics Draperies, curtains including cubicle curtains and loosely hanging fabric or films shall be in accordance with 10.3.1. Excluding curtains and draperies: at showers and baths; on windows in patient sleeping room located in sprinklered compartments; and in non-patient sleeping rooms in sprinklered compartments where individual drapery or curtain panels do not exceed 48 square feet or total area does not exceed 20 percent of the wall. 18.7.5.1, 18.3.5.11, 19.7.5.1, 19.3.5.11, 10.3.1				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K752	Upholstered Furniture and Mattresses				
	Newly introduced upholstered furniture meets Class I or char length, and heat release criteria in accordance with 10.3.2.1 and 10.3.3, unless the building is fully sprinklered.				
	Newly introduced mattresses shall meet char length and heat release criteria in accordance with 10.3.2.2 and 10.3.4, unless the building is fully sprinklered.				
	Upholstered furniture and mattresses belonging to nursing home residents do not have to meet these requirements as all nursing homes are required to be fully sprinklered.				
	Newly introduced upholstered furniture and mattresses means purchased on or after the LSC final rule effective date.				
	18.7.5.2, 18.7.5.4, 19.7.5.2, 19.7.5.4				
K753	Combustible Decorations				
	Combustible decorations shall be prohibited unless one of the following is met:				
	Flame retardant or treated with approved fire-retardant coating that is listed and labeled for product.				
	Decorations meet NFPA 701.				
	 Decorations exhibit heat release less than 100 kilowatts in accordance with NFPA 289. 				
	• Decorations, such as photographs, paintings and other art are attached to the walls, ceilings and non-fire-rated doors in accordance with 18.7.5.6(4) or 19.7.5.6(4).				
	 The decorations in existing occupancies are in such limited quantities that a hazard of fire development or spread is not present. 18.7.5.6, 19.7.5.6 				
K761	Maintenance, Inspection & Testing - Doors				
	Fire doors assemblies are inspected and tested annually in accordance with NFPA 80 Standard for Fire Doors and Other Opening Protectives.				
	Fire doors that are not located in required fire barriers, including corridor doors to patient rooms and smoke barrier doors, are routinely inspected as part of the facility maintenance program.				
	Individuals performing the door inspection and testing have an understanding of the operating components of the doors. Written records of inspection and testing are maintained and are available for review.				
	18.7.6, 19.7.6, 8.3.3.1 (LSC), 5.2, 5.2.3 (NFPA 80)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K754	Soiled Linen and Trash Containers				
	Soiled linen or trash collection receptacles shall not exceed 32 gallons in capacity. The average density of container capacity in a room or space shall not exceed 0.5 gallons/square feet. A total container capacity of 32 gallons shall not be exceeded within any 64 square feet area. Mobile soiled linen or trash collection receptacles with capacities greater than 32 gallons shall be located in a room protected as a hazardous area when not attended. Containers used solely for recycling are permitted to be excluded from the				
	above requirements where each container is ≤ 96 gal. unless attended, and containers for combustibles are labeled and listed as meeting FM Approval Standard 6921 or equivalent.				
	18.7.5.7, 19.7.5.7				
K771	Engineer Smoke Control Systems 2012 EXISTING When installed, engineered smoke control systems are tested in accordance with established engineering principles. Test documentation is maintained on the premises.				
	19.7.7				
	2012 NEW When installed, engineered smoke control systems are tested in accordance with NFPA 92, <i>Standard for Smoke Control Systems</i> . Test documentation is maintained on the premises. 18.7.7				
K781	Portable Space Heaters Portable space heating devices shall be prohibited in all health care occupancies. Unless used in nonsleeping staff and employee areas where the heating elements do not exceed 212 degrees Fahrenheit (100 degrees Celsius). 18.7.8, 19.7.8				
K791	Construction, Repair, and Improvement Operations				
	Construction, repair, and improvement operations shall comply with 4.6.10. Any means of egress in any area undergoing construction, repair, or improvements shall be inspected daily to ensure its ability to be used instantly in case of emergency and compliance with NFPA 241.				
	18.7.9, 19.7.9, 4.6.10, 7.1.10.1				

ID PREFIX		MET	NOT MET	N/A	REMARKS
TREFFX	PART II – HEALTH CARE FACILITIES CODE REQUIREMENTS		IVIL		
K900	Health Care Facilities Code - Other List in the REMARKS section any NFPA 99 requirements (excluding Chapter 7, 8, 12, and 13) that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Health Care Facilities Code or NFPA standard citation, should be included on Form CMS-2567.				
K901	Fundamentals – Building System Categories Building systems are designed to meet Category 1 through 4 requirements as detailed in NFPA 99. Categories are determined by a formal and documented risk assessment procedure performed by qualified personnel. Chapter 4 (NFPA 99)				
K902	Gas and Vacuum Piped Systems – Other List in the REMARKS section any NFPA 99 Chapter 5 Gas and Vacuum Systems requirements that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567. Chapter 5 (NFPA 99)				
K903	Gas and Vacuum Piped Systems – Categories Medical gas, medical air, surgical vacuum, WAGD, and air supply systems are designated: □ Category 1. Systems in which failure is likely to cause major injury or death. □ Category 2. Systems in which failure is likely to cause minor injury. □ Category 3. Systems in which failure is not likely to cause injury, but can cause discomfort. Deep sedation and general anesthesia are not to be administered using a Category 3 medical gas system. 5.1.1.1, 5.2.1, 5.3.1.1, 5.3.1.5 (NFPA 99)				
K904	Gas and Vacuum Piped Systems – Warning Systems All master, area, and local alarm systems used for medical gas and vacuum systems comply with appropriate Category warning system requirements, as applicable. 5.1.9, 5.2.9, 5.3.6.2.2 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K905	Gas and Vacuum Piped Systems – Central Supply System Identification and Labeling Containers, cylinders and tanks are designed, fabricated, tested, and marked in accordance with 5.1.3.1.1 through 5.1.3.1.7. Locations				
	containing only oxygen or medical air have doors labeled with "Medical Gases, NO Smoking or Open Flame". Locations containing other gases have doors labeled "Positive Pressure Gases, NO Smoking or Open Flame, Room May Have Insufficient Oxygen, Open Door and Allow Room to Ventilate Before Opening."				
14000	5.1.3.1, 5.2.3.1, 5.3.10 (NFPA 99)				
K906	Gas and Vacuum Piped Systems – Central Supply System Operations Adaptors or conversion fittings are prohibited. Cylinders are handled in accordance with 11.6.2. Only cylinders, reusable shipping containers, and their accessories are stored in rooms containing central supply systems or cylinders. No flammable materials are stored with cylinders. Cryogenic liquid storage units intended to supply the facility are not used to transfill. Cylinders are kept away from sources of heat. Valve protection caps are secured in place, if supplied, unless cylinder is in use. Cylinders are not stored in tightly closed spaces. Cylinders in use and storage are prevented from exceeding 130°F, and nitrous oxide and carbon dioxide cylinders are prevented from reaching temperatures lower than manufacture recommendations or 20°F. Full or empty cylinders, when not connected, are stored in locations complying with 5.1.3.3.2 through 5.1.3.3.3, and are not stored in enclosures containing motor-driven machinery, unless for instrument air reserve headers. 5.1.3.2, 5.1.3.3.17, 5.1.3.3.1.8, 5.1.3.3.4, 5.2.3.2, 5.2.3.3, 5.3.6.20.4, 5.6.20.5, 5.3.6.20.7, 5.3.6.20.8, 5.3.6.20.9 (NFPA 99)				
K907	Gas and Vacuum Piped Systems – Maintenance Program				
	Medical gas, vacuum, WAGD, or support gas systems have documented maintenance programs. The program includes an inventory of all source systems, control valves, alarms, manufactured assemblies, and outlets. Inspection and maintenance schedules are established through risk assessment considering manufacturer recommendations. Inspection procedures and testing methods are established through risk assessment. Persons maintaining systems are qualified as demonstrated by training and certification or credentialing to the requirements of AASE 6030 or 6040. 5.1.14.2.1, 5.1.14.2.2, 5.1.15, 5.2.14, 5.3.13.4.2 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K908	Gas and Vacuum Piped Systems – Inspection and Testing Operations				
	The gas and vacuum systems are inspected and tested as part of a maintenance program and include the required elements. Records of the inspections and testing are maintained as required. 5.1.14.2.3, B.5.2, 5.2.13, 5.3.13, 5.3.13.4 (NFPA 99)				
K909	Gas and Vacuum Piped Systems – Information and Warning Signs				
	Piping is labeled by stencil or adhesive markers identifying the gas or vacuum system, including the name of system or chemical symbol, color code (Table 5.1.11), and operating pressure if other than standard. Labels are at intervals not more than 20 feet, are in every room, at both sides of wall penetrations, and on every story traversed by riser. Piping is not painted. Shutoff valves are identified with the name or chemical symbol of the gas or vacuum system, room or area served, and caution to not use the valve except in emergency. 5.1.14.3, 5.1.11.1, 5.1.11.2, 5.2.11, 5.3.13.3, 5.3.11 (NFPA 99)				
K910	Gas and Vacuum Piped Systems – Modifications				
	Whenever modifications are made that breach the pipeline, any necessary installer and verification test specified in 5.1.2 is conducted on the downstream portion of the medical gas piping system. Permanent records of all tests required by system verification tests are maintained. 5.1.14.4.1, 5.1.14.4.6, 5.2.13, 5.3.13.4.3 (NFPA 99)				
K911	Electrical Systems – Other				
	List in the REMARKS section any NFPA 99 Chapter 6 Electrical Systems requirements that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567. Chapter 6 (NFPA 99)				
K912	Electrical Systems – Receptacles				
	Power receptacles have at least one, separate, highly dependable grounding pole capable of maintaining low-contact resistance with its mating plug. In pediatric locations, receptacles in patient rooms, bathrooms, play rooms, and activity rooms, other than nurseries, are listed tamper-resistant or employ a listed cover.				
	If used in patient care room, ground-fault circuit interrupters (GFCI) are listed.				
	6.3.2.2.6.2 (F), 6.3.2.2.4.2 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K913	Electrical Systems – Wet Procedure Locations Operating rooms are considered wet procedure locations, unless otherwise determined by a risk assessment conducted by the facility governing body. Operating rooms defined as wet locations are protected by either isolated power or ground-fault circuit interrupters. A written record of the risk assessment is maintained and available for inspection. 6.3.2.2.8.4, 6.3.2.2.8.7, 6.4.4.2				
K914	Electrical Systems – Maintenance and Testing Hospital-grade receptacles at patient bed locations and where deep sedation or general anesthesia is administered, are tested after initial installation, replacement or servicing. Additional testing is performed at intervals defined by documented performance data. Receptacles not listed as hospital-grade at these locations are tested at intervals not exceeding 12 months. Line isolation monitors (LIM), if installed, are tested at intervals of ≤ 1 month by actuating the LIM test switch per 6.3.2.6.3.6, which activates both visual and audible alarm. For LIM circuits with automated self-testing, this manual test is performed at intervals ≤ 12 months. LIM circuits are tested per 6.3.3.3.2 after any repair or renovation to the electric distribution system. Records are maintained of required tests and associated repairs or modifications, containing date, room or area tested, and results. 6.3.4 (NFPA 99)				
K915	Electrical Systems – Essential Electric System Categories ☐ Critical care rooms (Category 1) in which electrical system failure is likely to cause major injury or death of patients, including all rooms where electric life support equipment is required, are served by a Type 1 EES. ☐ General care rooms (Category 2) in which electrical system failure is likely to cause minor injury to patients (Category 2) are served by a Type 1 or Type 2 EES. ☐ Basic care rooms (Category 3) in which electrical system failure is not likely to cause injury to patients and rooms other than patient care rooms are not required to be served by an EES. Type 3 EES life safety branch has an alternate source of power that will be effective for 1 1/2 hours. 3.3.138, 6.3.2.2.10, 6.6.2.2.2, 6.6.3.1.1 (NFPA 99), TIA 12-3				

	MET	NOT MET	N/A	REMARKS
Electrical Systems – Essential Electric System Alarm Annunciator		IVIEI		
A remote annunciator that is storage battery powered is provided to operate outside of the generating room in a location readily observed by operating personnel. The annunciator is hard-wired to indicate alarm conditions of the emergency power source. A centralized computer system (e.g., building information system) is not to be substituted for the alarm annunciator.				
· · ·				
Electrical Systems – Essential Electric System Receptacles Electrical receptacles or cover plates supplied from the life safety and critical branches have a distinctive color or marking. 6.4.2.2.6, 6.5.2.2.4.2, 6.6.2.2.3.2 (NFPA 99)				
Electrical Systems – Essential Electric System Maintenance and Testing				
The generator or other alternate power source and associated equipment is capable of supplying service within 10 seconds. If the 10-second criterion is not met during the monthly test, a process shall be provided to annually confirm this capability for the life safety and critical branches. Maintenance and testing of the generator and transfer switches are performed in accordance with NFPA 110.				
Generator sets are inspected weekly, exercised under load 30 minutes 12 times a year in 20-40 day intervals, and exercised once every 36 months for 4 continuous hours. Scheduled test under load conditions include a complete simulated cold start and automatic or manual transfer of all EES loads, and are conducted by competent personnel. Maintenance and testing of stored energy power sources (Type 3 EES) are in accordance with NFPA 111. Main and feeder circuit breakers are inspected annually, and a program for periodically exercising the components is established according to manufacturer requirements. Written records of maintenance and testing are maintained and readily available. EES electrical panels and circuits are marked, readily identifiable, and separate from normal power circuits. Minimizing the possibility of damage of the emergency power source is a design consideration for new installations. 6.4.4, 6.5.4, 6.6.4 (NFPA 99), NFPA 110, NFPA 111, 700.10 (NFPA 70)				
	outside of the generating room in a location readily observed by operating personnel. The annunciator is hard-wired to indicate alarm conditions of the emergency power source. A centralized computer system (e.g., building information system) is not to be substituted for the alarm annunciator. 6.4.1.1.17, 6.4.1.1.17.5 (NFPA 99) Electrical Systems – Essential Electric System Receptacles Electrical receptacles or cover plates supplied from the life safety and critical branches have a distinctive color or marking. 6.4.2.2.6, 6.5.2.2.4.2, 6.6.2.2.3.2 (NFPA 99) Electrical Systems – Essential Electric System Maintenance and Testing The generator or other alternate power source and associated equipment is capable of supplying service within 10 seconds. If the 10-second criterion is not met during the monthly test, a process shall be provided to annually confirm this capability for the life safety and critical branches. Maintenance and testing of the generator and transfer switches are performed in accordance with NFPA 110. Generator sets are inspected weekly, exercised under load 30 minutes 12 times a year in 20-40 day intervals, and exercised once every 36 months for 4 continuous hours. Scheduled test under load conditions include a complete simulated cold start and automatic or manual transfer of all EES loads, and are conducted by competent personnel. Maintenance and testing of stored energy power sources (Type 3 EES) are in accordance with NFPA 111. Main and feeder circuit breakers are inspected annually, and a program for periodically exercising the components is established according to manufacturer requirements. Written records of maintenance and testing are maintained and readily available. EES electrical panels and circuits are marked, readily identifiable, and separate from normal power circuits. Minimizing the possibility of damage of the emergency power source is a design consideration for new installations.	Electrical Systems – Essential Electric System Alarm Annunciator A remote annunciator that is storage battery powered is provided to operate outside of the generating room in a location readily observed by operating personnel. The annunciator is hard-wired to indicate alarm conditions of the emergency power source. A centralized computer system (e.g., building information system) is not to be substituted for the alarm annunciator. 6.4.1.1.17, 6.4.1.1.17.5 (NFPA 99) Electrical Systems – Essential Electric System Receptacles Electrical receptacles or cover plates supplied from the life safety and critical branches have a distinctive color or marking. 6.4.2.2.6, 6.5.2.2.4.2, 6.6.2.2.3.2 (NFPA 99) Electrical Systems – Essential Electric System Maintenance and Testing The generator or other alternate power source and associated equipment is capable of supplying service within 10 seconds. If the 10-second criterion is not met during the monthly test, a process shall be provided to annually confirm this capability for the life safety and critical branches. Maintenance and testing of the generator and transfer switches are performed in accordance with NFPA 110. Generator sets are inspected weekly, exercised under load 30 minutes 12 times a year in 20-40 day intervals, and exercised once every 36 months for 4 continuous hours. Scheduled test under load conditions include a complete simulated cold start and automatic or manual transfer of all EES loads, and are conducted by competent personnel. Maintenance and testing of stored energy power sources (Type 3 EES) are in accordance with NFPA 111. Main and feeder circuit breakers are inspected annually, and a program for periodically exercising the components is established according to manufacturer requirements. Written records of maintenance and testing are maintained and readily available. EES electrical panels and circuits are marked, readily identifiable, and separate from normal power circuits. Minimizing the possibility of damage of the emergency power source is a	Electrical Systems – Essential Electric System Alarm Annunciator A remote annunciator that is storage battery powered is provided to operate outside of the generating room in a location readily observed by operating personnel. The annunciator is hard-wired to indicate alarm conditions of the emergency power source. A centralized computer system (e.g., building information system) is not to be substituted for the alarm annunciator. 6.4.1.1.17, 6.4.1.1.17.5 (NFPA 99) Electrical Systems – Essential Electric System Receptacles Electrical receptacles or cover plates supplied from the life safety and critical branches have a distinctive color or marking. 6.4.2.2.6, 6.5.2.2.4.2, 6.6.2.2.3.2 (NFPA 99) Electrical Systems – Essential Electric System Maintenance and Testing The generator or other alternate power source and associated equipment is capable of supplying service within 10 seconds. If the 10-second criterion is not met during the monthly test, a process shall be provided to annually confirm this capability for the life safety and critical branches. Maintenance and testing of the generator and transfer switches are performed in accordance with NFPA 110. Generator sets are inspected weekly, exercised under load 30 minutes 12 times a year in 20-40 day intervals, and exercised once every 36 months for 4 continuous hours. Scheduled test under load conditions include a complete simulated cold start and automatic or manual transfer of all EES loads, and are conducted by competent personnel. Maintenance and testing of stored energy power sources (Type 3 EES) are in accordance with NFPA 111. Main and feeder circuit breakers are inspected annually, and a program for periodically exercising the components is established according to manufacturer requirements. Written records of maintenance and circuits are maintained and readily available. EES electrical panels and circuits are marked, readily identifiable, and separate from normal power circuits. Minimizing the possibility of damage of the emergency power source	Electrical Systems – Essential Electric System Alarm Annunciator A remote annunciator that is storage battery powered is provided to operate outside of the generating room in a location readily observed by operating personnel. The annunciator is hard-wired to indicate alarm conditions of the emergency power source. A centralized computer system (e.g., building information system) is not to be substituted for the alarm annunciator. 6.4.1.1.17, 6.4.1.1.17.5 (NFPA 99) Electrical Systems – Essential Electric System Receptacles Electrical branches have a distinctive color or marking. 6.4.2.2.6, 6.5.2.2.4.2, 6.6.2.2.3.2 (NFPA 99) Electrical Systems – Essential Electric System Maintenance and Testing The generator or other alternate power source and associated equipment is capable of supplying service within 10 seconds. If the 10-second criterion is not met during the monthly test, a process shall be provided to annually confirm this capability for the life safety and critical branches. Maintenance and testing of the generator and transfer switches are performed in accordance with NFPA 110. Generator sets are inspected weekly, exercised under load 30 minutes 12 times a year in 20-40 day intervals, and exercised once every 36 months for 4 continuous hours. Scheduled test under load conditions include a complete simulated cold start and automatic or manual transfer of all EES loads, and are conducted by competent personnel. Maintenance and testing of stored energy power sources (Type 3 EES) are in accordance with NFPA 111. Main and feeder circuit breakers are inspected annually, and a program for periodically exercising the components is established according to manufacturer requirements. Written records of maintenance and testing are maintained and readily available. EES electrical panels and circuits are marked, readily identifiable, and separate from normal power circuits. Minimizing the possibility of damage of the emergency power source is a design consideration for new installations.

ID PREFIX		MET	NOT MET	N/A	REMARKS
K919	Electrical Equipment – Other List in the REMARKS section any NFPA 99 Chapter 10, Electrical Equipment, requirements that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567. Chapter 10 (NFPA 99)				
K920	Electrical Equipment – Power Cords and Extension Cords Power strips in a patient care vicinity are only used for components of movable patient-care-related electrical equipment (PCREE) assembles that have been assembled by qualified personnel and meet the conditions of 10.2.3.6. Power strips in the patient care vicinity may not be used for non-PCREE (e.g., personal electronics), except in long-term care resident rooms that do not use PCREE. Power strips for PCREE meet UL 1363A or UL 60601-1. Power strips for non-PCREE in the patient care rooms (outside of vicinity) meet UL 1363. In non-patient care rooms, power strips meet other UL standards. All power strips are used with general precautions. Extension cords are not used as a substitute for fixed wiring of a structure. Extension cords used temporarily are removed immediately upon completion of the purpose for which it was installed and meets the conditions of 10.2.4. 10.2.3.6 (NFPA 99), 10.2.4 (NFPA 99), 400-8 (NFPA 70), 590.3(D) (NFPA 70), TIA 12-5				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K921	Electrical Equipment – Testing and Maintenance Requirements				
	The physical integrity, resistance, leakage current, and touch current tests for fixed and portable patient-care related electrical equipment (PCREE) is performed as required in 10.3. Testing intervals are established with policies and protocols. All PCREE used in patient care rooms is tested in accordance with 10.3.5.4 or 10.3.6 before being put into service and after any repair or modification. Any system consisting of several electrical appliances demonstrates compliance with NFPA 99 as a complete system. Service manuals, instructions, and procedures provided by the manufacturer include information as required by 10.5.3.1.1 and are considered in the development of a program for electrical equipment maintenance. Electrical equipment instructions and maintenance manuals are readily available, and safety labels and condensed operating instructions on the appliance are legible. A record of electrical equipment tests, repairs, and modifications is maintained for a period of time to demonstrate compliance in accordance with the facility's policy. Personnel responsible for the testing, maintenance and use of electrical appliances receive continuing training. 10.3, 10.5.2.1, 10.5.2.1.2, 10.5.2.5, 10.5.3, 10.5.6, 10.5.8				
K922	Gas Equipment – Other				
NJZZ	List in the REMARKS section any NFPA 99 Chapter 11 Gas Equipment requirements that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567. Chapter 11 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K923	Gas Equipment – Cylinder and Container Storage				
	≥ 3,000 cubic feet				
	Storage locations are designed, constructed, and ventilated in accordance with 5.1.3.3.2 and 5.1.3.3.3.				
	> 300 but <3,000 cubic feet				
	Storage locations are outdoors in an enclosure or within an enclosed interior space of non- or limited- combustible construction, with door (or gates outdoors) that can be secured. Oxidizing gases are not stored with flammables, and are separated from combustibles by 20 feet (5 feet if sprinklered) or enclosed in a cabinet of noncombustible construction having a minimum 1/2 hr. fire protection rating.				
	≤ 300 cubic feet				
	In a single smoke compartment, individual cylinders available for immediate use in patient care areas with an aggregate volume of ≤ 300 cubic feet are not required to be stored in an enclosure. Cylinders must be handled with precautions as specified in 11.6.2.				
	A precautionary sign readable from 5 feet is on each door or gate of a cylinder storage room, where the sign includes the wording as a minimum "CAUTION: OXIDIZING GAS(ES) STORED WITHIN NO SMOKING".				
	Storage is planned so cylinders are used in order of which they are received from the supplier. Empty cylinders are segregated from full cylinders. When facility employs cylinders with integral pressure gauge, a threshold pressure considered empty is established. Empty cylinders are marked to avoid confusion. Cylinders stored in the open are protected from weather.				
K924	11.3.1, 11.3.2, 11.3.3, 11.3.4, 11.6.5 (NFPA 99) Gas Equipment – Testing and Maintenance Requirements				
N924	Anesthesia apparatus are tested at the final path to patient after any adjustment, modification or repair. Before the apparatus is returned to service, each connection is checked to verify proper gas and an oxygen analyzer is used to verify oxygen concentration. Defective equipment is immediately removed from service. Areas designated for servicing of oxygen equipment are clean and free of oil, grease, or other flammables. Manufacturer service manuals are used to maintain equipment and a scheduled maintenance program is followed. 11.4.1.3, 11.5.1.3, 11.6.2.5, 11.6.2.6 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K925	Gas Equipment – Respiratory Therapy Sources of Ignition				
	Smoking materials are removed from patients receiving respiratory therapy. When a nasal cannula is delivering oxygen outside of a patient's room, no sources of ignition are within in the site of intentional expulsion (1-foot). When other oxygen deliver equipment is used or oxygen is delivered inside a patient's room, no sources of ignition are within the area are of administration (15-feet). Solid fuel-burning appliances is not in the area of administration. Nonmedical appliances with hot surfaces or sparking mechanisms are not within oxygen-delivery equipment or site of intentional expulsion. 11.5.1.1, TIA 12-6 (NFPA 99)				
K926	Gas Equipment – Qualifications and Training of Personnel				
	Personnel concerned with the application, maintenance and handling of medical gases and cylinders are trained on the risk. Facilities provide continuing education, including safety guidelines and usage requirements. Equipment is serviced only by personnel trained in the maintenance and operation of equipment. 11.5.2.1 (NFPA 99)				
K927	Gas Equipment – Transfilling Cylinders				
	Transfilling of oxygen from one cylinder to another is in accordance with CGA P-2.5, <i>Transfilling of High Pressure Gaseous Oxygen Used for Respiration</i> . Transfilling of any gas from one cylinder to another is prohibited in patient care rooms. Transfilling to liquid oxygen containers or to portable containers over 50 psi comply with conditions under 11.5.2.3.1 (NFPA 99). Transfilling to liquid oxygen containers or to portable containers under 50 psi comply with conditions under 11.5.2.3.2 (NFPA 99). 11.5.2.2 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K928	Gas Equipment – Labeling Equipment and Cylinders				
	Equipment listed for use in oxygen-enriched atmospheres are so labeled. Oxygen metering equipment and pressure reducing regulators are labeled "OXYGEN-USE NO OIL". Flowmeters, pressure reducing regulators, and oxygen-dispensing apparatus are clearly and permanently labeled designating the gases for which they are intended. Oxygen-metering equipment, pressure reducing regulators, humidifiers, and nebulizers are labeled with name of manufacturer or supplier. Cylinders and containers are labeled in accordance with CGA C-7. Color coding is not utilized as the primary method of determining cylinder or container contents. All labeling is durable and withstands cleaning or disinfecting.				
K929	11.5.3.1 (NFPA 99) Gas Equipment – Precautions for Handling Oxygen Cylinders and Manifolds				
	Handling of oxygen cylinders and manifolds is based on CGA G-4, Oxygen. Oxygen cylinders, containers, and associated equipment are protected from contact with oil and grease, from contamination, protected from damage, and handled with care in accordance with precautions provided under 11.6.2.1 through 11.6.2.4 (NFPA 99).				
K930	Gas Equipment – Liquid Oxygen Equipment				
	The storage and use of liquid oxygen in base reservoir containers and portable containers comply with sections 11.7.2 through 11.7.4 (NFPA 99). 11.7 (NFPA 99)				
K931	Hyperbaric Facilities				
	All occupancies containing hyperbaric facilities comply with construction, equipment, administration, and maintenance requirements of NFPA 99. Chapter 14 (NFPA 99)				
K932	Features of Fire Protection – Other List in the REMARKS section any NFPA 99 Chapter 15 Features of Fire Protection requirements that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567. Chapter 15 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K933	Peatures of Fire Protection – Fire Loss Prevention in Operating Rooms Periodic evaluations are made of hazards that could be encountered during surgical procedures, and fire prevention procedures are established. When flammable germicides or antiseptics are employed during surgeries utilizing electrosurgery, cautery or lasers: • packaging is non-flammable. • applicators are in unit doses. • Preoperative "time-out" is conducted prior the initiation of any surgical procedure to verify: • application site is dry prior to draping and use of surgical equipment. • pooling of solution has not occurred or has been corrected. • solution-soaked materials have been removed from the OR prior to draping and use of surgical devices. • policies and procedures are established outlining safety precautions related to the use of flammable germicide or antiseptic use. Procedures are established for operating room emergencies including alarm activation, evacuation, equipment shutdown, and control operations. Emergency procedures include the control of chemical spills, and extinguishment of drapery, clothing and equipment fires. Training is provided to new OR personnel (including surgeons), continuing education is provided, incidents are reviewed monthly, and procedures are reviewed annually. 15.13 (NFPA 99)				

Name of Facility 20	2012 LIFE SAFETY CODE
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PART III – RECOMMENDATION FOR WAIVER OF SPECIFIC LIFE SAFETY CODE PROVISIONS

For each item of the Life Safety Code recommended for waiver, list the survey report form item number and state the reason for the conclusion that: (a) the specific provisions of the code, if rigidly applied, would result in unreasonable hardship on the facility, and (b) the waiver of such unmet provisions will not adversely affect the health and safety of the patients. If additional space is required, attach additional sheet(s).

PROVISION NUMBER(S)	JUSTIFICATION
K400	

Title	Office	Date
Title	Office	Dete
ritie	Office	Date
	Title Title	

2012 LIFE SAFETY CODE

Form Approved OMB Exempt

FIRE SAFETY SURVEY REPORT HEALTHO	E				1. (B) MEDICAID I.D. NO.					
PART I — Life Safety Code, New and Existing PART II — Health Care Facilities Code, New and Existing PART III — Recommendation for Waiver PART IV – Crucial Data Extract OPTIONAL — Chapter 4 – NFPA 101A - Fire Safety Evaluation System for Health Care Occupancies – CMS-2786T										
Identifying information as shown in applicable r	ecords. Enter changes, if any, alon	gside each	item,	giving date o	f change.					
2. NAME OF FACILITY 2. (A)	MULTIPLE CONSTRUCTION (BLDGS) A. BUILDING B. WING C. FLOOR	2. (B) ADDRE	ESS OF	FACILITY (STR	EET, CITY, STA	TE, ZIP CODE)	A. Fully Sprinklered (All required areas are sprinklered) B. Partially Sprinklered (Not all required areas are sprinklered) C. None (No sprinkler system)			
3. SURVEY FOR 4. DA	TE OF SURVEY	DATE OF PL	AN APP	ROVAL	SURVEY UNDE	R	110100			
☐ MEDICARE ☐ MEDICAID		K6	52012 I			12 EXISTING 6. 2012 NEW				
5. SURVEY FOR CERTIFICATION OF 1. HOSPITAL 2. SKILLED/NURSING	FACILITY 4. ICF/IID UND	DER HEALTH	CARE	5.	HOSPICE					
IF "2" OR "5" ABOVE IS MARKED, CHECK APPROPRIAT 1 ENTIRE FACILITY 2 DISTINCT PART OF	` ,		_	3. IF DIST	_	OSPITAL, IS HO	SPITAL ACCREDITED?			
6. BED COMPOSITION a. TOTAL NO. OF BEDS IN THE FACILITY b. NUMBER OF HOSPI CERTIFIED FOR MEI				UMBER OF SKI ERTIFIED FOR	ILLED BEDS MEDICAID		R OF NF or ICF/IID BEDS ED FOR MEDICAID			
7. A. THE FACILITY MEETS THE STANDARD, BASED UPON (CHECK ALL APPROPRIATE BOXES) 1. COMPLIANCE WITH ALL PROVISIONS 2. ACCEPTANCE OF A PLAN OF CORRECTION 3. RECOMMENDED WAIVERS 4. FSES 5. PERFORMANCE BASED DESIGN B. THE FACILITY DOES NOT MEET THE STANDARD										
surveyor (E Kimberly Swenson	TITLE	OFFI	CE			DATE				
SURVEYOR ID K10										
FIRE AUTHORITY OFFICIAL (Signature)	TITLE	OFFI	CE			DATE				
CMS FORMS SHALL BE COMPLETED AND RETAINED	AS PART OF THE SURVEY RECORD.	·				•				

ID PREFIX		MET	NOT MET	N/A	REMARKS	
	PART I – NFPA 101 LSC REQUIREMENTS (Items in italics relate to the FSES)					
	SECTION 1 – GENERAL REQUIREMENTS					
K100	General Requirements – Other					
	List in the REMARKS section any LSC Section 18.1 and 19.1 General Requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567.					
K111	Building Rehabilitation					
	Repair, Renovation, Modification, or Reconstruction					
	Any building undergoing repair, renovation, modification, or reconstruction complies with both of the following:					
	Requirements of Chapter 18 and 19.					
	Requirements of the applicable Sections 43.3, 43.4, 43.5, and 43.6.					
	18.1.1.4.3, 19.1.1.4.3, 43.1.2.1					
	Change of Use or Change of Occupancy					
	Any building undergoing change of use or change of occupancy classification complies with the requirements of Section 43.7, unless permitted by 18.1.1.4.2 or 19.1.1.4.2.					
	18.1.1.4.2 (4.6.7 and 4.6.11), 19.1.1.4.2 (4.6.7 and 4.6.11), 43.1.2.2 (43.7)					
	Additions					
	Any building undergoing an addition shall comply with the requirements of Section 43.8. If the building has a common wall with a nonconforming building, the common wall is a fire barrier having at least a two hour fire resistance rating constructed of materials as required for the addition.					
	Communicating openings occur only in corridors and are protected by approved self-closing fire doors with at least a 1-1/2 hour fire resistance rating. Additions comply with the requirements of Section 43.8.					
	18.1.1.4.1 (4.6.7 and 4.6.11), 18.1.1.4.1.1 (8.3), 18.1.1.4.1.2, 18.1.1.4.1.3, 19.1.1.4.1 (4.6.7 and 4.6.11), 19.1.1.4.1.1 (8.3), 19.1.1.4.1.2, 19.1.1.4.1.3, 43.1.2.3(43.8)					

ID PREFIX		MET	NOT MET	N/A	REMARKS
K112	Sprinkler Requirements for Major Rehabilitation If a nonsprinklered smoke compartment has undergone major rehabilitation the automatic sprinkler requirements of 18.3.5 have been applied to the smoke compartment. In cases where the building is not protected throughout by a sprinkler system, the requirements of 18.4.3.2, 18.4.3.3, and 18.4.3.8 are also met. Note: Major rehabilitation involves the modification of more than 50 percent, or more than 4500 ft² of the area of the smoke compartment. 18.1.1.4.3.3, 19.1.1.4.3.3				
K131	 Multiple Occupancies – Sections of Health Care Facilities Sections of health care facilities classified as other occupancies meet all of the following: They are not intended to serve four or more inpatients for purposes of housing, treatment, or customary access. They are separated from areas of health care occupancies by construction having a minimum two hour fire resistance rating in accordance with Chapter 8. The entire building is protected throughout by an approved, supervised automatic sprinkler system in accordance with Section 9.7. Hospital outpatient surgical departments are required to be classified as an Ambulatory Health Care Occupancy regardless of the number of patients served. 18.1.3.3, 19.1.3.3, 42 CFR 482.41, 42 CFR 485.623 				
K132	Multiple Occupancies – Contiguous Non-Health Care Occupancies Non-health care occupancies that are located immediately next to a Health Care Occupancy, but are primarily intended to provide outpatient services are permitted to be classified as Business or Ambulatory Health Care Occupancies, provided the facilities are separated by construction having not less than two hour fire resistance-rated construction, and are not intended to provide services simultaneously for four or more inpatients. Outpatient surgical departments must be classified as Ambulatory Health Care Occupancy regardless of the number of patients served. 18.1.3.4.1, 19.1.3.4.1				

ID PREFIX					MET	NOT MET	N/A	REMARKS
K133	Multiple Occupancies – Construction Type							
	Where separated occupancies are in accordance with 18/19.1.3.2 or 18/19.1.3.4, the most stringent construction type is provided throughout the building, unless a two hour separation is provided in accordance with 8.2.1.3, in which case the construction type is determined as follows:							
	•	occu acco	pancy is based on the story indance with 18/19.1.6 and Ta					
	•	occu	pancies shall be based on th	s of the building enclosing the other e applicable occupancy chapters.				
14404			, 19.1.3.5, 8.2.1.3	• • •				
K161		_	Construction Type and He	eight				
	2012 EXISTING Building construction type and stories meets Table 19.1.6.1, unless							
			e permitted by 19.1.6.2 throu					
			, 19.1.6.5					
			Construction Type					
	/	1	l (442), l (332), ll (222)	Any number of stories non-sprinklered or sprinklered				
	2	2	II (111)	One story non-sprinklered Maximum 3 stories sprinklered				
	3	3	II (000)					
	2	4	III (211)	Not allowed non-sprinklered				
	5	5	IV (2HH)	Maximum 2 stories sprinklered				
	6	3	V (111)					
		7	III (200)	Not allowed non-sprinklered				
	8	8	V (000)	Maximum 1 story sprinklered				
	Sprinklered stories must be sprinklered throughout by an approved, supervised automatic system in accordance with section 9.7. (See 19.3.5)							
	Give a brief description, in REMARKS, of the construction, the number of stories, including basements, floors on which patients are located, location of smoke or fire barriers and dates of approval. Complete sketch or attach small floor plan of the building as appropriate.							

ID PREFIX				MET	NOT MET	N/A	REMARKS
K161	2012 NEW Building construction type and stories meets Table 18.1.6.1, unless otherwise permitted by 18.1.6.2 through 18.1.6.7 18.1.6.4, 18.1.6.5						
		Construction Type					
	1	I (442), I (332), II (222)	Not allowed non-sprinklered Any number of stories sprinklered				
	2	II (111)	Not allowed non-sprinklered Maximum 3 stories sprinklered				
	3	II (000)					
	4	III (211)	Not allowed non-sprinklered				
	5	IV (2HH)	Maximum 1 story sprinklered				
	6	V (111)					
	7	III (200)	Not allowed non-sprinklered				
	8	V (000)	·				
	Sprinklered stories must be sprinklered throughout by an approved, supervised automatic system in accordance with section 9.7. (See 18.3.5) Give a brief description, in REMARKS, of the construction, the number of stories, including basements, floors on which patients are located, location of smoke or fire barriers and dates of approval. Complete sketch or attach small floor plan of the building as appropriate.						
K162	Roofing Systems Involving Combustibles 2012 EXISTING						
	Buildings of Type I (442), Type I (332), Type II (222), or Type II (111) having roof systems employing combustible roofing supports, decking or roofing meet the following:						
		of covering meets Class C requ					
		ouilding portions with a sing not less than 2½ inches concrete					
	attic or other space is either unoccupied or protected throughout by an approved automatic sprinkler system.						
	19.1.6.2*, ASTM E108, ANSI/UL 790						

ID PREFIX		MET	NOT MET	N/A	REMARKS
K162	2012 NEW				
	Buildings of Type I (442), Type I (332), Type II (222), Type II (111) having roof systems employing combustible roofing supports, decking or roofing meet the following:				
	roof covering meets Class A requirements.				
	2. roof is separated from occupied building portions with 2 hour fire resistive noncombustible floor assembly using not less than 2½ inches concrete or gypsum fill.				
	the structural elements supporting the rated floor assembly meet the required fire resistance rating of the building.				
	18.1.6.2, ASTM E108, ANSI/UL 790				
K163	Interior Nonbearing Wall Construction				
	Interior nonbearing walls in Type I or II construction are constructed of noncombustible or limited-combustible materials.				
	Interior nonbearing walls required to have a minimum 2 hour fire resistance rating are permitted to be fire-retardant-treated wood enclosed within noncombustible or limited-combustible materials, provided they are not used as shaft enclosures.				
	18.1.6.4, 18.1.6.5, 19.1.6.4, 19.1.6.5				
	SECTION 2 – MEANS OF EGRESS REQUIREMENTS				
K200	Means of Egress Requirements – Other				
	List in the REMARKS section any LSC Section 18.2 and 19.2 Means of Egress requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567.				
	18.2, 19.2				
K211	Means of Egress – General				
	Aisles, passageways, corridors, exit discharges, exit locations, and accesses are in accordance with Chapter 7, and the means of egress is continuously maintained free of all obstructions to full use in case of emergency, unless modified by 18/19.2.2 through 18/19.2.11.				
	18.2.1, 19.2.1, 7.1.10.1				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K221	Patient Sleeping Room Doors Locks on patient sleeping room doors are not permitted unless the keylocking device that restricts access from the corridor does not restrict egress from the patient room, or the locking arrangement is permitted for patient clinical, security or safety needs in accordance with 18.2.2.2.5 or 19.2.2.2.5. 18.2.2.2, 19.2.2.2, TIA 12-4				
K222	Egress Doors Doors in a required means of egress shall not be equipped with a latch or a lock that requires the use of a tool or key from the egress side unless using one of the following special locking arrangements:				
	□ CLINICAL NEEDS OR SECURITY THREAT LOCKING Where special locking arrangements for the clinical security needs of the patient are used, only one locking device shall be permitted on each door and provisions shall be made for the rapid removal of occupants by: remote control of locks; keying of all locks or keys carried by staff at all times; or other such reliable means available to the staff at all times. 18.2.2.2.5.1, 18.2.2.2.6, 19.2.2.2.5.1, 19.2.2.2.6				
	□ SPECIAL NEEDS LOCKING ARRANGEMENTS Where special locking arrangements for the safety needs of the patient are used, all of the Clinical or Security Locking requirements are being met. In addition, the locks must be electrical locks that fail safely so as to release upon loss of power to the device; the building is protected by a supervised automatic sprinkler system and the locked space is protected by a complete smoke detection system (or is constantly monitored at an attended location within the locked space); and both the sprinkler and detection systems are arranged to unlock the doors upon activation. 18.2.2.2.5.2, 19.2.2.2.5.2, TIA 12-4				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K222	□ DELAYED-EGRESS LOCKING ARRANGEMENTS Approved, listed delayed-egress locking systems installed in accordance with 7.2.1.6.1 shall be permitted on door assemblies serving low and ordinary hazard contents in buildings protected throughout by an approved, supervised automatic fire detection system or an approved, supervised automatic sprinkler system. 18.2.2.2.4, 19.2.2.2.4 □ ACCESS-CONTROLLED EGRESS LOCKING ARRANGEMENTS Access-Controlled Egress Door assemblies installed in accordance with 7.2.1.6.2 shall be permitted. 18.2.2.2.4, 19.2.2.2.4 □ ELEVATOR LOBBY EXIT ACCESS LOCKING ARRANGEMENTS Elevator lobby exit access door locking in accordance with 7.2.1.6.3 shall be permitted on door assemblies in buildings protected throughout by an approved, supervised automatic fire detection system and an approved, supervised automatic sprinkler system. 18.2.2.2.4, 19.2.2.2.4				
K223	Doors with Self-Closing Devices Doors in an exit passageway, stairway enclosure, or horizontal exit, smoke barrier, or hazardous area enclosure are self-closing and kept in the closed position, unless held open by a release device complying with 7.2.1.8.2 that automatically closes all such doors throughout the smoke compartment or entire facility upon activation of: Required manual fire alarm system; and Local smoke detectors designed to detect smoke passing through the opening or a required smoke detection system; and Automatic sprinkler system, if installed; and Loss of power. 18.2.2.2.7, 18.2.2.2.8, 19.2.2.2.7, 19.2.2.2.8				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K224	Horizontal-Sliding Doors				
	Horizontal-sliding doors permitted by 7.2.1.14 that are not automatic-closing are limited to a single leaf and shall have a latch or other mechanism to ensure the door will not rebound.				
	Horizontal-sliding doors serving an occupant load fewer than 10 shall be permitted, providing all of the following criteria are met:				
	Area served by the door has no high hazard contents.				
	Door is operable from either side without special knowledge or effort.				
	 Force required to operate the door in the direction of travel is ≤ 30 lbf to set the door in motion and ≤ 15 lbf to close or open to the required width. 				
	Assembly is appropriately fire rated, and where rated, is self-or automatic-closing by smoke detection per 7.2.1.8, and installed per NFPA 80.				
	Where required to latch, the door has a latch or other mechanism to ensure the door will not rebound.				
	18.2.2.2.10, 19.2.2.2.10				
K225	Stairways and Smokeproof Enclosures				
	Stairways and Smokeproof enclosures used as exits are in accordance with 7.2.				
	18.2.2.3, 18.2.2.4, 19.2.2.3, 19.2.2.4, 7.2				
K226	Horizontal Exits				
	Horizontal exits, if used, are in accordance with 7.2.4 and the provisions of 18.2.2.5.1 through 18.2.2.5.7, or 19.2.2.5.1 through 19.2.2.5.4.				
	18.2.2.5, 19.2.2.5				
K227	Ramps and Other Exits				
	Ramps, exit passageways, fire and slide escapes, alternating tread devices, and areas of refuge are in accordance with the provisions 7.2.5 through 7.2.12.				
	18.2.2.6 to 18.2.2.10 or 19.2.2.6 to 19.2.2.10				
K231	Means of Egress Capacity				
	The capacity of required means of egress is in accordance with 7.3. 18.2.3.1, 19.2.3.1				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K232	Aisle, Corridor or Ramp Width 2012 EXISTING				
	The width of aisles or corridors (clear or unobstructed) serving as exit access shall be at least 4 feet and maintained to provide the convenient removal of nonambulatory patients on stretchers, except as modified by 19.2.3.4, exceptions 1-5.				
	19.2.3.4, 19.2.3.5 2012 NEW				
	The width of aisles or corridors (clear and unobstructed) serving as exit access in hospitals and nursing homes shall be at least 8 feet. In limited care facility and psychiatric hospitals, width of aisles or corridors shall be at least 6 feet, except as modified by the 18.2.3.4 or 18.2.3.5 exceptions.				
K233	18.2.3.4, 18.2.3.5 Clear Width of Exit and Exit Access Doors				
N233	2012 EXISTING				
	Exit access doors and exit doors are of the swinging type and are at least 32 inches in clear width. Exceptions are provided for existing 34-inch doors and for existing 28-inch doors where the fire plan does not require evacuation by bed, gurney, or wheelchair. 19.2.3.6, 19.2.3.7				
	2012 NEW				
	Exit access doors and exit doors are of the swinging type and are at least 41.5 inches in clear width. In psychiatric hospitals or limited care facilities, doors are at least 32 inches wide. Doors not subject to patient use, in exit stairway enclosures, or serving newborn nurseries shall be no less than 32 inches in clear width. If using a pair of doors, the doors shall be provided with a rabbet, bevel, or astragal at the meeting edge, at least one of the doors shall provide 32 inches in clear width, and the inactive leaf of the pair shall be secured with automatic flush bolts. 18.2.3.6, 18.2.3.7				
K241	Number of Exits – Story and Compartment				
	Not less than two exits, remote from each other, and accessible from every part of every story are provided for each story. Each smoke compartment shall likewise be provided with two distinct egress paths to exits that do not require the entry into the same adjacent smoke compartment. 18.2.4.1-18.2.4.4, 19.2.4.1-19.2.4.4				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K251	Dead-End Corridors and Common Path of Travel 2012 EXISTING				
	Dead-end corridors shall not exceed 30 feet. Existing dead-end corridors greater than 30 feet shall be permitted to be continued to be used if it is impractical and unfeasible to alter them.				
	19.2.5.2				
K251	2012 NEW				
	Dead-end corridors shall not exceed 30 feet. Common path of travel shall not exceed 100 feet.				
	18.2.5.2, 18.2.5.3				
K252	Number of Exits – Corridors				
	Every corridor shall provide access to not less than two approved exits in accordance with Sections 7.4 and 7.5 without passing through any intervening rooms or spaces other than corridors or lobbies.				
	18.2.5.4, 19.2.5.4				
K253	Number of Exits – Patient Sleeping and Non-Sleeping Rooms				
	Patient sleeping rooms of more than 1,000 square feet or nonsleeping rooms of more than 2,500 square feet have at least two exit access doors remotely located from each other.				
	18.2.5.5.1, 18.2.5.5.2, 19.2.5.5.1, 19.2.5.5.2				
K254	Corridor Access				
	All habitable rooms not within suites have a door leading directly outside to grade or have a door leading to an exit access corridor. Patient sleeping rooms with less than eight patient beds may have one room intervening to reach an exit access corridor provided the intervening room is equipped with an approved automatic smoke detection system.				
	18.2.5.6.1 through 18.2.5.6.4, 19.2.5.6.1 through 19.2.5.6.4				
K255	Suite Separation, Hazardous Content, and Subdivision				
	All suites are separated from the remainder of the building (including from other suites) by construction meeting the separation provisions for corridor construction (18.3.6.2-18.3.6.5 or 19.3.6.2-19.3.6.5). Existing approved barriers shall be allowed to continue to be used provided they limit the transfer of smoke. Intervening rooms have no hazardous areas and hazardous areas within suites comply with 18/19.2.5.7.1.3. Subdivision of suites shall be by noncombustible or limited-combustible construction. 18.2.5.7.1.2 through 18.2.5.7.1.4, 19.2.5.7.1.2, 19.2.5.7.1.3, 19.2.5.7.1.4				

ID PREFIX		MET	NOT MET	N/A	REMARKS
	Sleeping Suites Occupants shall have exit access to a corridor or direct access to a horizontal exit. Where ≥ 2 exits are required, one exit access door may be to a stairway, passageway or to the exterior. Suites shall be provided with constant staff supervision. Staff shall have direct visual supervision of patient sleeping rooms, from a constantly attended location or the room shall be provided with an automatic smoke detection system. Suites more than 1,000 ft² shall have 2 or more remote exits. One means of egress from the suite shall be to a corridor and one may be into an adjacent suite separated in accordance with corridor requirements. Suites shall not exceed the following size limitations: • 5,000 square feet if the suite is not fully smoke detected or fully sprinklered. • 7,500 square feet if the suite is either fully smoke detected and fully sprinklered and the sleeping rooms have direct supervision from a constantly attended location. Travel distance between any point in a suite to exit access shall not exceed 100 feet and distance to an exit shall not exceed 150 feet (200 feet if	MET		N/A	REMARKS
	building is fully sprinklered). 18.2.5.7.2, 19.2.5.7.2				
K257	Non-Sleeping Suites Occupants shall have exit access to a corridor or direct access to a horizontal exit. Where ≥ 2 exits are required, one exit access door may be to a stairway, passageway or to the exterior. Suites more than 2,500 ft² shall have 2 or more remote exits. One means of egress from the suite shall be to a corridor and one may be into an adjacent suite separated in accordance with corridor requirements. Suites shall not exceed 10,000 ft². Travel distance between any point in a suite to exit access shall not exceed 100 feet and distance to an exit shall not exceed 150 feet (200 feet if building is fully sprinklered). 18.2.5.7.3, 19.2.5.7.3				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K261	Travel Distance to Exits				
	Travel distance (excluding suites) to exits are measured in accordance with 7.6.				
	 From any point in the room or suite to exit less than or equal to 150 feet (less than or equal to 200 feet if the building is fully sprinklered). 				
	 Point in a room to room door less than or equal to 50 feet. 				
	18.2.6, 19.2.6				
K271	Discharge from Exits				
	Exit discharge is arranged in accordance with 7.7, provides a level walking surface meeting the provisions of 7.1.7 with respect to changes in elevation and shall be maintained free of obstructions. Additionally, the exit discharge shall be a hard packed all-weather travel surface. 18.2.7, 19.2.7				
K281	Illumination of Means of Egress				
	Illumination of means of egress, including exit discharge, is arranged in accordance with 7.8 and shall be either continuously in operation or capable of automatic operation without manual intervention. 18.2.8, 19.2.8				
K291	Emergency Lighting				
	Emergency lighting of at least 1-1/2 hour duration is provided automatically in accordance with 7.9.				
	18.2.9.1, 19.2.9.1				
K292	Life Support Means of Egress 2012 NEW (INDICATE N/A FOR EXISTING)				
	Buildings equipped with or requiring the use of life support systems (electro- mechanical or inhalation anesthetics) have illumination of means of egress, emergency lighting equipment, exit, and directional signs supplied by the life safety branch of the electrical system described in NFPA 99.				
	(Indicate N/A if life support equipment is for emergency purposes only.) 18.2.9.2, 18.2.10.5				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K293	Exit Signage 2012 EXISTING Exit and directional signs are displayed in accordance with 7.10 with continuous illumination also served by the emergency lighting system. 19.2.10.1 (Indicate N/A in one-story existing occupancies with less than 30 occupants where the line of exit travel is obvious.) 2012 NEW	IVILI	MET	IVA	INLIVIANNO
	Exit and directional signs are displayed in accordance with 7.10 with continuous illumination also served by the emergency lighting system. 18.2.10.1				
	SECTION 3 – PROTECTION				
K300	Protection – Other List in the REMARKS section any LSC Section 18.3 and 19.3 Protection requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567.				
K311	Vertical Openings – Enclosure 2012 EXISTING Stairways, elevator shafts, light and ventilation shafts, chutes, and other vertical openings between floors are enclosed with construction having a fire resistance rating of at least 1-hour. An atrium may be used in accordance with 8.6. 19.3.1.1 through 19.3.1.6 If all vertical openings are properly enclosed with construction providing at least a 2 hour fire resistance rating, also check this box. □				
	2012 NEW Stairways, elevator shafts, light and ventilation shafts, chutes, and other vertical openings between floors are enclosed with construction having a fire resistance rating of at least 2 hours connecting four or more stories. (1-hour for single story building and buildings up to three stories in height.) An atrium may be used in accordance with 8.6.7. 18.3.1 through 18.3.1.5				

ID PREFIX						MET	NOT MET	N/A	REMARKS
K321	Hazardous Areas – Enclosure 2012 EXISTING Hazardous areas are protected by a fire barrier having 1-hour fire resistance rating (with ¾ hour fire rated doors) or an automatic fire extinguishing system in accordance with 8.7.1 or 19.3.5.9. When the approved automatic fire extinguishing system option is used, the areas shall be separated from other spaces by smoke resisting partitions and doors in accordance with 8.4. Doors shall be self-closing or automatic-closing and permitted to have nonrated or field-applied protective plates that do not exceed 48 inches from the bottom of the door. Describe the floor and zone locations of hazardous areas that are deficient in REMARKS. 19.3.2.1, 19.3.5.9								
	Area Automatic Sprinkler Separation N/A								
	a. Boiler and Fuel-Fired Heater Rooms								
	b. Laundries (larger than 100 sq. ft.)				-				
	c. Repair, Maintenance, and Paint Shops								
	d. Soiled Linen Rooms (exceeding 64 gal.) e. Trash Collection Rooms (exceeding 64 gal.)				-				
	f. Combustible Storage Rooms/Spaces (over 50 sq. ft.)				-				

ID PREFIX						MET	NOT MET	N/A	REMARKS
K321	2012 NEW								
	shall be enclosed with a 1-hour fire door without windows (in accordant closing or automatic-closing in acc	Hazardous areas are protected in accordance with 18.3.2.1. The areas shall be enclosed with a 1-hour fire-rated barrier, with a ¾ hour fire-rated door without windows (in accordance with 8.7.1.1). Doors shall be self-closing or automatic-closing in accordance with 7.2.1.8. Hazardous areas are protected by a sprinkler system in accordance with 9.7, 18.3.2.1, and 8.4.							
	Describe the floor and zone locations of hazardous areas that are deficient in REMARKS.								
	18.3.2.1, 7.2.1.8, 8.4, 8.7, 9.7								
	Area	Automatic Sprinkler	Separation	N/A					
	a. Boiler and Fuel-Fired Heater Rooms								
	b. Laundries (larger than 100 sq. ft.)								
	c. Repair, Maintenance, and Paint Shops								
	d. Soiled Linen Rooms (exceeding 64 gal.)								
	e. Trash Collection Rooms (exceeding 64 gal.)								
	f. Combustible Storage Rooms/Spaces (over 50 and less than 100 sq. ft.)								
	g. Combustible Storage Rooms/Spaces (over 100 sq. ft.)								
	h. Laboratories (if classified as Severe Hazard - see K322)								

ID PREFIX		MET	NOT MET	N/A	REMARKS
ID PREFIX	Laboratories Laboratories employing quantities of flammable, combustible, or hazardous materials that are considered a severe hazard are protected by 1-hour fire resistance-rated separation, automatic sprinkler system, and are in accordance with 8.7 and with NFPA 99. Laboratories not considered a severe hazard are protected as hazardous areas (see K321). Laboratories using chemicals are in accordance with NFPA 45, Standard on Fire Protection for Laboratories Using Chemicals. Gas appliances are of appropriate design and installed in accordance with NFPA 54. Shutoff valves are marked to identify material they control. Devices requiring medical grade oxygen from the piped distribution system meet the requirements under 11.4.2.2 (NFPA 99).	MET	NOT MET	N/A	REMARKS
	18.3.2.2, 19.3.2.2, 8.7, 8.7.4.1 (LSC) 9.3.1.2, 11.4.3.2, 15.4 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K323	Anesthetizing Locations				
	Areas designated for administration of general anesthesia (i.e., inhalation anesthetics) are in accordance with 8.7 and NFPA 99.				
	Zone valves are: located immediately outside each life-support, critical care, and anesthetizing location of moderate sedation, deep sedation, or general anesthesia for medical gas or vacuum; readily accessible in an emergency; and arranged so shutting off any one anesthetizing location will not affect others.				
	Area alarm panels are provided to monitor all medical gas, medical- surgical vacuum, and piped WAGD systems. Panels are at locations that provide for surveillance, indicate medical gas pressure decreases of 20 percent and vacuum decreases of 12 inch gauge HgV, and provide visual and audible indication. Alarm sensors are installed either on the source side of individual room zone valve box assemblies or on the patient/use side of each of the individual zone box valve assemblies.				
	The EES critical branch supplies power for task illumination, fixed equipment, select receptacles, and select power circuits, and EES equipment system supplies power to ventilation system.				
	Heating, cooling, and ventilation are in accordance with ASHRAE 170. Medical supply and equipment manufacturer's instructions for use are considered before reducing humidity levels to those allowed by ASHRAE, per S&C 13-58.				
	18.3.2.3, 19.3.2.3 (LSC) 5.1.4.8.7, 5.1.4.8.7.2, 5.1.9.3, 5.1.9.3.4, 6.4.2.2.4.2 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K324	Cooking Facilities				
	Cooking equipment is protected in accordance with NFPA 96, Standard for Ventilation Control and Fire Protection of Commercial Cooking Operations, unless:				
	 residential cooking equipment (i.e., small appliances such as microwaves, hot plates, toasters) are used for food warming or limited cooking in accordance with 18.3.2.5.2, 19.3.2.5.2. 				
	 cooking facilities open to the corridor in smoke compartments with 30 or fewer patients comply with the conditions under 18.3.2.5.3, 19.3.2.5.3, or 				
	 cooking facilities in smoke compartments with 30 or fewer patients comply with conditions under 18.3.2.5.4, 19.3.2.5.4. 				
	Cooking facilities protected according to NFPA 96 per 9.2.3 are not required to be enclosed as hazardous areas, but shall not be open to the corridor.				
	18.3.2.5.1 through 18.3.2.5.4, 19.3.2.5.1 through 19.3.2.5.5, 9.2.3, TIA 12-2				
K325	Alcohol Based Hand Rub Dispenser (ABHR)				
	ABHRs are protected in accordance with 8.7.3.1, unless all conditions are met:				
	Corridor is at least 6 feet wide.				
	 Maximum individual dispenser capacity is 0.32 gallons (0.53 gallons in suites) of fluid and 18 ounces of Level 1 aerosols. 				
	Dispensers shall have a minimum of four foot horizontal spacing.				
	Not more than an aggregate of 10 gallons of fluid or 1135 ounces of aerosol are used in a single smoke compartment outside a storage cabinet, excluding one individual dispenser per room.				
	Storage in a single smoke compartment greater than 5 gallons complies with NFPA 30.				
	Dispensers are not installed within 1 inch of an ignition source.				
	Dispensers over carpeted floors are in sprinklered smoke compartments.				
	ABHR does not exceed 95 percent alcohol.				
	Operation of the dispenser shall comply with Section 18.3.2.6(11) or 19.3.2.6(11).				
	ABHR is protected against inappropriate access.				
	18.3.2.6, 19.3.2.6, 42 CFR Parts 403, 418, 460, 482, 483, and 485				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K331	Interior Wall and Ceiling Finish 2012 EXISTING Interior wall and ceiling finishes, including exposed interior surfaces of buildings such as fixed or movable walls, partitions, columns, and have a flame spread rating of Class A or Class B. The reduction in class of interior finish for a sprinkler system as prescribed in 10.2.8.1 is permitted. 10.2, 19.3.3.1, 19.3.3.2 Indicate flame spread rating(s).				
	Interior wall and ceiling finishes, including exposed interior surfaces of buildings such as fixed or movable walls, partitions and columns have a flame spread rating of Class A. The reduction in class of interior finish for a sprinkler system as prescribed in 10.2.8.1 is permitted. Individual rooms not exceeding four persons may have a Class A or B finish. Lower half of corridor walls, not exceeding 4 feet in height, may have a Class A or B flame spread rating. 10.2, 18.3.3.1, 18.3.3.2 Indicate flame spread rating(s).				
K332	Interior Floor Finish 2012 NEW (Indicate N/A for 2012 EXISTING) Interior finishes shall comply with 10.2. Floor finishes in exit enclosures and exit access corridors and spaces not separated by walls that resist the passage of smoke shall be Class I or II. 18.3.3.3.1, 18.3.3.3.2, 18.3.3.3.3, 10.2, 10.2.7.1, 10.2.7.2				
K341	Fire Alarm System – Installation A fire alarm system is installed with systems and components approved for the purpose in accordance with NFPA 70, National Electric Code, and NFPA 72, National Fire Alarm Code to provide effective warning of fire in any part of the building. In areas not continuously occupied, detection is installed at each fire alarm control unit. In new occupancy, detection is also installed at notification appliance circuit power extenders, and supervising station transmitting equipment. Fire alarm system wiring or other transmission paths are monitored for integrity. 18.3.4.1, 19.3.4.1, 9.6, 9.6.1.8				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K342	Fire Alarm System – Initiation				
	Initiation of the fire alarm system is by manual means and by any required sprinkler system alarm, detection device, or detection system. Manual alarm boxes are provided in the path of egress near each required exit. Manual alarm boxes in patient sleeping areas shall not be required at exits if manual alarm boxes are located at all nurse's stations or other continuously attended staff location, provided alarm boxes are visible, continuously accessible, and 200' travel distance is not exceeded. 18.3.4.2.1, 18.3.4.2.2, 19.3.4.2.1, 19.3.4.2.2, 9.6.2.5				
K343	Fire Alarm – Notification				
	2012 EXISTING				
	Positive alarm sequence in accordance with 9.6.3.4 are permitted in buildings protected throughout by a sprinkler system. Occupant notification is provided automatically in accordance with 9.6.3 by audible and visual signals.				
	In critical care areas, visual alarms are sufficient. The fire alarm system transmits the alarm automatically to notify emergency forces in the event of a fire.				
	19.3.4.3, 19.3.4.3.1, 19.3.4.3.2, 9.6.4, 9.7.1.1(1)				
	2012 NEW				
	Positive alarm sequence in accordance with 9.6.3.4 are permitted. Occupant notification is provided automatically in accordance with 9.6.3 by audible and visual signals.				
	In critical care areas, visual alarms are sufficient. The fire alarm system transmits the alarm automatically to notify emergency forces in the event of a fire.				
	Annunciation and annunciation zoning for fire alarm and sprinklers shall be provided by audible and visual indicators and zones shall not be larger than 22,500 square feet per zone.				
	18.3.4.3 through 18.3.4.3.3, 9.6.4				
K344	Fire Alarm – Control Functions				
	The fire alarm automatically activates required control functions and is provided with an alternative power supply in accordance with NFPA 72.				
	18.3.4.4, 19.3.4.4, 9.6.1, 9.6.5, NFPA 72				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K345	Fire Alarm System – Testing and Maintenance A fire alarm system is tested and maintained in accordance with an approved program complying with the requirements of NFPA 70, National Electric Code, and NFPA 72, National Fire Alarm and Signaling Code. Records of system acceptance, maintenance and testing are readily available.				
K346	9.6.1.3, 9.6.1.5, NFPA 70, NFPA 72 Fire Alarm – Out of Service Where required fire alarm system is out of services for more than 4 hours in a 24 hour period, the authority having jurisdiction shall be notified, and the building shall be evacuated or an approved fire watch shall be provided for all parties left unprotected by the shutdown until the fire alarm system has been returned to service. 9.6.1.6				
K347	Smoke Detection 2012 EXISTING Smoke detection systems are provided in spaces open to corridors as required by 19.3.6.1. 19.3.4.5.2				
	2012 NEW Smoke detection systems are provided in spaces open to corridors as required by 18.3.6.1 In nursing homes, an automatic smoke detection system is installed in the corridors of all smoke compartments containing resident sleeping rooms, unless the resident sleeping rooms have: • smoke detection, or • automatic door closing devices with integral smoke detectors on the room side that provide occupant notification. Such detectors are electrically interconnected to the fire alarm system. 18.3.4.5.2, 18.3.4.5.3				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K351	Sprinkler System – Installation				
	2012 EXISTING				
	Nursing homes, and hospitals where required by construction type, are protected throughout by an approved automatic sprinkler system in accordance with NFPA 13, Standard for the Installation of Sprinkler Systems.				
	In Type I and II construction, alternative protection measures are permitted to be substituted for sprinkler protection in specific areas where state or local regulations prohibit sprinklers.				
	In hospitals, sprinklers are not required in clothes closets of patient sleeping rooms where the area of the closet does not exceed 6 ft² and sprinkler coverage covers the closet footprint as required by NFPA 13, Standard for Installation of Sprinkler Systems.				
	19.3.5.1, 19.3.5.2, 19.3.5.3, 19.3.5.4, 19.3.5.5, 19.4.2, 19.3.5.10, 9.7, 9.7.1.1(1)				
	2012 NEW				
	Buildings are to be protected throughout by an approved automatic sprinkler system in accordance with NFPA 13, Standard for the Installation of Sprinkler Systems.				
	In Type I and II construction, alternative protection measures are permitted to be substituted for sprinkler protection in specific areas where State and local regulations prohibit sprinklers.				
	Listed quick-response or listed residential sprinklers are used throughout smoke compartments with patient sleeping rooms.				
	In hospitals, sprinklers are not required in clothes closets of patient sleeping rooms where the area of the closet does not exceed 6 ft² and sprinkler coverage covers the closet footprint as required by NFPA 13, Standard for Installation of Sprinkler Systems.				
	18.3.5.1, 18.3.5.4, 18.3.5.5, 18.3.5.6, 9.7, 9.7.1.1(1), 18.3.5.10				
K352	Sprinkler System – Supervisory Signals				
	Automatic sprinkler system supervisory attachments are installed and monitored for integrity in accordance with NFPA 72, <i>National Fire Alarm and Signaling Code</i> , and provide a signal that sounds and is displayed at a continuously attended location or approved remote facility when sprinkler operation is impaired.				
	9.7.2.1, NFPA 72				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K353	Sprinkler System – Maintenance and Testing Automatic sprinkler and standpipe systems are inspected, tested, and maintained in accordance with NFPA 25, Standard for the Inspection, Testing, and Maintaining of Water-based Fire Protection Systems. Records of system design, maintenance, inspection and testing are maintained in a secure location and readily available. a) Date sprinkler system last checked. b) Who provided system test. c) Water system supply source. Provide in REMARKS information on coverage for any non-required or partial automatic sprinkler system. 9.7.5, 9.7.7, 9.7.8, and NFPA 25				
K354	Sprinkler System – Out of Service Where the sprinkler system is impaired, the extent and duration of the impairment has been determined, areas or buildings involved are inspected and risks are determined, recommendations are submitted to management or designated representative, and the fire department and other authorities having jurisdiction have been notified. Where the sprinkler system is out of service for more than 10 hours in a 24 hour period, the building or portion of the building affected are evacuated or an approved fire watch is provided until the sprinkler system has been returned to service. 18.3.5.1, 19.3.5.1, 9.7.5, 15.5.2 (NFPA 25) Portable Fire Extinguishers Portable fire extinguishers are selected, installed, inspected, and				
K361	maintained in accordance with NFPA 10, <i>Standard for Portable Fire Extinguishers</i> . 18.3.5.12, 19.3.5.12, NFPA 10 Corridors – Areas Open to Corridor Spaces (other than patient sleeping rooms, treatment rooms and hazardous areas), waiting areas, nurse's stations, gift shops, and cooking facilities, open to the corridor are in accordance with the criteria under 18.3.6.1 and 19.3.6.1. 18.3.6.1, 19.3.6.1				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K362	Corridors – Construction of Walls 2012 EXISTING Corridors are separated from use areas by walls constructed with at least ½ hour fire resistance rating. In fully sprinklered smoke compartments, partitions are only required to resist the transfer of smoke. In nonsprinklered buildings, walls extend to the underside of the floor or roof deck above the ceiling. Corridor walls may terminate at the underside of ceilings where specifically permitted by Code. Fixed fire window assemblies in corridor walls are in accordance with Section 8.3, but in sprinklered compartments there are no restrictions in area or fire resistance of glass or frames. If the walls have a fire resistance rating, give the rating if the walls terminate at the underside of the ceiling, give brief description in	=	MET		
	REMARKS, describing the ceiling throughout the floor area. 19.3.6.2, 19.3.6.2.7 2012 NEW Corridor walls shall form a barrier to limit the transfer of smoke. Such walls shall be permitted to terminate at the ceiling where the ceiling is constructed to limit the transfer of smoke. No fire resistance rating is required for the corridor walls. 18.3.6.2				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K363	Corridor – Doors 2012 EXISTING Doors protecting corridor openings in other than required enclosures of vertical openings, exits, or hazardous areas resist the passage of smoke and are made of 1¾ inch solid-bonded core wood or other material capable of resisting fire for at least 20 minutes. Doors in fully sprinklered smoke compartments are only required to resist the passage of smoke. Corridor doors and doors to rooms containing flammable or combustible materials have positive latching hardware. Roller latches are prohibited by CMS regulation. These requirements do not apply to auxiliary spaces that do not contain flammable or combustible material. Powered doors complying with 7.2.1.9 are permissible if provided with a device capable of keeping the door closed when a force of 5lbf is applied, whether or not power is applied. Clearance between bottom of door and floor covering is not exceeding 1 inch. There is no impediment to the closing of the doors. Hold open devices that release when the door is pushed or pulled are permitted. Nonrated protective plates of unlimited height are permitted. Dutch doors meeting 19.3.6.3.6 are permitted. Door frames shall be labeled and made of steel or other materials in compliance with 8.3, unless the smoke compartment is sprinklered. Fixed fire window assemblies are allowed per 8.3. In sprinklered compartments there are no restrictions in area or fire resistance of glass or frames in window assemblies. 19.3.6.3, 42 CFR Parts 403, 418, 460, 482, 483, and 485 Show in REMARKS details of doors such as fire protection ratings, automatics closing devices, etc.				
	Doors protecting corridor openings shall be constructed to resist the passage of smoke. Corridor doors and doors to rooms containing flammable or combustible materials have self-latching and positive latching hardware. Roller latches are prohibited by CMS regulation. These requirements do not apply to auxiliary spaces that do not contain flammable or combustible material. Powered doors complying with 7.2.1.9 are permissible if provided with a device capable of keeping the door closed when a force of 5lbf is applied, whether or not power is applied. Clearance between bottom of door and floor covering is not exceeding 1 inch. There is no impediment to the closing of the doors. Hold open devices that release when the door is pushed or pulled are permitted. Nonrated protective plates of unlimited height are permitted. Dutch doors meeting 18.3.6.3.6 are permitted. 18.3.6.3, 42 CFR Parts 403, 418, 460, 482, 483, and 485 Show in REMARKS details of doors such as fire protection ratings, automatic closing devices, etc.				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K364	Corridor – Openings				
	Transfer grilles are not used in corridor walls or doors. Auxiliary spaces that do not contain flammable or combustible materials are permitted to have louvers or be undercut.				
	In other than smoke compartments containing patient sleeping rooms, miscellaneous openings are permitted in vision panels or doors, provided the openings per room do not exceed 20 in² and are at or below half the distance from floor to ceiling. In sprinklered rooms, the openings per room do not exceed 80 in².				
	Vision panels in corridor walls or doors shall be fixed window assemblies in approved frames. (In fully sprinklered smoke compartments, there are no restrictions in the area and fire resistance of glass and frames.) 18.3.6.5.1, 19.3.6.5.2, 8.3				
K371	Subdivision of Building Spaces – Smoke Compartments				
	2012 EXISTING				
	Smoke barriers shall be provided to form at least two smoke compartments on every sleeping floor with a 30 or more patient bed capacity. Size of compartments cannot exceed 22,500 square feet or a 200-foot travel distance from any point in the compartment to a door in the smoke barrier.				
	19.3.7.1, 19.3.7.2				
	Detail in REMARKS zone dimensions including length of zones and deadend corridors.				
	2012 NEW				
	Smoke barriers shall be provided to form at least two smoke compartments on every floor used by inpatients for sleeping or treatment, and on every floor with an occupant load of 50 or more persons, regardless of use.				
	Size of compartments cannot exceed 22,500 square feet or a 200-foot travel distance from any point in the compartment to a door in the smoke barrier.				
	Smoke subdivision requirements do not apply to any of the stories or areas described in 18.3.7.2.				
	18.3.7.1, 18.3.7.2				
	Detail in REMARKS zone dimensions including length of zones and deadend corridors.				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K372	Subdivision of Building Spaces – Smoke Barrier Construction		IVILI		
	2012 EXISTING				
	Smoke barriers shall be constructed to a ½ hour fire resistance rating per 8.5. Smoke barriers shall be permitted to terminate at an atrium wall. Smoke dampers are not required in duct penetrations in fully ducted HVAC systems where an approved sprinkler system is installed for smoke compartments adjacent to the smoke barrier.				
	19.3.7.3, 8.6.7.1(1)				
	Describe any mechanical smoke control system in REMARKS.				
	2012 NEW				
	Smoke barriers shall be constructed to provide at least a 1-hour fire resistance rating and constructed in accordance with 8.5. Smoke barriers shall be permitted to terminate at an atrium wall. Smoke dampers are not required in duct penetrations of fully ducted HVAC systems.				
	18.3.7.3, 18.3.7.4, 18.3.7.5, 8.3				
	Describe any mechanical smoke control system in REMARKS.				
K373	Subdivision of Building Spaces – Accumulation Space Space shall be provided on each side of smoke barriers to adequately accommodate the total number of occupants in adjoining compartments. 18.3.7.5.1, 18.3.7.5.2, 19.3.7.5.1, 19.3.7.5.2				
K374	Subdivision of Building Spaces – Smoke Barrier Doors 2012 EXISTING Doors in smoke barriers are 1¾-inch thick solid bonded wood-core doors or of construction that resists fire for 20 minutes. Nonrated protective plates of unlimited height are permitted. Doors are permitted to have fixed fire window assemblies per 8.5. Doors are self-closing or automatic-closing, do not require latching, and are not required to swing in the direction of egress travel. Door opening provides a minimum clear width of 32 in for swinging or horizontal doors. 19.3.7.6, 19.3.7.8, 19.3.7.9				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K374	2012 NEW				
	Doors in smoke barriers have at least a 20-minute fire protection rating or are at least 1¾-inch thick solid bonded core wood.				
	Required clear widths are provided per 18.3.7.6(4) and (5).				
	Nonrated protective plates of unlimited height are permitted. Horizontal- sliding doors comply with 7.2.1.14. Swinging doors shall be arranged so that each door swings in an opposite direction.				
	Doors shall be self-closing and rabbets, bevels, or astragals are required at the meeting edges. Positive latching is not required.				
	18.3.7.6, 18.3.7.7, 18.3.7.8				
K379	Smoke Barrier Door Glazing				
	2012 EXISTING				
	Openings in smoke barrier doors shall be fire-rated glazing or wired glass panels in steel frames.				
	19.3.7.6, 19.3.7.6.2, 8.5				
	2012 NEW				
	Windows in smoke barrier doors shall be installed in each cross corridor swinging or horizontal-sliding door protected by fire-rated glazing or by wired glass panels in approved frames.				
	18.3.7.9				
K381	Sleeping Room Outside Windows and Doors				
	Every patient sleeping room has an outside window or outside door. In new occupancies, sill height does not exceed 36 inches above the floor. Windows in atrium walls are considered outside windows. Newborn nurseries and rooms intended for occupancy less than 24 hours have no outside window or door requirements. Window sills in special nursing care areas (e.g., ICU, CCU, hemodialysis, neonatal) do not exceed 60 inches above the floor.				
	42 CFR 403, 418, 460, 482, 483, and 485				
	SECTION 4 – SPECIAL PROVISIONS				
K400	Special Provisions – Other				
	List in the REMARKS section any LSC Section 18.4 and 19.4 Special Provisions requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or				
	NFPA standard citation, should be included on Form CMS-2567.				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K421	High-Rise Buildings 2012 EXISTING High-rise buildings are protected throughout by an approved, supervised automatic sprinkler system in accordance with Section 9.7 within 12 years of LSC final rule effective date. 19.4.2				
	2012 NEW High-rise buildings comply with section 11.8. 18.4.2				
	SECTION 5 – BUILDING SERVICES				
K500	Building Services – Other List in the REMARKS section any LSC Section 18.5 and 19.5 Building Services requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567.				
K511	Utilities – Gas and Electric				
	Equipment using gas or related gas piping complies with NFPA 54, <i>National Fuel Gas Code</i> , electrical wiring and equipment complies with NFPA 70, <i>National Electric Code</i> . Existing installations can continue in service provided no hazard to life.				
	18.5.1.1, 19.5.1.1, 9.1.1, 9.1.2				
K521	HVAC Heating, ventilation, and air conditioning shall comply with 9.2 and shall be installed in accordance with the manufacturer's specifications. 18.5.2.1, 19.5.2.1, 9.2				
K522	HVAC – Any Heating Device				
	Any heating device, other than a central heating plant, is designed and installed so combustible materials cannot be ignited by device, and has a safety feature to stop fuel and shut down equipment if there is excessive temperature or ignition failure. If fuel fired, the device also:				
	is chimney or vent connected.				
	takes air for combustion from outside.				
	 provides for a combustion system separate from occupied area atmosphere. 18.5.2.2, 19.5.2.2 				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K523	HVAC - Suspended Unit Heaters				
	Suspended unit heaters are permitted provided the following are met:				
	Not located in means of egress or in patient rooms.				
	Located high enough to be out of reach of people in the area.				
	 Has a safety feature to stop fuel and shut down equipment if there is excessive temperature or ignition failure. 				
	18.5.2.3(1), 19.5.2.3(1)				
K524	HVAC - Direct-Vent Gas Fireplaces				
	Direct-vent gas fireplaces, as defined in NFPA 54, inside of all smoke compartments containing patient sleeping areas comply with the requirements of 18.5.2.3(2), 19.5.2.3(2). 18.5.2.3(2), 19.5.2.3(2), NFPA 54				
K525	HVAC - Solid Fuel-Burning Fireplaces				
	Solid fuel-burning fireplaces are permitted in other than patient sleeping areas provided:				
	Areas are separated by 1-hour fire resistance construction.				
	• Fireplace complies with 9.2.2.				
	 Fireplace enclosure resists breakage up to 650°F and has heat- tempered glass. 				
	 Room has supervised CO detection per 9.8. 				
	18.5.2.3(3) and 19.5.2.3(3)				
K531	Elevators				
	2012 EXISTING				
	Elevators comply with the provision of 9.4. Elevators are inspected and tested as specified in ASME A17.1, Safety Code for Elevators and Escalators. Firefighter's Service is operated monthly with a written record.				
	Existing elevators conform to ASME/ANSI A17.3, <i>Safety Code for Existing Elevators and Escalators</i> . All existing elevators, having a travel distance of 25 feet or more above or below the level that best serves the needs of emergency personnel for firefighting purposes, conform with Firefighter's Service Requirements of ASME/ANSI A17.3. (Includes firefighter's service Phase I key recall and smoke detector automatic recall, firefighter's service Phase II emergency in-car key operation, machine room smoke detectors, and elevator lobby smoke detectors.)				
	19.5.3, 9.4.2, 9.4.3				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K531	2012 NEW Elevators comply with the provision of 9.4. Elevators are inspected and tested as specified in ASME A17.1, <i>Safety Code for Elevators and Escalators</i> . Firefighter's Service is operated monthly with a written record. New elevators conform to ASME/ANSI A17.1, <i>Safety Code for Elevators and Escalators</i> , including Firefighter's Service Requirements. (Includes firefighter's Phase I key recall and smoke detector automatic recall, firefighter's service Phase II emergency in-car key operation, machine room smoke detectors, and elevator lobby smoke detectors.) 18.5.3, 9.4.2, 9.4.3				
K532	Escalators, Dumbwaiters, and Moving Walks 2012 EXISTING Escalators, dumbwaiters, and moving walks comply with the provisions of 9.4. All existing escalators, dumbwaiters, and moving walks conform to the requirements of ASME/ANSI A17.3, Safety Code for Existing Elevators and Escalators. (Includes escalator emergency stop buttons and automatic skirt obstruction stop. For power dumbwaiters, includes hoistway door locking to keep doors closed except for floor where car is being loaded or unloaded.) 19.5.3, 9.4.2.2				
	2012 NEW Escalators, dumbwaiters, and moving walks comply with the provisions of 9.4. 18.5.3, 9.4.2.2				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K541	Rubbish Chutes, Incinerators, and Laundry Chutes				
	2012 EXISTING				
	(1) Any existing linen and trash chute, including pneumatic rubbish and linen systems, that opens directly onto any corridor shall be sealed by fire resistive construction to prevent further use or shall be provided with a fire door assembly having a fire protection rating of 1-hour. All new chutes shall comply with 9.5.				
	(2) Any rubbish chute or linen chute, including pneumatic rubbish and linen systems, shall be provided with automatic extinguishing protection in accordance with 9.7.				
	(3) Any trash chute shall discharge into a trash collection room used for no other purpose and protected in accordance with 8.4. (Existing laundry chutes permitted to discharge into same room are protected by automatic sprinklers in accordance with 19.3.5.9 or 19.3.5.7.)				
	(4) Existing fuel-fed incinerators shall be sealed by fire resistive construction to prevent further use.				
	19.5.4, 9.5, 8.4, NFPA 82				
	2012 NEW				
	Rubbish chutes, incinerators, and laundry chutes shall comply with the provisions of Section 9.5, unless otherwise specified in 18.5.4.2.				
	The fire resistance rating of chute charging room shall not be required to exceed 1-hour.				
	 Any rubbish chute or linen chute shall be provided with automatic extinguishing protection in accordance with Section 9.7. 				
	 Chutes shall discharge into a trash collection room used for no other purpose and shall be protected in accordance with 8.7. 18.5.4.2, 8.7, 9.5, 9.7, NFPA 82 				
	SECTION 6 – RESERVED				
	SECTION 6 - RESERVED SECTION 7 - OPERATING FEATURES				
1/700					
K700	Operating Features – Other List in the REMARKS section any LSC Section 18.7 and 19.7 Operating				
	Features requirements that are not addressed by the provided K-tags, but are deficient. This information, along with the applicable Life Safety Code or				
	NFPA standard citation, should be included in Form CMS-2567.				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K711	Evacuation and Relocation Plan				
	There is a written plan for the protection of all patients and for their evacuation in the event of an emergency.				
	Employees are periodically instructed and kept informed with their duties under the plan, and a copy of the plan is readily available with telephone operator or with security. The plan addresses the basic response required of staff per 18/19.7.2.1.2 and provides for all of the fire safety plan components per 18/19.7.2.2.				
	18.7.1.1 through 18.7.1.3, 18.7.2.1.2, 18.7.2.2, 18.7.2.3, 19.7.1.1 through 19.7.1.3, 19.7.2.1.2, 19.7.2.2, 19.7.2.3				
K712	Fire Drills				
K712	Fire Drills Fire drills include the transmission of a fire alarm signal and simulation of emergency fire conditions. Fire drills are held at expected and unexpected times under varying conditions, at least quarterly on each shift. The staff is familiar with procedures and is aware that drills are part of established routine. Where drills are conducted between 9:00 PM and 6:00 AM, a coded announcement may be used instead of audible alarms. 18.7.1.4 through 18.7.1.7, 19.7.1.4 through 19.7.1.7				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K741	 Smoking Regulations Smoking regulations shall be adopted and shall include not less than the following provisions: Smoking shall be prohibited in any room, ward, or compartment where flammable liquids, combustible gases, or oxygen is used or stored and in any other hazardous location, and such area shall be posted with signs that read NO SMOKING or shall be posted with the international symbol for no smoking. In health care occupancies where smoking is prohibited and signs are prominently placed at all major entrances, secondary signs with language that prohibits smoking shall not be required. Smoking by patients classified as not responsible shall be prohibited. The requirement of 18.7.4(3) shall not apply where the patient is under direct supervision. Ashtrays of noncombustible material and safe design shall be provided in all areas where smoking is permitted. Metal containers with self-closing cover devices into which ashtrays can be emptied shall be readily available to all areas where smoking is permitted. 		MEI		
K751	Draperies, Curtains, and Loosely Hanging Fabrics Draperies, curtains including cubicle curtains and loosely hanging fabric or films shall be in accordance with 10.3.1. Excluding curtains and draperies: at showers and baths; on windows in patient sleeping room located in sprinklered compartments; and in non-patient sleeping rooms in sprinklered compartments where individual drapery or curtain panels do not exceed 48 square feet or total area does not exceed 20 percent of the wall. 18.7.5.1, 18.3.5.11, 19.7.5.1, 19.3.5.11, 10.3.1				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K752	Upholstered Furniture and Mattresses				
	Newly introduced upholstered furniture meets Class I or char length, and heat release criteria in accordance with 10.3.2.1 and 10.3.3, unless the building is fully sprinklered.				
	Newly introduced mattresses shall meet char length and heat release criteria in accordance with 10.3.2.2 and 10.3.4, unless the building is fully sprinklered.				
	Upholstered furniture and mattresses belonging to nursing home residents do not have to meet these requirements as all nursing homes are required to be fully sprinklered.				
	Newly introduced upholstered furniture and mattresses means purchased on or after the LSC final rule effective date.				
	18.7.5.2, 18.7.5.4, 19.7.5.2, 19.7.5.4				
K753	Combustible Decorations				
	Combustible decorations shall be prohibited unless one of the following is met:				
	Flame retardant or treated with approved fire-retardant coating that is listed and labeled for product.				
	Decorations meet NFPA 701.				
	 Decorations exhibit heat release less than 100 kilowatts in accordance with NFPA 289. 				
	• Decorations, such as photographs, paintings and other art are attached to the walls, ceilings and non-fire-rated doors in accordance with 18.7.5.6(4) or 19.7.5.6(4).				
	 The decorations in existing occupancies are in such limited quantities that a hazard of fire development or spread is not present. 18.7.5.6, 19.7.5.6 				
K761	Maintenance, Inspection & Testing - Doors				
	Fire doors assemblies are inspected and tested annually in accordance with NFPA 80 Standard for Fire Doors and Other Opening Protectives.				
	Fire doors that are not located in required fire barriers, including corridor doors to patient rooms and smoke barrier doors, are routinely inspected as part of the facility maintenance program.				
	Individuals performing the door inspection and testing have an understanding of the operating components of the doors. Written records of inspection and testing are maintained and are available for review.				
	18.7.6, 19.7.6, 8.3.3.1 (LSC), 5.2, 5.2.3 (NFPA 80)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K754	Soiled Linen and Trash Containers				
	Soiled linen or trash collection receptacles shall not exceed 32 gallons in capacity. The average density of container capacity in a room or space shall not exceed 0.5 gallons/square feet. A total container capacity of 32 gallons shall not be exceeded within any 64 square feet area. Mobile soiled linen or trash collection receptacles with capacities greater than 32 gallons shall be located in a room protected as a hazardous area when not attended. Containers used solely for recycling are permitted to be excluded from the				
	above requirements where each container is ≤ 96 gal. unless attended, and containers for combustibles are labeled and listed as meeting FM Approval Standard 6921 or equivalent.				
	18.7.5.7, 19.7.5.7				
K771	Engineer Smoke Control Systems 2012 EXISTING When installed, engineered smoke control systems are tested in accordance with established engineering principles. Test documentation is maintained on the premises.				
	19.7.7				
	2012 NEW When installed, engineered smoke control systems are tested in accordance with NFPA 92, <i>Standard for Smoke Control Systems</i> . Test documentation is maintained on the premises. 18.7.7				
K781	Portable Space Heaters Portable space heating devices shall be prohibited in all health care occupancies. Unless used in nonsleeping staff and employee areas where the heating elements do not exceed 212 degrees Fahrenheit (100 degrees Celsius). 18.7.8, 19.7.8				
K791	Construction, Repair, and Improvement Operations				
	Construction, repair, and improvement operations shall comply with 4.6.10. Any means of egress in any area undergoing construction, repair, or improvements shall be inspected daily to ensure its ability to be used instantly in case of emergency and compliance with NFPA 241.				
	18.7.9, 19.7.9, 4.6.10, 7.1.10.1				

ID PREFIX		MET	NOT MET	N/A	REMARKS
TREID	PART II – HEALTH CARE FACILITIES CODE REQUIREMENTS		IVIL		
K900	Health Care Facilities Code - Other List in the REMARKS section any NFPA 99 requirements (excluding Chapter 7, 8, 12, and 13) that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Health Care Facilities Code or NFPA standard citation, should be included on Form CMS-2567.				
K901	Fundamentals – Building System Categories Building systems are designed to meet Category 1 through 4 requirements as detailed in NFPA 99. Categories are determined by a formal and documented risk assessment procedure performed by qualified personnel. Chapter 4 (NFPA 99)				
K902	Gas and Vacuum Piped Systems – Other List in the REMARKS section any NFPA 99 Chapter 5 Gas and Vacuum Systems requirements that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567. Chapter 5 (NFPA 99)				
K903	Gas and Vacuum Piped Systems – Categories Medical gas, medical air, surgical vacuum, WAGD, and air supply systems are designated: □ Category 1. Systems in which failure is likely to cause major injury or death. □ Category 2. Systems in which failure is likely to cause minor injury. □ Category 3. Systems in which failure is not likely to cause injury, but can cause discomfort. Deep sedation and general anesthesia are not to be administered using a Category 3 medical gas system. 5.1.1.1, 5.2.1, 5.3.1.1, 5.3.1.5 (NFPA 99)				
K904	Gas and Vacuum Piped Systems – Warning Systems All master, area, and local alarm systems used for medical gas and vacuum systems comply with appropriate Category warning system requirements, as applicable. 5.1.9, 5.2.9, 5.3.6.2.2 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K905	Gas and Vacuum Piped Systems – Central Supply System Identification and Labeling Containers, cylinders and tanks are designed, fabricated, tested, and marked in accordance with 5.1.3.1.1 through 5.1.3.1.7. Locations				
	containing only oxygen or medical air have doors labeled with "Medical Gases, NO Smoking or Open Flame". Locations containing other gases have doors labeled "Positive Pressure Gases, NO Smoking or Open Flame, Room May Have Insufficient Oxygen, Open Door and Allow Room to Ventilate Before Opening."				
14000	5.1.3.1, 5.2.3.1, 5.3.10 (NFPA 99)				
K906	Gas and Vacuum Piped Systems – Central Supply System Operations Adaptors or conversion fittings are prohibited. Cylinders are handled in accordance with 11.6.2. Only cylinders, reusable shipping containers, and their accessories are stored in rooms containing central supply systems or cylinders. No flammable materials are stored with cylinders. Cryogenic liquid storage units intended to supply the facility are not used to transfill. Cylinders are kept away from sources of heat. Valve protection caps are secured in place, if supplied, unless cylinder is in use. Cylinders are not stored in tightly closed spaces. Cylinders in use and storage are prevented from exceeding 130°F, and nitrous oxide and carbon dioxide cylinders are prevented from reaching temperatures lower than manufacture recommendations or 20°F. Full or empty cylinders, when not connected, are stored in locations complying with 5.1.3.3.2 through 5.1.3.3.3, and are not stored in enclosures containing motor-driven machinery, unless for instrument air reserve headers. 5.1.3.2, 5.1.3.3.17, 5.1.3.3.1.8, 5.1.3.3.4, 5.2.3.2, 5.2.3.3, 5.3.6.20.4, 5.6.20.5, 5.3.6.20.7, 5.3.6.20.8, 5.3.6.20.9 (NFPA 99)				
K907	Gas and Vacuum Piped Systems – Maintenance Program				
	Medical gas, vacuum, WAGD, or support gas systems have documented maintenance programs. The program includes an inventory of all source systems, control valves, alarms, manufactured assemblies, and outlets. Inspection and maintenance schedules are established through risk assessment considering manufacturer recommendations. Inspection procedures and testing methods are established through risk assessment. Persons maintaining systems are qualified as demonstrated by training and certification or credentialing to the requirements of AASE 6030 or 6040. 5.1.14.2.1, 5.1.14.2.2, 5.1.15, 5.2.14, 5.3.13.4.2 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K908	Gas and Vacuum Piped Systems – Inspection and Testing Operations				
	The gas and vacuum systems are inspected and tested as part of a maintenance program and include the required elements. Records of the inspections and testing are maintained as required. 5.1.14.2.3, B.5.2, 5.2.13, 5.3.13, 5.3.13.4 (NFPA 99)				
K909	Gas and Vacuum Piped Systems – Information and Warning Signs				
	Piping is labeled by stencil or adhesive markers identifying the gas or vacuum system, including the name of system or chemical symbol, color code (Table 5.1.11), and operating pressure if other than standard. Labels are at intervals not more than 20 feet, are in every room, at both sides of wall penetrations, and on every story traversed by riser. Piping is not painted. Shutoff valves are identified with the name or chemical symbol of the gas or vacuum system, room or area served, and caution to not use the valve except in emergency. 5.1.14.3, 5.1.11.1, 5.1.11.2, 5.2.11, 5.3.13.3, 5.3.11 (NFPA 99)				
K910	Gas and Vacuum Piped Systems – Modifications				
	Whenever modifications are made that breach the pipeline, any necessary installer and verification test specified in 5.1.2 is conducted on the downstream portion of the medical gas piping system. Permanent records of all tests required by system verification tests are maintained. 5.1.14.4.1, 5.1.14.4.6, 5.2.13, 5.3.13.4.3 (NFPA 99)				
K911	Electrical Systems – Other				
	List in the REMARKS section any NFPA 99 Chapter 6 Electrical Systems requirements that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567. Chapter 6 (NFPA 99)				
K912	Electrical Systems – Receptacles				
	Power receptacles have at least one, separate, highly dependable grounding pole capable of maintaining low-contact resistance with its mating plug. In pediatric locations, receptacles in patient rooms, bathrooms, play rooms, and activity rooms, other than nurseries, are listed tamper-resistant or employ a listed cover.				
	If used in patient care room, ground-fault circuit interrupters (GFCI) are listed.				
	6.3.2.2.6.2 (F), 6.3.2.2.4.2 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K913	Electrical Systems – Wet Procedure Locations Operating rooms are considered wet procedure locations, unless otherwise determined by a risk assessment conducted by the facility governing body. Operating rooms defined as wet locations are protected by either isolated power or ground-fault circuit interrupters. A written record of the risk assessment is maintained and available for inspection. 6.3.2.2.8.4, 6.3.2.2.8.7, 6.4.4.2				
K914	Electrical Systems – Maintenance and Testing Hospital-grade receptacles at patient bed locations and where deep sedation or general anesthesia is administered, are tested after initial installation, replacement or servicing. Additional testing is performed at intervals defined by documented performance data. Receptacles not listed as hospital-grade at these locations are tested at intervals not exceeding 12 months. Line isolation monitors (LIM), if installed, are tested at intervals of ≤ 1 month by actuating the LIM test switch per 6.3.2.6.3.6, which activates both visual and audible alarm. For LIM circuits with automated self-testing, this manual test is performed at intervals ≤ 12 months. LIM circuits are tested per 6.3.3.3.2 after any repair or renovation to the electric distribution system. Records are maintained of required tests and associated repairs or modifications, containing date, room or area tested, and results. 6.3.4 (NFPA 99)				
K915	Electrical Systems – Essential Electric System Categories ☐ Critical care rooms (Category 1) in which electrical system failure is likely to cause major injury or death of patients, including all rooms where electric life support equipment is required, are served by a Type 1 EES. ☐ General care rooms (Category 2) in which electrical system failure is likely to cause minor injury to patients (Category 2) are served by a Type 1 or Type 2 EES. ☐ Basic care rooms (Category 3) in which electrical system failure is not likely to cause injury to patients and rooms other than patient care rooms are not required to be served by an EES. Type 3 EES life safety branch has an alternate source of power that will be effective for 1 1/2 hours. 3.3.138, 6.3.2.2.10, 6.6.2.2.2, 6.6.3.1.1 (NFPA 99), TIA 12-3				

	MET	NOT MET	N/A	REMARKS
Electrical Systems – Essential Electric System Alarm Annunciator		IVIEI		
A remote annunciator that is storage battery powered is provided to operate outside of the generating room in a location readily observed by operating personnel. The annunciator is hard-wired to indicate alarm conditions of the emergency power source. A centralized computer system (e.g., building information system) is not to be substituted for the alarm annunciator.				
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Electrical Systems – Essential Electric System Receptacles Electrical receptacles or cover plates supplied from the life safety and critical branches have a distinctive color or marking. 6.4.2.2.6, 6.5.2.2.4.2, 6.6.2.2.3.2 (NFPA 99)				
Electrical Systems – Essential Electric System Maintenance and Testing				
The generator or other alternate power source and associated equipment is capable of supplying service within 10 seconds. If the 10-second criterion is not met during the monthly test, a process shall be provided to annually confirm this capability for the life safety and critical branches. Maintenance and testing of the generator and transfer switches are performed in accordance with NFPA 110.				
Generator sets are inspected weekly, exercised under load 30 minutes 12 times a year in 20-40 day intervals, and exercised once every 36 months for 4 continuous hours. Scheduled test under load conditions include a complete simulated cold start and automatic or manual transfer of all EES loads, and are conducted by competent personnel. Maintenance and testing of stored energy power sources (Type 3 EES) are in accordance with NFPA 111. Main and feeder circuit breakers are inspected annually, and a program for periodically exercising the components is established according to manufacturer requirements. Written records of maintenance and testing are maintained and readily available. EES electrical panels and circuits are marked, readily identifiable, and separate from normal power circuits. Minimizing the possibility of damage of the emergency power source is a design consideration for new installations. 6.4.4, 6.5.4, 6.6.4 (NFPA 99), NFPA 110, NFPA 111, 700.10 (NFPA 70)				
	outside of the generating room in a location readily observed by operating personnel. The annunciator is hard-wired to indicate alarm conditions of the emergency power source. A centralized computer system (e.g., building information system) is not to be substituted for the alarm annunciator. 6.4.1.1.17, 6.4.1.1.17.5 (NFPA 99) Electrical Systems – Essential Electric System Receptacles Electrical receptacles or cover plates supplied from the life safety and critical branches have a distinctive color or marking. 6.4.2.2.6, 6.5.2.2.4.2, 6.6.2.2.3.2 (NFPA 99) Electrical Systems – Essential Electric System Maintenance and Testing The generator or other alternate power source and associated equipment is capable of supplying service within 10 seconds. If the 10-second criterion is not met during the monthly test, a process shall be provided to annually confirm this capability for the life safety and critical branches. 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ID PREFIX		MET	NOT MET	N/A	REMARKS
K919	Electrical Equipment – Other List in the REMARKS section any NFPA 99 Chapter 10, Electrical Equipment, requirements that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567. Chapter 10 (NFPA 99)				
K920	Electrical Equipment – Power Cords and Extension Cords Power strips in a patient care vicinity are only used for components of movable patient-care-related electrical equipment (PCREE) assembles that have been assembled by qualified personnel and meet the conditions of 10.2.3.6. Power strips in the patient care vicinity may not be used for non-PCREE (e.g., personal electronics), except in long-term care resident rooms that do not use PCREE. Power strips for PCREE meet UL 1363A or UL 60601-1. Power strips for non-PCREE in the patient care rooms (outside of vicinity) meet UL 1363. In non-patient care rooms, power strips meet other UL standards. All power strips are used with general precautions. Extension cords are not used as a substitute for fixed wiring of a structure. Extension cords used temporarily are removed immediately upon completion of the purpose for which it was installed and meets the conditions of 10.2.4. 10.2.3.6 (NFPA 99), 10.2.4 (NFPA 99), 400-8 (NFPA 70), 590.3(D) (NFPA 70), TIA 12-5				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K921	Electrical Equipment – Testing and Maintenance Requirements				
	The physical integrity, resistance, leakage current, and touch current tests for fixed and portable patient-care related electrical equipment (PCREE) is performed as required in 10.3. Testing intervals are established with policies and protocols. All PCREE used in patient care rooms is tested in accordance with 10.3.5.4 or 10.3.6 before being put into service and after any repair or modification. Any system consisting of several electrical appliances demonstrates compliance with NFPA 99 as a complete system. Service manuals, instructions, and procedures provided by the manufacturer include information as required by 10.5.3.1.1 and are considered in the development of a program for electrical equipment maintenance. Electrical equipment instructions and maintenance manuals are readily available, and safety labels and condensed operating instructions on the appliance are legible. A record of electrical equipment tests, repairs, and modifications is maintained for a period of time to demonstrate compliance in accordance with the facility's policy. Personnel responsible for the testing, maintenance and use of electrical appliances receive continuing training. 10.3, 10.5.2.1, 10.5.2.1.2, 10.5.2.5, 10.5.3, 10.5.6, 10.5.8				
K922	Gas Equipment – Other				
NJZZ	List in the REMARKS section any NFPA 99 Chapter 11 Gas Equipment requirements that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567. Chapter 11 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K923	Gas Equipment – Cylinder and Container Storage				
	≥ 3,000 cubic feet				
	Storage locations are designed, constructed, and ventilated in accordance with 5.1.3.3.2 and 5.1.3.3.3.				
	> 300 but <3,000 cubic feet				
	Storage locations are outdoors in an enclosure or within an enclosed interior space of non- or limited- combustible construction, with door (or gates outdoors) that can be secured. Oxidizing gases are not stored with flammables, and are separated from combustibles by 20 feet (5 feet if sprinklered) or enclosed in a cabinet of noncombustible construction having a minimum 1/2 hr. fire protection rating.				
	≤ 300 cubic feet				
	In a single smoke compartment, individual cylinders available for immediate use in patient care areas with an aggregate volume of ≤ 300 cubic feet are not required to be stored in an enclosure. Cylinders must be handled with precautions as specified in 11.6.2.				
	A precautionary sign readable from 5 feet is on each door or gate of a cylinder storage room, where the sign includes the wording as a minimum "CAUTION: OXIDIZING GAS(ES) STORED WITHIN NO SMOKING".				
	Storage is planned so cylinders are used in order of which they are received from the supplier. Empty cylinders are segregated from full cylinders. When facility employs cylinders with integral pressure gauge, a threshold pressure considered empty is established. Empty cylinders are marked to avoid confusion. Cylinders stored in the open are protected from weather.				
K924	11.3.1, 11.3.2, 11.3.3, 11.3.4, 11.6.5 (NFPA 99) Gas Equipment – Testing and Maintenance Requirements				
N924	Anesthesia apparatus are tested at the final path to patient after any adjustment, modification or repair. Before the apparatus is returned to service, each connection is checked to verify proper gas and an oxygen analyzer is used to verify oxygen concentration. Defective equipment is immediately removed from service. Areas designated for servicing of oxygen equipment are clean and free of oil, grease, or other flammables. Manufacturer service manuals are used to maintain equipment and a scheduled maintenance program is followed. 11.4.1.3, 11.5.1.3, 11.6.2.5, 11.6.2.6 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K925	Gas Equipment – Respiratory Therapy Sources of Ignition				
	Smoking materials are removed from patients receiving respiratory therapy. When a nasal cannula is delivering oxygen outside of a patient's room, no sources of ignition are within in the site of intentional expulsion (1-foot). When other oxygen deliver equipment is used or oxygen is delivered inside a patient's room, no sources of ignition are within the area are of administration (15-feet). Solid fuel-burning appliances is not in the area of administration. Nonmedical appliances with hot surfaces or sparking mechanisms are not within oxygen-delivery equipment or site of intentional expulsion. 11.5.1.1, TIA 12-6 (NFPA 99)				
K926	Gas Equipment – Qualifications and Training of Personnel				
	Personnel concerned with the application, maintenance and handling of medical gases and cylinders are trained on the risk. Facilities provide continuing education, including safety guidelines and usage requirements. Equipment is serviced only by personnel trained in the maintenance and operation of equipment. 11.5.2.1 (NFPA 99)				
K927	Gas Equipment – Transfilling Cylinders				
	Transfilling of oxygen from one cylinder to another is in accordance with CGA P-2.5, <i>Transfilling of High Pressure Gaseous Oxygen Used for Respiration</i> . Transfilling of any gas from one cylinder to another is prohibited in patient care rooms. Transfilling to liquid oxygen containers or to portable containers over 50 psi comply with conditions under 11.5.2.3.1 (NFPA 99). Transfilling to liquid oxygen containers or to portable containers under 50 psi comply with conditions under 11.5.2.3.2 (NFPA 99). 11.5.2.2 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K928	Gas Equipment – Labeling Equipment and Cylinders				
	Equipment listed for use in oxygen-enriched atmospheres are so labeled. Oxygen metering equipment and pressure reducing regulators are labeled "OXYGEN-USE NO OIL". Flowmeters, pressure reducing regulators, and oxygen-dispensing apparatus are clearly and permanently labeled designating the gases for which they are intended. Oxygen-metering equipment, pressure reducing regulators, humidifiers, and nebulizers are labeled with name of manufacturer or supplier. Cylinders and containers are labeled in accordance with CGA C-7. Color coding is not utilized as the primary method of determining cylinder or container contents. All labeling is durable and withstands cleaning or disinfecting.				
K929	11.5.3.1 (NFPA 99) Gas Equipment – Precautions for Handling Oxygen Cylinders and Manifolds				
	Handling of oxygen cylinders and manifolds is based on CGA G-4, Oxygen. Oxygen cylinders, containers, and associated equipment are protected from contact with oil and grease, from contamination, protected from damage, and handled with care in accordance with precautions provided under 11.6.2.1 through 11.6.2.4 (NFPA 99).				
K930	Gas Equipment – Liquid Oxygen Equipment				
	The storage and use of liquid oxygen in base reservoir containers and portable containers comply with sections 11.7.2 through 11.7.4 (NFPA 99). 11.7 (NFPA 99)				
K931	Hyperbaric Facilities				
	All occupancies containing hyperbaric facilities comply with construction, equipment, administration, and maintenance requirements of NFPA 99. Chapter 14 (NFPA 99)				
K932	Features of Fire Protection – Other List in the REMARKS section any NFPA 99 Chapter 15 Features of Fire Protection requirements that are not addressed by the provided K-Tags, but are deficient. This information, along with the applicable Life Safety Code or NFPA standard citation, should be included on Form CMS-2567. Chapter 15 (NFPA 99)				

ID PREFIX		MET	NOT MET	N/A	REMARKS
K933	Peatures of Fire Protection – Fire Loss Prevention in Operating Rooms Periodic evaluations are made of hazards that could be encountered during surgical procedures, and fire prevention procedures are established. When flammable germicides or antiseptics are employed during surgeries utilizing electrosurgery, cautery or lasers: • packaging is non-flammable. • applicators are in unit doses. • Preoperative "time-out" is conducted prior the initiation of any surgical procedure to verify: • application site is dry prior to draping and use of surgical equipment. • pooling of solution has not occurred or has been corrected. • solution-soaked materials have been removed from the OR prior to draping and use of surgical devices. • policies and procedures are established outlining safety precautions related to the use of flammable germicide or antiseptic use. Procedures are established for operating room emergencies including alarm activation, evacuation, equipment shutdown, and control operations. Emergency procedures include the control of chemical spills, and extinguishment of drapery, clothing and equipment fires. Training is provided to new OR personnel (including surgeons), continuing education is provided, incidents are reviewed monthly, and procedures are reviewed annually. 15.13 (NFPA 99)				

Name of Facility 20	2012 LIFE SAFETY CODE
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PART III – RECOMMENDATION FOR WAIVER OF SPECIFIC LIFE SAFETY CODE PROVISIONS

For each item of the Life Safety Code recommended for waiver, list the survey report form item number and state the reason for the conclusion that: (a) the specific provisions of the code, if rigidly applied, would result in unreasonable hardship on the facility, and (b) the waiver of such unmet provisions will not adversely affect the health and safety of the patients. If additional space is required, attach additional sheet(s).

PROVISION NUMBER(S)	JUSTIFICATION
K400	

Title	Office	Date
Title	Office	Dete
ritie	Office	Date
	Title Title	

PART IV - FIRE SAFETY SURVEY REPORT CRUCIAL DATA EXTRACT (TO BE USED WITH CMS 2786 FORMS)

Provider Number		Facility Name		Survey Date						
K1						*K4				
			I			1 131				
		OF PLAN	K3 MULT	IPLE CONSTRUCTION	ON A. BUILDING					
	APPF	ROVAL	TOTAL NUME	BER OF BUILDINGS		」 B. WING				
					C. FLOOR					
			NUMBER OF	THIS BUILDING		D. APARTMEN	T UNIT			
LSC	FORM	M INDICATOR	· I		COMPLETE IF I	ICF/IID IS SURVEYE	D UNDER CHAPTER 33,			
		HEALTH	CARE FORM		EXISTING					
	12	2786R	2012 EXISTIN	G	SMALL (1	6 BEDS OR LESS)				
	13	2786R	2012 NEW			1. PROMP	Т			
					K8	2. SLOW 3. IMPRAC	CTICAL			
		AHC	O FORM		LARGE					
	14	2786U	2012 EXISTIN	G						
	15	2786U	2012 NEW			4. PROMP 5. SLOW	Т			
					K8	6. IMPRAC	CTICAL			
ICF/IID FORM				APARTMENT HOUSE						
	16	2786V, W, X	2012 EXISTIN	G	AIAKIMENI	⊓ 7. PROMP	т			
	17	2786V, W, X	2012 NEW		K8	8. SLOW	1			
						□ 9. IMPRAC	CTICAL			
*K7				SED FROM ABOVE						
	3	SELECT NUMB	ER OF FORM O	SED FROM ABOVE						
(Cho	ok if k	(221 or K251 or	e marked as not	annliaahla	COMPLETE IF ICF/IID IS SURVEYED UNDER CHAPTER 33, EXISTING					
•		6 M, R, T, U, V,		аррисавіе	ENTER E – SCORE					
			,		ENTER E - SCORE					
		K321:	K351:		K5:	e.g. 2.5				
*K9	ΕΛ	CILITY MEETS	S I SC BASED O	N (Check all that App	[/v)					
			Г		·y/					
	A′	1.	A2.	A3	3.	A4.	A5.			
		MP. WITH ALL ROVISIONS)	(ACCEP	TABLE POC)	(WAIVERS)	(FSES)	(PERFORMANCE BASED DESIGN)			
FAC	ILITY	DOES NOT ME	ET LSC	K0180						
				A.	В.		C			
B. FULLY SPRINKLE			SED DVDTIVI	LY SPRINKLERED) NONE					
			_	(All required areas ar		Il required areas are	(No sprinkler system)			
*840	ND A T	ODV		sprinklered)		sprinklered)				
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GOLDEN LIVINGCENTER - DELANO 433 COUNTY ROAD 30 DELANO, MN 55328

Smoke Barrier
Fire Seperation
Required EXIT

Fully Sprinkled

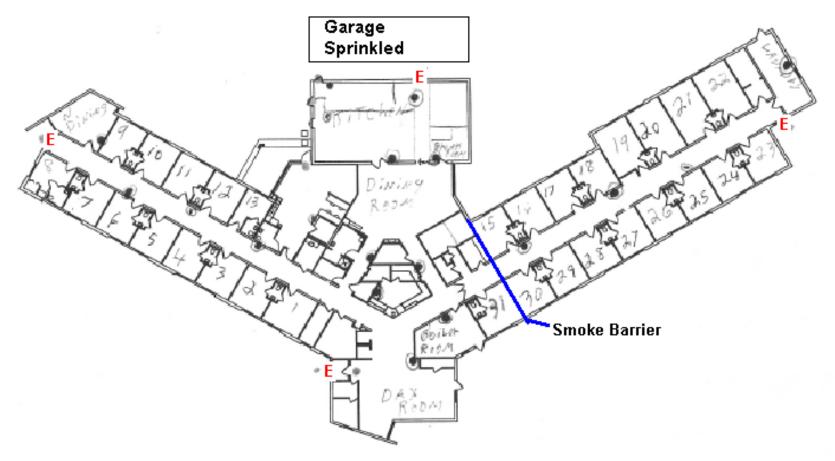


1st Floor



GLS 07/19/12

JAA 03/25/2014



Minnesota	State Fire Mars	hal Division-CMS Survey Draft Statemen	nt of Deficiencies	Page of				
PROJEC	T NUMBER:	PROVIDER NAME		SURVEY DATE				
Adminis	strator:	I.	Phone Numb	DÈT:				
Email a	Email address:							
State Fire Inspector:								
These are preliminary findings only. A complete and final Statement of Deficiencies 2567 report will be provided by US Mail.								
At the time of this inspection, this facility was found to comply with the requirements of the 2012 Life Safety Code applicable to: SNF/NF Hospital CFMR ASC Facilities participating in the Medicare/Medicaid programs. The following fire/life safety deficiencies were found during this inspection:								
K TAG S& S		Summary of Deficiency(ies)	Revisit	☐ Clearance				