

DEPARTMENT OF HEALTH AND HUMAN SERVICES

CENTERS FOR MEDICARE & MEDICAID SERVICES

MEDICARE/MEDICAID CERTIFICATION AND TRANSMITTAL

ID: KMBS

PART I - TO BE COMPLETED BY THE STATE SURVEY AGENCY

Facility ID: 00907

1. MEDICARE/MEDICAID PROVIDER NO. (L1) 245212 2.STATE VENDOR OR MEDICAID NO. (L2) 623840800	3. NAME AND ADDRESS OF FACILITY (L3) ESSENTIA HEALTH OAK CROSSING (L4) 1040 LINCOLN AVENUE (L5) DETROIT LAKES, MN (L6) 56501	4. TYPE OF ACTION: <u>7</u> (L8) <div style="display: flex; justify-content: space-between;"> <div> 1. Initial 3. Termination 5. Validation 7. On-Site Visit </div> <div> 2. Recertification 4. CHOW 6. Complaint 9. Other </div> </div> 8. Full Survey After Complaint FISCAL YEAR ENDING DATE: (L35) 06/30
5. EFFECTIVE DATE CHANGE OF OWNERSHIP (L9) 6. DATE OF SURVEY 08/25/2015 (L34) 8. ACCREDITATION STATUS: (L10) 0 Unaccredited 1 TJC 2 AOA 3 Other	7. PROVIDER/SUPPLIER CATEGORY <u>02</u> (L7) 01 Hospital 05 HHA 09 ESRD 13 PTIP 22 CLIA 02 SNF/NF/Dual 06 PRTF 10 NF 14 CORF 03 SNF/NF/Distinct 07 X-Ray 11 ICF/IID 15 ASC 04 SNF 08 OPT/SP 12 RHC 16 HOSPICE	
11. LTC PERIOD OF CERTIFICATION From (a) : To (b) : 12.Total Facility Beds 96 (L18) 13.Total Certified Beds 96 (L17)	10.THE FACILITY IS CERTIFIED AS: X A. In Compliance With <u>And/Or Approved Waivers Of The Following Requirements:</u> Program Requirements Compliance Based On: ___ 1. Acceptable POC ___ 2. Technical Personnel ___ 3. 24 Hour RN ___ 4. 7-Day RN (Rural SNF) ___ 5. Life Safety Code ___ 6. Scope of Services Limit ___ 7. Medical Director ___ 8. Patient Room Size ___ 9. Beds/Room B. Not in Compliance with Program Requirements and/or Applied Waivers: * Code: A (L12)	
14. LTC CERTIFIED BED BREAKDOWN <div style="display: flex; justify-content: space-around;"> <div>18 SNF (L37)</div> <div>18/19 SNF 96 (L38)</div> <div>19 SNF (L39)</div> <div>ICF (L42)</div> <div>IID (L43)</div> </div>		15. FACILITY MEETS 1861 (e) (1) or 1861 (j) (1): (L15)
16. STATE SURVEY AGENCY REMARKS (IF APPLICABLE SHOW LTC CANCELLATION DATE):		
17. SURVEYOR SIGNATURE <u>Gail Anderson, Unit Supervisor</u>	Date : 08/25/2015 (L19)	18. STATE SURVEY AGENCY APPROVAL <u>Mark Meath, Enforcement Specialist</u>
Date: 08/25/2015 (L20)		
PART II - TO BE COMPLETED BY HCFA REGIONAL OFFICE OR SINGLE STATE AGENCY		
19. DETERMINATION OF ELIGIBILITY <input checked="" type="checkbox"/> 1. Facility is Eligible to Participate <input type="checkbox"/> 2. Facility is not Eligible (L21)	20. COMPLIANCE WITH CIVIL RIGHTS ACT:	21. 1. Statement of Financial Solvency (HCFA-2572) 2. Ownership/Control Interest Disclosure Stmt (HCFA-1513) 3. Both of the Above : _____
22. ORIGINAL DATE OF PARTICIPATION 11/01/1976 (L24)	23. LTC AGREEMENT BEGINNING DATE (L41)	24. LTC AGREEMENT ENDING DATE (L25)
25. LTC EXTENSION DATE: (L27)	27. ALTERNATIVE SANCTIONS A. Suspension of Admissions: (L44) B. Rescind Suspension Date: (L45)	
26. TERMINATION ACTION: (L30) VOLUNTARY 00 01-Merger, Closure 02-Dissatisfaction W/ Reimbursement 03-Risk of Involuntary Termination 04-Other Reason for Withdrawal	INVOLUNTARY 05-Fail to Meet Health/Safety 06-Fail to Meet Agreement OTHER 07-Provider Status Change 00-Active	
28. TERMINATION DATE:	29. INTERMEDIARY/CARRIER NO. 03001 (L28) (L31)	30. REMARKS
31. RO RECEIPT OF CMS-1539 (L32)	32. DETERMINATION OF APPROVAL DATE 08/19/2015 (L33)	
DETERMINATION APPROVAL		



Protecting, Maintaining and Improving the Health of Minnesotans

CMS Certification Number (CCN): 245212

August 25, 2015

Ms. Christy Brinkman, Administrator
Essentia Health Oak Crossing
1040 Lincoln Avenue
Detroit Lakes, Minnesota 56501

Dear Ms. Brinkman:

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 August 14, 2015 the above facility is certified for:

96 Skilled Nursing Facility/Nursing Facility Beds

Your facility's Medicare approved area consists of all 96 skilled nursing facility beds.

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 Medicaid provider agreement may be subject to non-renewal or termination.

Feel free to contact me if you have questions related to this eNotice.

Sincerely,

A handwritten signature in black ink that reads "Mark Meath".

Mark Meath, Enforcement Specialist
Program Assurance Unit
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Email: mark.meath@state.mn.us
Telephone: (651) 201-4118 Fax: (651) 215-9697



Protecting, Maintaining and Improving the Health of Minnesotans

Electronically delivered
August 25, 2015

Ms. Christy Brinkman, Administrator
Essentia Health Oak Crossing
1040 Lincoln Avenue
Detroit Lakes, Minnesota 56501

RE: Project Number S5212024

Dear Ms. Brinkman:

On July 24, 2015, we informed you that we would recommend enforcement remedies based on the deficiencies cited by this Department for a standard survey, completed on July 10, 2015. This survey found the most serious deficiencies to be a pattern of deficiencies that constituted no actual harm with potential for more than minimal harm that was not immediate jeopardy (Level E), whereby corrections were required.

On August 25, 2015, the Minnesota Department of Health completed a Post Certification Revisit (PCR) by review of your plan of correction to verify that your facility had achieved and maintained compliance with federal certification deficiencies issued pursuant to a standard survey, completed on July 10, 2015. We presumed, based on your plan of correction, that your facility had corrected these deficiencies as of August 14, 2015. Based on our PCR, we have determined that your facility has corrected the deficiencies issued pursuant to our standard survey, completed on July 10, 2015, effective August 14, 2015 and therefore remedies outlined in our letter to you dated July 24, 2015, will not be imposed.

Please note, it is your responsibility to share the information contained in this letter and the results of this visit with the President of your facility's Governing Body.

Feel free to contact me if you have questions related to this eNotice.

Sincerely,

A handwritten signature in black ink that reads "Mark Meath".

Mark Meath, Enforcement Specialist
Program Assurance Unit
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Email: mark.meath@state.mn.us
Telephone: (651) 201-4118 Fax: (651) 215-9697

Post-Certification Revisit Report

Public reporting 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 CMS, Office of Financial Management, P.O. Box 26684, Baltimore, MD 21207; and to the Office of Management and Budget, Paperwork Reduction Project (0938-0390), Washington, D.C. 20503.

(Y1) Provider / Supplier / CLIA / Identification Number 245212	(Y2) Multiple Construction A. Building B. Wing	(Y3) Date of Revisit 8/25/2015
Name of Facility ESSENTIA HEALTH OAK CROSSING		Street Address, City, State, Zip Code 1040 LINCOLN AVENUE DETROIT LAKES, MN 56501

This report is completed by a qualified State surveyor for the Medicare, Medicaid and/ or Clinical Laboratory Improvement Amendments program, to show those deficiencies previously reported on the CMS-2567, Statement of Deficiencies and Plan of Correction that have been corrected and the date such corrective action was accomplished. Each deficiency should be fully identified using either the regulation or LSC provision number and the identification prefix code previously shown on the CMS-2567 (prefix codes shown to the left of each requirement on the survey report form).

(Y4) Item	(Y5) Date	(Y4) Item	(Y5) Date	(Y4) Item	(Y5) Date
ID Prefix F0167 Reg. # 483.10(g)(1) LSC	Correction Completed 08/14/2015	ID Prefix F0176 Reg. # 483.10(n) LSC	Correction Completed 08/14/2015	ID Prefix F0278 Reg. # 483.20(g) - (i) LSC	Correction Completed 08/14/2015
ID Prefix F0371 Reg. # 483.35(i) LSC	Correction Completed 08/14/2015	ID Prefix F0441 Reg. # 483.65 LSC	Correction Completed 08/14/2015	ID Prefix Reg. # LSC	Correction Completed
ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed
ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed
ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed

Reviewed By State Agency	Reviewed By GA/mm	Date: 08/25/2015	Signature of Surveyor: 28034	Date: 08/25/2015		
Reviewed By CMS RO	Reviewed By	Date:	Signature of Surveyor:	Date:		
Followup to Survey Completed on: 7/10/2015		Check for any Uncorrected Deficiencies. Was a Summary of Uncorrected Deficiencies (CMS-2567) Sent to the Facility? <table border="0"> <tr> <td>YES</td> <td>NO</td> </tr> </table>			YES	NO
YES	NO					

ID: KMBS
Facility ID: 00907

PART II - TO BE COMPLETED BY HCFA REGIONAL OFFICE OR SINGLE STATE AGENCY

020499



Protecting, Maintaining and Improving the Health of Minnesotans

Electronically delivered
July 24, 2015

Ms. Christy Brinkman, Administrator
Essentia Health Oak Crossing
1040 Lincoln Avenue
Detroit Lakes, Minnesota 56501

RE: Project Number S5212024

Dear Ms. Brinkman:

On July 10, 2015, a standard survey was completed at your facility by the Minnesota Departments 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 a pattern of deficiencies that constitute no actual harm with potential for more than minimal harm that is not immediate jeopardy (Level E), as evidenced by the attached CMS-2567 whereby corrections are required. A copy of the Statement of Deficiencies (CMS-2567) is enclosed.

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.

This letter provides important information regarding your response to these deficiencies and addresses the following issues:

Opportunity to Correct - the facility is allowed an opportunity to correct identified deficiencies before remedies are imposed;

Electronic Plan of Correction - when a plan of correction will be due and the information to be contained in that document;

Remedies - the type of remedies that will be imposed with the authorization of the Centers for Medicare and Medicaid Services (CMS) if substantial compliance is not attained at the time of a revisit;

Potential Consequences - the consequences of not attaining substantial compliance 3 and 6 months after the survey date; and

Informal Dispute Resolution - your right to request an informal reconsideration to dispute the attached deficiencies.

Please note, it is your responsibility to share the information contained in this letter and the results of this visit with the President of your facility's Governing Body.

DEPARTMENT CONTACT

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

Gail Anderson, Unit Supervisor
Minnesota Department of Health
1505 Pebble Lake Road #300
Fergus Falls, Minnesota 56537
gail.anderson@state.mn.us
Telephone: (218) 332-5140
Fax: (218) 332-5196

OPPORTUNITY TO CORRECT - DATE OF CORRECTION - REMEDIES

As of January 14, 2000, CMS policy requires that facilities will not be given an opportunity to correct before remedies will be imposed when actual harm was cited at the last standard or intervening survey and also cited at the current survey. Your facility does not meet this criterion. Therefore, if your facility has not achieved substantial compliance by August 19, 2015, the Department of Health will impose the following remedy:

- State Monitoring. (42 CFR 488.422)

ELECTRONIC PLAN OF CORRECTION (ePoC)

An ePoC for the deficiencies must be submitted within **ten calendar days** of your receipt of this letter. Your ePoC must:

- 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;
- Address what measures will be put into place or systemic changes made to ensure that the deficient practice will not recur;
- Indicate how the facility plans to monitor its performance to make sure that solutions are

sustained. The facility must develop a plan for ensuring that correction is achieved and sustained. This plan must be implemented, and the corrective action evaluated for its effectiveness. The plan of correction is integrated into the quality assurance system;

- Include dates when corrective action will be completed. The corrective action completion dates must be acceptable to the State. If the plan of correction is unacceptable for any reason, the State will notify the facility. If the plan of correction is acceptable, the State will notify the facility. Facilities should be cautioned that they are ultimately accountable for their own compliance, and that responsibility is not alleviated in cases where notification about the acceptability of their plan of correction is not made timely. The plan of correction will serve as the facility's allegation of compliance; and,
- Submit electronically to acknowledge your receipt of the electronic 2567, your review and your ePoC submission.

The state agency may, in lieu of a revisit, determine correction and compliance by accepting the facility's ePoC if the ePoC is reasonable, addresses the problem and provides evidence that the corrective action has occurred.

If an acceptable ePoC is not received within 10 calendar days from the receipt of this letter, we will recommend to the CMS Region V Office that one or more of the following remedies be imposed:

- Optional denial of payment for new Medicare and Medicaid admissions (42 CFR 488.417 (a));
- Per day civil money penalty (42 CFR 488.430 through 488.444).

Failure to submit an acceptable ePoC could also result in the termination of your facility's Medicare and/or Medicaid agreement.

PRESUMPTION OF COMPLIANCE - CREDIBLE ALLEGATION OF COMPLIANCE

The facility's ePoC will serve as your allegation of compliance upon the Department's acceptance. Your signature at the bottom of the first page of the CMS-2567 form will be used as verification of compliance. 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, Licensing and Certification Program staff and/or the Department of Public Safety, State Fire Marshal Division staff, if your ePoC for the respective deficiencies (if any) is acceptable.

VERIFICATION OF SUBSTANTIAL COMPLIANCE

Upon receipt of an acceptable ePoC, an onsite revisit of your facility may be conducted to validate that substantial compliance with the regulations has been attained in accordance with your verification. A Post Certification Revisit (PCR) will occur after the date you identified that compliance was achieved

in your plan of correction.

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.

Original deficiencies not corrected

If your facility has not achieved substantial compliance, we will impose the remedies described above. If the level of noncompliance worsened to a point where a higher category of remedy may be imposed, we will recommend to the CMS Region V Office that those other remedies be imposed.

Original deficiencies not corrected and new deficiencies found during the revisit

If new deficiencies are identified at the time of the revisit, those deficiencies may be disputed through the informal dispute resolution process. However, the remedies specified in this letter will be imposed for original deficiencies not corrected. If the deficiencies identified at the revisit require the imposition of a higher category of remedy, we will recommend to the CMS Region V Office that those remedies be imposed.

Original deficiencies corrected but new deficiencies found during the revisit

If new deficiencies are found at the revisit, the remedies specified in this letter will be imposed. If the deficiencies identified at the revisit require the imposition of a higher category of remedy, we will recommend to the CMS Region V Office that those remedies be imposed. You will be provided the required notice before the imposition of a new remedy or informed if another date will be set for the imposition of these remedies.

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

If substantial compliance with the regulations is not verified by October 10, 2015 (three months after the identification of noncompliance), the CMS Region V Office must deny payment for new admissions as mandated by the Social Security Act (the Act) at Sections 1819(h)(2)(D) and 1919(h)(2)(C) and Federal regulations at 42 CFR Section 488.417(b). This mandatory denial of payments will be based on the failure to comply with deficiencies originally contained in the Statement of Deficiencies, upon the identification of new deficiencies at the time of the revisit, or if deficiencies have been issued as the result of a complaint visit or other survey conducted after the original statement of deficiencies was issued. This mandatory denial of payment is in addition to any remedies that may still be in effect as of this date.

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 January 10, 2016 (six months after the

identification of noncompliance) if your facility does not achieve substantial compliance. This action is mandated by the Social Security Act at Sections 1819(h)(2)(C) and 1919(h)(3)(D) and Federal regulations at 42 CFR Sections 488.412 and 488.456.

INFORMAL DISPUTE RESOLUTION

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: http://www.health.state.mn.us/divs/fpc/profinfo/lrc/lrc_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 plan of correction. A copy of the Department's informal dispute resolution policies are posted on the MDH Information Bulletin website at: <http://www.health.state.mn.us/divs/fpc/profinfo/infobul.htm>

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.

Feel free to contact me if you have questions.

Sincerely,



Kamala Fiske-Downing, Program Specialist
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Kamala.Fiske-Downing@state.mn.us
Telephone: (651) 201-4112 Fax: (651) 215-9697

DEPARTMENT OF HEALTH AND HUMAN SERVICES
CENTERS FOR MEDICARE & MEDICAID SERVICES

PRINTED: 08/06/2015
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245212	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED 07/10/2015
NAME OF PROVIDER OR SUPPLIER ESSENTIA HEALTH OAK CROSSING			STREET ADDRESS, CITY, STATE, ZIP CODE 1040 LINCOLN AVENUE DETROIT LAKES, MN 56501		
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)		(X5) COMPLETION DATE
F 000	INITIAL COMMENTS The facility's plan of correction (POC) will serve as your allegation of compliance upon the Department's acceptance. Because you are enrolled in ePOC, your signature is not required at the bottom of the first page of the CMS-2567 form. Your electronic submission of the POC will be used as verification of compliance. Upon receipt of an acceptable electronic POC, an on-site revisit of your facility may be conducted to validate that substantial compliance with the regulations has been attained in accordance with your verification.	F 000			
F 167 SS=C	483.10(g)(1) RIGHT TO SURVEY RESULTS - READILY ACCESSIBLE A resident has the right to examine the results of the most recent survey of the facility conducted by Federal or State surveyors and any plan of correction in effect with respect to the facility. The facility must make the results available for examination and must post in a place readily accessible to residents and must post a notice of their availability. This REQUIREMENT is not met as evidenced by: Based on observation, interview and document review the facility failed to ensure the most recent federal survey results were readily available and accessible to residents, families and visitors of the facility. This deficient practice had the potential to affect all 83 residents residing in the facility, families and/or visitors.	F 167	Previous year survey results were posted immediately when provider notified (July 10, 2015) Facility will secure survey documents in the resident information center located outside administration.		8/14/15

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE

TITLE

(X6) DATE

Electronically Signed

07/27/2015

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.

DEPARTMENT OF HEALTH AND HUMAN SERVICES
CENTERS FOR MEDICARE & MEDICAID SERVICES

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NAME OF PROVIDER OR SUPPLIER ESSENTIA HEALTH OAK CROSSING			STREET ADDRESS, CITY, STATE, ZIP CODE 1040 LINCOLN AVENUE DETROIT LAKES, MN 56501		
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F 167	Continued From page 1 Findings include: During the initial tour of the facility on 7/7/15, at 1:30 p.m., a wooden, wall mounted three-tiered holder was observed on the wall to the left of the administration hallway, in the main level of the facility. The lowest section of the wall mounted three-tiered holder had letters that spelled out survey information on the front of the holder. Inside the lowest section, a post certification revisit dated 12/3/2013, and two separate reports/letters from the Office of Health Facility Complaints dated, 4/1/14 and 4/11/14 were observed. However, the holder lacked a copy of the results of the most recent recertification survey conducted in the facility. Continued observations on 7/8/15 at 1:33 p.m., and 7/10/15 at 3:22 p.m. revealed the same documents in the survey information section of the three-tiered letter holder attached to the wall. No federal survey results were observed in any of the sections of the letter holder or the surrounding area. During an interview on 7/10/15, at 3:23 p.m., director of nursing (DON) verified the federal survey results for the most recent recertification survey were missing. The DON indicated she was not aware of where the survey results were and indicated it could have been possible a resident may have taken the survey results. A policy was requested, but not provided.	F 167	Resident will be notified upon admission the whereabouts of these documents Weekly audit will be performed to assure documents are in place Audits will go to facilities QAPI to review audit findings (10-5-15) and to determine future audit requirements.		
F 176 SS=D	483.10(n) RESIDENT SELF-ADMINISTER DRUGS IF DEEMED SAFE	F 176			8/14/15

DEPARTMENT OF HEALTH AND HUMAN SERVICES
CENTERS FOR MEDICARE & MEDICAID SERVICES

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STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245212	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED 07/10/2015
NAME OF PROVIDER OR SUPPLIER ESSENTIA HEALTH OAK CROSSING			STREET ADDRESS, CITY, STATE, ZIP CODE 1040 LINCOLN AVENUE DETROIT LAKES, MN 56501		
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)		(X5) COMPLETION DATE
F 176	<p>Continued From page 2</p> <p>An individual resident may self-administer drugs if the interdisciplinary team, as defined by §483.20(d)(2)(ii), has determined that this practice is safe.</p> <p>This REQUIREMENT is not met as evidenced by: Based on observation, interview, and document review, the facility failed to ensure a compressive self administration medication assessment was completed to determine ability of safe self medication administration for 1 of 1 residents (R196) observed to self administer medications during the survey.</p> <p>Findings include:</p> <p>During a medication pass observation on 7/7/15, at 5:38 p.m. licensed practical nurse (LPN)-A dispensed (a blood pressure medication) metoprolol 100 mg (milligrams) into a white paper cup. LPN-A placed the white paper medication cup on the dining table next to R196, and walked back to medication cart. LPN- A signed the medication as given on the electronic medication administration form and then proceeded with her duties. LPN-A did not observe or return to R196 during the entire observation. At 5:53 p.m. LPN-A left the dining room and pushed the medication cart down a hall of the facility, out of view of R196.</p> <p>R196's admission Minimum Data Set (MDS) care area assessment (CAA), dated 6/30/15, identified R196 had severe cognitive impairment, with diagnoses which included dementia, had confusion, disorientation, forgetfulness, difficulty with recall and short and long term memory.</p>	F 176	<p>DON re-affirmed for R-196 that she was not appropriate (per policy) for self-administration of medications (7-10-15.) This Resident has discharged back home with her husband.</p> <p>All current Residents will be assessed for ability to self-administer medications by 8-14-15.</p> <p>Facility reviewed current procedure and assessment for self-administration of drugs. Revisions made to assess every Resident upon admission for ability to self-administer medications.</p> <p>LPN-A was educated in the facilities Policy and Procedure for self-administration.</p> <p>All Nurses will review facilities policy and procedure for self-administration of medications and demonstrate competency per a post test.</p> <p>Random audits of medication passes will be completed weekly in all neighborhoods by RN clinical Coordinator.</p> <p>Findings will be reviewed by facilities QAPI and determination of future audit</p>		

DEPARTMENT OF HEALTH AND HUMAN SERVICES
CENTERS FOR MEDICARE & MEDICAID SERVICES

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NAME OF PROVIDER OR SUPPLIER ESSENTIA HEALTH OAK CROSSING			STREET ADDRESS, CITY, STATE, ZIP CODE 1040 LINCOLN AVENUE DETROIT LAKES, MN 56501		
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)	(X5) COMPLETION DATE	
F 176	<p>Continued From page 3</p> <p>During an interview on 7/07/2015, at 6:00 p.m. LPN-A stated, " [R196] can manage meds (medications) a little bit." LPN-A indicated she felt R196 would take the medications independently. LPN-A indicated a visitor was always with R196; therefore, no follow up was needed to ensure R196 received the medication.</p> <p>During an interview on 7/09/2015, at 8:56 a.m. LPN-B verified the facility practice of residents who self administer medications have initial assessments, a physician order, and were reviewed monthly for continued appropriateness of self medication administration.</p> <p>Review of R196's electronic and paper chart identified the following: The current physician's orders dated 6/10/15 - 7/10/15 identified an order for metoprolol 100 mg, oral twice a day. However, the record lacked a physician's order for the resident to self administer medications. The record lacked documentation a self administration of medication assessment had been completed, to determine if R196 was safe to self administer medications. R196s' care plan revised 7/8/15, did not identify R196 was safe to self administer medications.</p> <p>During an interview on 7/10/2015, 5:13 p.m. the director of nursing (DON) verified a resident self administration assessment was required to be completed in order for medications to be left at a residents side. The DON stated "the [facility] policy would prohibit" self administration without an assessment for safety completed.</p> <p>The undated facility policy titled Routine Orders</p>	F 176	requirements will occur on 10-5-15.		

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F 176	Continued From page 4	F 176			
F 278 SS=D	<p>identified, Self-Administration of Medications identified "May initiate after assessment and care plan.</p> <p>483.20(g) - (j) ASSESSMENT ACCURACY/COORDINATION/CERTIFIED</p> <p>The assessment must accurately reflect the resident's status.</p> <p>A registered nurse must conduct or coordinate each assessment with the appropriate participation of health professionals.</p> <p>A registered nurse must sign and certify that the assessment is completed.</p> <p>Each individual who completes a portion of the assessment must sign and certify the accuracy of that portion of the assessment.</p> <p>Under Medicare and Medicaid, an individual who willfully and knowingly certifies a material and false statement in a resident assessment is subject to a civil money penalty of not more than \$1,000 for each assessment; or an individual who willfully and knowingly causes another individual to certify a material and false statement in a resident assessment is subject to a civil money penalty of not more than \$5,000 for each assessment.</p> <p>Clinical disagreement does not constitute a material and false statement.</p> <p>This REQUIREMENT is not met as evidenced by: Based on interview and document review, the</p>	F 278			8/14/15
			A correction MDS for R-60 was submitted		

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F 278	<p>Continued From page 5</p> <p>facility failed to ensure the Minimum Data Set (MDS) accurately reflected the pressure ulcer status for 1 of 2 residents (R60) identified with pressure ulcers.</p> <p>Findings include:</p> <p>R60's admission MDS dated 3/6/15, identified R60 was at risk for development of pressure ulcer, however did not identify R60 had a unhealed pressure ulcer and did not identify the presence of a stage 2 pressure ulcer upon admission.</p> <p>R60's Physician Order Report dated 2/27/15 to 7/10/15 identified the following diagnoses: surgery aftercare, malignant brain and breast cancer, neuropathy and urinary incontinency.</p> <p>Review of R60's admission body observation dated 2/28/15, identified R60's right buttock had a pressure ulcer present upon admission and both buttocks were discolored from previous sores.</p> <p>Review of R60's Tissue Tolerance assessment dated 3/1/15, indicated R60 continued to have a pressure sore on left buttock, open area small eraser size</p> <p>Review of R60's Resident progress noted dated 3/2/15 indicated healing stage 2 area on right buttock.</p> <p>R60's pressure ulcer Care Area Assessment (CAA) dated 3/6/15, identified R60 was at risk for development of pressure ulcer, required extensive assistance with bed mobility and required a special mattress or seat cushion to reduce or relieve pressure. However, the CAA</p>	F 278	<p>on 7-15-15 (Admission and 14 day)</p> <p>A 100% record review was completed on current Residents as of 7-16-15 and no changes to section "M" were noted</p> <p>Training for all MDS Coordinators to include Leading Age module on Section M. DON reviewed facilities current Policy and Procedure and outlined document sources in our EMR (Matrix Care), both of which will be trained to MDS coordinators.</p> <p>Audits will be completed on all new admissions between July 17, 2015 and September 30th 2015 to assure accurate coding of Section M to documentation.</p> <p>Findings will be reviewed at facilities QAPI 10-5-15, and will make recommendations for further audit requirements.</p>		

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F 278	Continued From page 6 did not identify the presence of R60's unhealed stage 2 pressure ulcer and lacked an analysis of current treatments and interventions for the unhealed stage 2 pressure ulcer. On 7/10/15, at 6:01 p.m. the director of nursing stated she expected all the skin issues should have been thoroughly assessed and documented during admission. DON confirmed R60's admission MDS and CAA and stated she expected the MDS and CAA to have identified R60's pressure ulcer upon admission. The facility's Care Conferences, Care Planning policy dated 4/1/15, directed staff to include the following sources of information for the MDS: -Review of the residents medical record -Observation of the resident -Interviews and communication with the resident -Communication with healthcare providers -Communication with physician -Communication with the resident's family members	F 278			
F 371 SS=D	483.35(i) FOOD PROCURE, STORE/PREPARE/SERVE - SANITARY The facility must - (1) Procure food from sources approved or considered satisfactory by Federal, State or local authorities; and (2) Store, prepare, distribute and serve food under sanitary conditions This REQUIREMENT is not met as evidenced	F 371			8/14/15

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F 371	<p>Continued From page 7</p> <p>by: Based on observation, Interview and record review the facility failed to clean and maintain food service equipment to prevent food borne illness for residents residing in 2 of 4 neighborhoods (Cedar Ridge, Meadowbrook) in the facility.</p> <p>Findings include:</p> <p>On 07/10/15, at 4:24 p.m. observations of the Meadow Brook neighborhood ice machine were conducted with Nursing Home Support (NHS-A) present. On the front of ice machine, a black spout with a pink substance inside the opening of the spout was observed. NHS-A reached out with a disposable cloth and wiped up inside the black spout covers. The disposable cloth had both brown sludge smears and pink smudges on the cloth. In addition, there were several white hard water scale stains observed covering the upper right 1/3 of drip tray, and covered the top 3 grates of the drip tray. NHS-A stated she was aware the ice machine had a build up of pink substance in the past and indicated an outside company cleaned and serviced the ice machines for the facility. She indicated homemakers were to routinely clean the outside of the ice machines, however, cleaning the spouts and the inside of the spouts were not the usual practice for facility staff. NHS-A confirmed she was aware the ice machine had a build up of the pink substance and stated she had not routinely cleaned the spouts of the ice machine.</p> <p>On 07/10/15, at 4:56 p.m. the director of dietary (DD) stated the facility utilized an outside company to service and clean the ice machines.</p>	F 371	<p>Ice machines chutes were cleaned by survey completion (7-10-15)</p> <p>Policy and Procedure developed and approved by QAPI on 7-27-15 for sanitation and cleaning of the ice machines.</p> <p>Homemakers and Neighborhood supports will be trained on the frequency and standard procedure for cleaning the drip tray and ice machine chute.</p> <p>Neighborhood cleaning check lists were revised to include this cleaning on a weekly basis.</p> <p>Audits will be performed in all 4 neighborhoods for cleanliness and documentation of the cleaning by RN Supervisor.</p> <p>Audit findings will be reviewed by facilities QAPI on 10-5-15 and recommendations for further audits will be made.</p>		

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F 371	<p>Continued From page 8</p> <p>She indicated facility staff did not service or maintain the equipment and indicated the outside company came to facility twice a year to service and clean the ice machines. DD stated the "Daily Homemaker Cleaning Duties Record" utilized by facility staff identified the clean the beverage area, however, did not identify what parts of the ice machine to clean and how to clean the ice machine. She confirmed the findings and indicated she had not been aware of the sludge and pink substance in the ice machines. She stated "that is not good" and indicated she felt there could be a risk for food borne illness.</p> <p>On 07/10/15, at 5:06 p.m. observations of the Cedar Ridge neighborhood ice machine were conducted with Nursing Assistant (NA-A) present. NA-A reached out with a white paper napkin and wiped the inside of the black spout of the ice/water dispenser. Upon removal of the white paper napkin from the black spout, two areas of pink sludge were noted on the napkin. Each area of pink sludge was 2 cm (centimeters) wide by 2 cm long. NA-A verified the two areas of pink sludge and indicated she routinely cleaned the outside of the ice machine and tray every day and stated she did not routinely clean the spouts or inside of the spouts.</p> <p>Review of the most current invoice from the outside service company dated 2/12/15, identified The machine was cleaned and sanitized and inspected and checked for proper operation."</p> <p>Review of the Manufacturer's cleaning and maintenance instructions, provided by the facility, revealed the manufacturer recommended cleaning and sanitizing the ice machine at least</p>	F 371			

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F 371	Continued From page 9	F 371			
F 441 SS=E	<p>twice a year. The instructions also identified more frequent cleaning may be required in some existing water conditions. In addition, the instructions included monthly cleaning instructions for the exterior of the ice machine included using a neutral cleaner to wipe off oil or dirt build up. Clean any chlorine staining (rust or colored spots) using a non-abrasive cleaner.</p> <p>483.65 INFECTION CONTROL, PREVENT SPREAD, LINENS</p> <p>The facility must establish and maintain an Infection Control Program designed to provide a safe, sanitary and comfortable environment and to help prevent the development and transmission of disease and infection.</p> <p>(a) Infection Control Program The facility must establish an Infection Control Program under which it - (1) Investigates, controls, and prevents infections in the facility; (2) Decides what procedures, such as isolation, should be applied to an individual resident; and (3) Maintains a record of incidents and corrective actions related to infections.</p> <p>(b) Preventing Spread of Infection (1) When the Infection Control Program determines that a resident needs isolation to prevent the spread of infection, the facility must isolate the resident. (2) The facility must prohibit employees with a communicable disease or infected skin lesions from direct contact with residents or their food, if direct contact will transmit the disease. (3) The facility must require staff to wash their hands after each direct resident contact for which</p>	F 441			8/14/15

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F 441	<p>Continued From page 10 hand washing is indicated by accepted professional practice.</p> <p>(c) Linens Personnel must handle, store, process and transport linens so as to prevent the spread of infection.</p> <p>This REQUIREMENT is not met as evidenced by: Based on observation, interview and document review the facility failed to ensure proper infection control practices for expired disinfectant wipes used to sanitize multi-use glucometers, on 2 of 4 facility resident communities, Harbor Springs and Meadow Brook. This deficient practice had the potential to affect 10 of 10 residents (R18, R134, R114, R35, R58, R73, R71, R75, R111, R125) currently using the multi-used glucometers.</p> <p>Findings include:</p> <p>Observation of the Harbor Springs medication cart and storage room was conducted on 7/9/15, at 12:56 p.m. with licensed practical nurse (LPN)-A present. The medication/ treatment cart contained 1 open box of 50 Sani-wipes with less than 10 packages missing and an expiration date of 4/2015. The storage room contained 1/2 box of 50 sani-wipes with an expiration dated of 4/2015, and a second 1/2 box of 50 sani-wipes with expiration dated of 5/2015.</p> <p>During an interview on 7/09/201, at 12:56 p.m. LPN-A verified the above findings. LPN-A indicated the expired Sani-wipes were currently</p>	F 441	<p>Facility audited all sani-wipes in facility to assure no further expired supplies.</p> <p>Facility reviewed supply ordering practices (dates) to determine root cause of expired wipe to prevent reoccurrence.</p> <p>Facility requested this supply be added to a PAR level and managed by materials management (for rotation of supply)</p> <p>All nurses were trained on where the expiration date is stamped on the individual wrapped sani-wipe.</p> <p>Audit will be performed of the supply in the med cart weekly @ random times and neighborhoods. The findings will be reviewed by facilities QAPI on 10-5-15 and recommendation will be made for further audits.</p> <p>Quarterly audit will occur in all nursing supply areas and reported through QAPI until determined no further audits are necessary.</p>		

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F 441	<p>Continued From page 11</p> <p>being used to clean the medication cart and the multi patient use glucometer. LPN-A stated, "three residents" currently used the shared glucometer at this time. LPN-A indicated she did not routinely check for expiration dates of the sani-wipes.</p> <p>During an interview on 7/10/2015, at 12:24 p.m. registered nurse (RN)-C verified the usual facility practice was to disinfect the multi use glucometers with the Sani-wipes. RN-C verified the use of expired Sani-wipes for sanitation of multi patient use glucometers would not be an acceptable infection control process. RN-C stated, " It would not be known if the expired wipes had the ability to sanitize."</p> <p>During review of the Meadow Brook medication cart and storage rooms on 7/10/2015, at 12:35 p.m. with LPN-B, the following was found: The medication/ treatment cart contained 21 of 22 Sani-wipes with an expiration date of 5/2015, The supply storage room contained 1 full box of 50 sani-wipes with expiration date of 5/2015. The medication storage room contained a box with 10 sani-wipes remaining in the box with an expiration date of 5/2015.</p> <p>During an interview on 7/10/2015, at 12:35 p.m. LPN-B. verified the above findings and indicated the expired wipes were currently utilized to disinfect the multi-use glucometers in the facility. LPN-B indicated she did not know where the expiration dates were printed on the Sani-wipe box and individual packages, and stated she did not check the expiration dates. LPN-B verified 10 residents currently used the multi used glucometer (R18, R134, R114, R35, R58,R73, R71, R75, R111, R125).</p>	F 441			

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F 441	<p>Continued From page 12</p> <p>During an interview on 7/10/2015, at 5:13 p.m. the director of nursing (DON) confirmed the current facility policy and indicated the use of expired Sani-wipes for multi use glucometers would not be an acceptable facility practice.</p> <p>The facility provided manufacturers instructions titled Nova Biomedical Customer Information Bulletin, dated 12/4/12, directed to disinfect the meter" #1 Using a new, fresh germicidal wipe."</p>	F 441			

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STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245212	(X2) MULTIPLE CONSTRUCTION A. BUILDING 02 - EXISTING BUILDING 02 B. WING _____		(X3) DATE SURVEY COMPLETED 07/09/2015
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K 000	<p>INITIAL COMMENTS</p> <p>FIRE SAFETY</p> <p>A Life Safety Code Survey was conducted by the Minnesota Department of Public Safety. At the time of this survey Essentia Health Oak Crossing 02 Main Building was found in substantial compliance with the requirements for participation in Medicare/Medicaid at 42 CFR, Subpart 483.70(a), Life Safety from Fire, and the 2000 edition of National Fire Protection Association (NFPA) Standard 101, Life Safety Code (LSC), Chapter 19 Existing Health Care.</p> <p>02 Main Building</p> <p>The facility was surveyed as two buildings: Essentia Health Oak Crossing is a 2-story building with a basement. The building was constructed at 3 different times. The original building (02) was constructed in 1968, is 2-story building with a small basement and was determined to be of Type II(000) construction due to the on going remodeling of this building. In 1999 an Administration / Entrance addition was constructed south of the original building and an addition to the hospital north of the original building. The entrance addition is Type V (111) construction, 2-stories without a basement and the hospital addition is Type II (111) construction, 1-story without a basement. In 2008 a 2-story building, without a basement, separated with two 2-hour fire barriers south of the entrance addition and was determined to be Type II (111) construction. The buildings are divided into 12 smoke zones (6 per floor) by 2- hour and 30 minute fire barriers.</p> <p>The facility has a complete automatic fire</p>	K 000			

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE

TITLE

(X6) DATE

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.

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K 000	<p>Continued From page 1</p> <p>sprinkler system in accordance with NFPA 13 Standard for the Installation of Sprinkler Systems 1999 edition with 2 systems. The facility has a fire alarm system with manual pull station near each exit door, smoke detection in the corridor system properly spaced and all common areas in accordance with NFPA 72 "The National Fire Alarm Code" (1999 edition). The fire alarm system is monitored for automatic fire department notification. Hazardous areas have either heat detection or smoke detection that are on the fire alarm system in accordance with the Minnesota State Fire Code (2007 edition).</p> <p>The facility has a capacity of 96 beds and had a census of 80 at the time of the survey.</p> <p>The requirement at 42 CFR, Subpart 483.70(a) is MET.</p>	K 000			

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OMB NO. 0938-0391

F5212023

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245212	(X2) MULTIPLE CONSTRUCTION A. BUILDING 03 - 2008 SOUTH B. WING _____		(X3) DATE SURVEY COMPLETED 07/09/2015
NAME OF PROVIDER OR SUPPLIER ESSENTIA HEALTH OAK CROSSING			STREET ADDRESS, CITY, STATE, ZIP CODE 1040 LINCOLN AVENUE DETROIT LAKES, MN 56501		
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)		(X5) COMPLETION DATE
K 000	<p>INITIAL COMMENTS</p> <p>FIRE SAFETY</p> <p>A Life Safety Code Survey was conducted by the Minnesota Department of Public Safety. At the time of this survey Essentia Health Oak Crossing 03 South Building was found in substantial compliance with the requirements for participation in Medicare/Medicaid at 42 CFR, Subpart 483.70(a), Life Safety from Fire, and the 2000 edition of National Fire Protection Association (NFPA) Standard 101, Life Safety Code (LSC), Chapter 18 New Health Care.</p> <p>03 South Building</p> <p>The facility was surveyed as two buildings Essentia Health Oak Crossing is a 2-story building with a basement. The building was constructed at 3 different times. The original building (02) was constructed in 1968, is 2-story building with a small basement and was determined to be of Type II(000) construction due to the on going remodeling of this building. In 1999 an Administration / Entrance addition was constructed south of the original building and an addition to the hospital north of the original building. The entrance addition is Type V (111) construction, 2-stories without a basement and the hospital addition is Type II (111) construction, 1-story without a basement. In 2008 a 2-story building, without a basement, separated with two 2-hour fire barriers south of the entrance addition and was determined to be Type II (111) construction. The buildings are divided into 12 smoke zones (6 per floor) by 2- hour and 30 minute fire barriers.</p> <p>The facility is completely protected with an</p>	K 000			

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE

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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.

DEPARTMENT OF HEALTH AND HUMAN SERVICES
CENTERS FOR MEDICARE & MEDICAID SERVICES

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K 000	<p>Continued From page 1</p> <p>automatic fire sprinkler system in accordance with NFPA 13 Standard for the Installation of Sprinkler Systems 1999 edition with 2 systems. The facility has a fire alarm system with manual pull station near each exit door, smoke detection in the corridor system properly spaced and all common areas in accordance with NFPA 72 "The National Fire Alarm Code" (1999 edition). The fire alarm system is monitored for automatic fire department notification. Hazardous areas have either heat detection or smoke detection that are on the fire alarm system in accordance with the Minnesota State Fire Code (2007 edition).</p> <p>The facility has a capacity of 96 beds and had a census of 80 at the time of the survey.</p> <p>The requirement at 42 CFR, Subpart 483.70(a) is MET.</p>	K 000		



Protecting, Maintaining and Improving the Health of Minnesotans

Electronically submitted
July 24, 2015

Ms. Christy Brinkman, Administrator
Essentia Health Oak Crossing
1040 Lincoln Avenue
Detroit Lakes, Minnesota 56501

Re: Enclosed State Nursing Home Licensing Orders - Project Number S5212024

Dear Ms. Brinkman:

The above facility was surveyed on July 7, 2015 through July 10, 2015 for the purpose of assessing compliance with Minnesota Department of Health Nursing Home Rules. At the time of the survey, the survey team from the Minnesota Department of Health, Health Regulation Division, noted one or more violations of these rules that are issued in accordance with Minnesota Stat. section 144.653 and/or Minnesota Stat. Section 144A.10. If, upon reinspection, it is found that the deficiency or deficiencies cited herein are not corrected, a civil fine for each deficiency not corrected shall be assessed in accordance with a schedule of fines promulgated by rule of the Minnesota Department of Health.

To assist in complying with the correction order(s), a "suggested method of correction" has been added. This provision is being suggested as one method that you can follow to correct the cited deficiency. Please remember that this provision is only a suggestion and you are not required to follow it. Failure to follow the suggested method will not result in the issuance of a penalty assessment. You are reminded, however, that regardless of the method used, correction of the deficiency within the established time frame is required. The "suggested method of correction" is for your information and assistance only.

You have agreed to participate in the electronic receipt of State licensure orders consistent with the Minnesota Department of Health Informational Bulletin 14-01, available at <http://www.health.state.mn.us/divs/fpc/profinfo/infobul.htm>. The State licensing orders are delineated on the attached Minnesota Department of Health orders being submitted to you electronically. The Minnesota Department of Health is documenting the State Licensing Correction Orders using federal software. Tag numbers have been assigned to Minnesota state statutes/rules for Nursing Homes.

The assigned tag number appears in the far left column entitled "ID Prefix Tag." The state statute/rule number and the corresponding text of the state statute/rule out of compliance is listed in the "Summary Statement of Deficiencies" column and replaces the "To Comply" portion of the correction order. This column also includes the findings that are in violation of the state statute after the statement, "This Rule

is not met as evidenced by." Following the surveyors findings are the Suggested Method of Correction and the Time Period For Correction.

PLEASE DISREGARD THE HEADING OF THE FOURTH COLUMN WHICH STATES, "PROVIDER'S PLAN OF CORRECTION." THIS APPLIES TO FEDERAL DEFICIENCIES ONLY. THIS WILL APPEAR ON EACH PAGE.

THERE IS NO REQUIREMENT TO SUBMIT A PLAN OF CORRECTION FOR VIOLATIONS OF MINNESOTA STATE STATUTES/RULES.

Although no plan of correction is necessary for State Statutes/Rules, please enter the word "corrected" in the box available for text. You must then indicate in the electronic State licensure process, under the heading completion date, the date your orders will be corrected prior to electronically submitting to the Minnesota Department of Health. We urge you to review these orders carefully, item by item, and if you find that any of the orders are not in accordance with your understanding at the time of the exit conference following the survey, you should contact:

Gail Anderson, Unit Supervisor
Minnesota Department of Health
1505 Pebble Lake Road #300
Fergus Falls, Minnesota 56537
gail.anderson@state.mn.us
Telephone: (218) 332-5140
Fax: (218) 332-5196

You may request a hearing on any assessments that may result from non-compliance with these orders provided that a written request is made to the Department within 15 days of receipt of a notice of assessment for non-compliance.

Please note it is your responsibility to share the information contained in this letter and the results of this visit with the President of your facility's Governing Body.

Please feel free to call me with any questions.

Sincerely,



Kamala Fiske-Downing, Program Specialist
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Kamala.Fiske-Downing@state.mn.us
Telephone: (651) 201-4112 Fax: (651) 215-9697

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00907	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____	(X3) DATE SURVEY COMPLETED 07/10/2015
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2 000	<p>Initial Comments</p> <p>*****ATTENTION*****</p> <p>NH LICENSING CORRECTION ORDER</p> <p>In accordance with Minnesota Statute, section 144A.10, this correction order has been issued pursuant to a survey. If, upon reinspection, it is found that the deficiency or deficiencies cited herein are not corrected, a fine for each violation not corrected shall be assessed in accordance with a schedule of fines promulgated by rule of the Minnesota Department of Health.</p> <p>Determination of whether a violation has been corrected requires compliance with all requirements of the rule provided at the tag number and MN Rule number indicated below. When a rule contains several items, failure to comply with any of the items will be considered lack of compliance. Lack of compliance upon re-inspection with any item of multi-part rule will result in the assessment of a fine even if the item that was violated during the initial inspection was corrected.</p> <p>You may request a hearing on any assessments that may result from non-compliance with these orders provided that a written request is made to the Department within 15 days of receipt of a notice of assessment for non-compliance.</p> <p>INITIAL COMMENTS: You have agreed to participate in the electronic receipt of State licensure orders consistent with the Minnesota Department of Health Informational Bulletin 14-01, available at http://www.health.state.mn.us/divs/fpc/profinfo/infobul.htm The State licensing orders are delineated on the attached Minnesota</p>	2 000		

Minnesota Department of Health
LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE

TITLE

(X6) DATE

Electronically Signed

07/27/15

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2 000	<p>Continued From page 1</p> <p>Department of Health orders being submitted to you electronically. Although no plan of correction is necessary for State Statutes/Rules, please enter the word "corrected" in the box available for text. You must then indicate in the electronic State licensure process, under the heading completion date, the date your orders will be corrected prior to electronically submitting to the Minnesota Department of Health.</p> <p>On 7/7, 7/8, 7/9 and 7/10/15 surveyors of this Department's staff, visited the above provider and the following correction orders are issued. Please indicate in your electronic plan of correction that you have reviewed these orders, and identify the date when they will be completed.</p> <p>Minnesota Department of Health is documenting the State Licensing Correction Orders using federal software. Tag numbers have been assigned to Minnesota state statutes/rules for Nursing Homes.</p> <p>The assigned tag number appears in the far left column entitled "ID Prefix Tag." The state statute/rule out of compliance is listed in the "Summary Statement of Deficiencies" column and replaces the "To Comply" portion of the correction order. This column also includes the findings which are in violation of the state statute after the statement, "This Rule is not met as evidence by." Following the surveyors findings are the Suggested Method of Correction and Time period for Correction.</p> <p>PLEASE DISREGARD THE HEADING OF THE FOURTH COLUMN WHICH STATES, "PROVIDER'S PLAN OF CORRECTION." THIS APPLIES TO FEDERAL DEFICIENCIES ONLY. THIS WILL APPEAR ON EACH PAGE.</p>	2 000		

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2 540	<p>THERE IS NO REQUIREMENT TO SUBMIT A PLAN OF CORRECTION FOR VIOLATIONS OF MINNESOTA STATE STATUTES/RULES.</p> <p>MN Rule 4658.0400 Subp. 1 & 2 Comprehensive Resident Assessment</p> <p>Subpart 1. Assessment. A nursing home must conduct a comprehensive assessment of each resident's needs, which describes the resident's capability to perform daily life functions and significant impairments in functional capacity. A nursing assessment conducted according to Minnesota Statutes, section 148.171, subdivision 15, may be used as part of the comprehensive resident assessment. The results of the comprehensive resident assessment must be used to develop, review, and revise the resident's comprehensive plan of care as defined in part 4658.0405.</p> <p>Subp. 2. Information gathered. The comprehensive resident assessment must include at least the following information:</p> <ul style="list-style-type: none"> A. medically defined conditions and prior medical history; B. medical status measurement; C. physical and mental functional status; D. sensory and physical impairments; E. nutritional status and requirements; F. special treatments or procedures; G. mental and psychosocial status; H. discharge potential; I. dental condition; J. activities potential; K. rehabilitation potential; L. cognitive status; M. drug therapy; and N. resident preferences. 	2 540		8/14/15

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2 540	<p>Continued From page 3</p> <p>This MN Requirement is not met as evidenced by: Based on interview and document review, the facility failed to ensure the Minimum Data Set (MDS) accurately reflected the pressure ulcer status for 1 of 2 residents (R60) identified with pressure ulcers.</p> <p>Findings include:</p> <p>R60's admission MDS dated 3/6/15, identified R60 was at risk for development of pressure ulcer, however did not identify R60 had a unhealed pressure ulcer and did not identify the presence of a stage 2 pressure ulcer upon admission.</p> <p>R60's Physician Order Report dated 2/27/15 to 7/10/15 identified the following diagnoses: surgery aftercare, malignant brain and breast cancer, neuropathy and urinary incontinency.</p> <p>Review of R60's admission body observation dated 2/28/15, identified R60's right buttock had a pressure ulcer present upon admission and both buttocks were discolored from previous sores.</p> <p>Review of R60's Tissue Tolerance assessment dated 3/1/15, indicated R60 continued to have a pressure sore on left buttock, open area small eraser size</p> <p>Review of R60's Resident progress noted dated 3/2/15 indicated healing stage 2 area on right buttock.</p> <p>R60's pressure ulcer Care Area Assessment (CAA) dated 3/6/15, identified R60 was at risk for development of pressure ulcer, required</p>	2 540	Completed	

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2 540	<p>Continued From page 4</p> <p>extensive assistance with bed mobility and required a special mattress or seat cushion to reduce or relieve pressure. However, the CAA did not identify the presence of R60's unhealed stage 2 pressure ulcer and lacked an analysis of current treatments and interventions for the unhealed stage 2 pressure ulcer.</p> <p>On 7/10/15, at 6:01 p.m. the director of nursing stated she expected all the skin issues should have been thoroughly assessed and documented during admission. DON confirmed R60's admission MDS and CAA and stated she expected the MDS and CAA to have identified R60's pressure ulcer upon admission.</p> <p>The facility's Care Conferences, Care Planning policy dated 4/1/15, directed staff to include the following sources of information for the MDS:</p> <ul style="list-style-type: none"> -Review of the residents medical record -Observation of the resident -Interviews and communication with the resident -Communication with healthcare providers -Communication with physician -Communication with the resident's family members <p>SUGGESTED METHOD FOR CORRECTION: The director of nursing (DON) and/or designee could review policy and provide education for staff regarding completion of an individualized comprehensive resident assessment. The Quality Assessment and Assurance (QAA) committee could do random audits to ensure compliance.</p> <p>TIME PERIOD FOR CORRECTION: Twenty-one</p>	2 540		

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2 540	Continued From page 5 (21) days.	2 540		
21015	<p>MN Rule 4658.0610 Subp. 7 Dietary Staff Requirements- Sanitary conditi</p> <p>Subp. 7. Sanitary conditions. Sanitary procedures and conditions must be maintained in the operation of the dietary department at all times.</p> <p>This MN Requirement is not met as evidenced by: Based on observation, Interview and record review the facility failed to clean and maintain food service equipment to prevent food borne illness for residents residing in 2 of 4 neighborhoods (Cedar Ridge, Meadowbrook) in the facility.</p> <p>Findings include:</p> <p>On 07/10/15, at 4:24 p.m. observations of the Meadow Brook neighborhood ice machine were conducted with Nursing Home Support (NHS-A) present. On the front of ice machine, a black spout with a pink substance inside the opening of the spout was observed. NHS-A reached out with a disposable cloth and wiped up inside the black spout covers. The disposable cloth had both brown sludge smears and pink smudges on the cloth. In addition, there were several white hard water scale stains observed covering the upper right 1/3 of drip tray, and covered the top 3 grates of the drip tray. NHS-A stated she was aware the ice machine had a build up of pink substance in the past and indicated an outside company cleaned and serviced the ice machines for the facility. She indicated homemakers were to</p>	21015	Completed	8/14/15

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21015	<p>Continued From page 6</p> <p>routinely clean the outside of the ice machines, however, cleaning the spouts and the inside of the spouts were not the usual practice for facility staff. NHS-A confirmed she was aware the ice machine had a build up of the pink substance and stated she had not routinely cleaned the spouts of the ice machine.</p> <p>On 07/10/15, at 4:56 p.m. the director of dietary (DD) stated the facility utilized an outside company to service and clean the ice machines. She indicated facility staff did not service or maintain the equipment and indicated the outside company came to facility twice a year to service and clean the ice machines. DD stated the "Daily Homemaker Cleaning Duties Record" utilized by facility staff identified the clean the beverage area, however, did not identify what parts of the ice machine to clean and how to clean the ice machine. She confirmed the findings and indicated she had not been aware of the sludge and pink substance in the ice machines. She stated "that is not good" and indicated she felt there could be a risk for food borne illness.</p> <p>On 07/10/15, at 5:06 p.m. observations of the Cedar Ridge neighborhood ice machine were conducted with Nursing Assistant (NA-A) present. NA-A reached out with a white paper napkin and wiped the inside of the black spout of the ice/water dispenser. Upon removal of the white paper napkin from the black spout, two areas of pink sludge were noted on the napkin. Each area of pink sludge was 2 cm (centimeters) wide by 2 cm long. NA-A verified the two areas of pink sludge and indicated she routinely cleaned the outside of the ice machine and tray every day and stated she did not routinely clean the spouts or</p>	21015		

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21015	Continued From page 7 inside of the spouts. Review of the most current invoice from the outside service company dated 2/12/15, identified The machine was cleaned and sanitized and inspected and checked for proper operation." Review of the Manufacturer's cleaning and maintenance instructions, provided by the facility, revealed the manufacturer recommended cleaning and sanitizing the ice machine at least twice a year. The instructions also identified more frequent cleaning may be required in some existing water conditions. In addition, the instructions included monthly cleaning instructions for the exterior of the ice machine included using a neutral cleaner to wipe off oil or dirt build up. Clean any chlorine staining (rust or colored spots) using a non-abrasive cleaner. SUGGESTED METHOD FOR CORRECTION: The director of nursing (DON) and/or designee could review policy and provide education for staff regarding routine cleaning and sanitation of the dietary equipment including the ice machines. The Quality Assessment and Assurance (QAA) committee could do random audits to ensure compliance. TIME PERIOD FOR CORRECTION: Twenty-one (21) days.	21015		
21390	MN Rule 4658.0800 Subp. 4 A-I Infection Control Subp. 4. Policies and procedures. The infection control program must include policies and procedures which provide for the following: A. surveillance based on systematic data	21390		8/14/15

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21390	<p>Continued From page 8</p> <p>collection to identify nosocomial infections in residents;</p> <p>B. a system for detection, investigation, and control of outbreaks of infectious diseases;</p> <p>C. isolation and precautions systems to reduce risk of transmission of infectious agents;</p> <p>D. in-service education in infection prevention and control;</p> <p>E. a resident health program including an immunization program, a tuberculosis program as defined in part 4658.0810, and policies and procedures of resident care practices to assist in the prevention and treatment of infections;</p> <p>F. the development and implementation of employee health policies and infection control practices, including a tuberculosis program as defined in part 4658.0815;</p> <p>G. a system for reviewing antibiotic use;</p> <p>H. a system for review and evaluation of products which affect infection control, such as disinfectants, antiseptics, gloves, and incontinence products; and</p> <p>I. methods for maintaining awareness of current standards of practice in infection control.</p> <p>This MN Requirement is not met as evidenced by: Based on observation, interview and document review the facility failed to ensure proper infection control practices for expired disinfectant wipes used to sanitize multi-use glucometers, on 2 of 4 facility resident communities, Harbor Springs and Meadow Brook. This deficient practice had the potential to affect 10 of 10 residents (R18, R134, R114, R35, R58, R73, R71, R75, R111, R125) currently using the multi-used glucometers.</p> <p>Findings include:</p>	21390	Completed	

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21390	<p>Continued From page 9</p> <p>Observation of the Harbor Springs medication cart and storage room was conducted on 7/9/15, at 12:56 p.m. with licensed practical nurse (LPN)-A present. The medication/ treatment cart contained 1 open box of 50 Sani-wipes with less than 10 packages missing and an expiration date of 4/2015.</p> <p>The storage room contained 1/2 box of 50 sani-wipes with an expiration dated of 4/2015, and a second 1/2 box of 50 sani-wipes with expiration dated of 5/2015.</p> <p>During an interview on 7/09/201, at 12:56 p.m. LPN-A verified the above findings. LPN-A indicated the expired Sani-wipes were currently being used to clean the medication cart and the multi patient use glucometer. LPN-A stated, "three residents" currently used the shared glucometer at this time. LPN-A indicated she did not routinely check for expiration dates of the sani-wipes.</p> <p>During an interview on 7/10/2015, at 12:24 p.m. registered nurse (RN)-C verified the usual facility practice was to disinfect the multi use glucometers with the Sani-wipes. RN-C verified the use of expired Sani-wipes for sanitation of multi patient use glucometers would not be an acceptable infection control process. RN-C stated, " It would not be known if the expired wipes had the ability to sanitize."</p> <p>During review of the Meadow Brook medication cart and storage rooms on 7/10/2015, at 12:35 p.m. with LPN-B, the following was found: The medication/ treatment cart contained 21 of 22 Sani-wipes with an expiration date of 5/2015, The supply storage room contained 1 full box of 50 sani-wipes with expiration date of 5/2015. The medication storage room contained a box</p>	21390		

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21390	<p>Continued From page 10</p> <p>with 10 sani-wipes remaining in the box with an expiration date of 5/2015.</p> <p>During an interview on 7/10/2015, at 12:35 p.m. LPN-B. verified the above findings and indicated the expired wipes were currently utilized to disinfect the multi-use glucometers in the facility. LPN-B indicated she did not know where the expiration dates were printed on the Sani-wipe box and individual packages, and stated she did not check the expiration dates. LPN-B verified 10 residents currently used the multi used glucometer (R18, R134, R114, R35, R58,R73, R71, R75, R111, R125).</p> <p>During an interview on 7/10/2015, at 5:13 p.m. the director of nursing (DON) confirmed the current facility policy and indicated the use of expired Sani-wipes for multi use glucometers would not be an acceptable facility practice.</p> <p>The facility provided manufacturers instructions titled Nova Biomedical Customer Information Bulletin, dated 12/4/12, directed to disinfect the meter" #1 Using a new, fresh germicidal wipe."</p> <p>SUGGESTED METHOD FOR CORRECTION: The director of nursing (DON) and/or designee could review policy and provide education for staff regarding proper infection control practices which included proper disinfection of the multi use glucometers. The Quality Assessment and Assurance (QAA) committee could do random audits to ensure compliance.</p> <p>TIME PERIOD FOR CORRECTION: Twenty-one (21) days.</p>	21390		

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21426	Continued From page 11	21426		
21426	<p>MN St. Statute 144A.04 Subd. 3 Tuberculosis Prevention And Control</p> <p>(a) A nursing home provider must establish and maintain a comprehensive tuberculosis infection control program according to the most current tuberculosis infection control guidelines issued by the United States Centers for Disease Control and Prevention (CDC), Division of Tuberculosis Elimination, as published in CDC's Morbidity and Mortality Weekly Report (MMWR). This program must include a tuberculosis infection control plan that covers all paid and unpaid employees, contractors, students, residents, and volunteers. The Department of Health shall provide technical assistance regarding implementation of the guidelines.</p> <p>(b) Written compliance with this subdivision must be maintained by the nursing home.</p> <p>This MN Requirement is not met as evidenced by: Based on interview and record review the facility failed to ensure a Tuberculin (TB) baseline screening had been completed for 5 of 5 residents (R39,R72,R195, R193, R66) upon admission. In addition, the facility failed to document the interpretation of the Tuberculosis Skin Test (TST) for 1 of 5 residents (R66) and for 1 of 5 newly hired employees(E1) reviewed for Tuberculosis (TB) program.</p> <p>Findings include:</p>	21426	Corrected	8/14/15

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21426	<p>Continued From page 12</p> <p>R39 was admitted to the facility in 2015. Review of the R39's immunization record revealed the TB baseline symptom screening form had not been completed upon admission to the facility.</p> <p>R72 was admitted to the facility in 2015. Review of R72's immunization record revealed the TB baseline symptom screening form had not been completed upon admission to the facility.</p> <p>R195 was admitted to the facility in 2015. Review of R195's immunization record revealed the TB baseline symptom screening form had not been completed upon admission to the facility.</p> <p>R193 was admitted to the facility in 2015. Review of R193's immunization record revealed the TB baseline symptom screening form had not been completed upon admission to the facility.</p> <p>R66 was admitted to the facility in 2015. Review of the R66's immunization record revealed the TB baseline symptom screening form had not been completed. In addition, the immunization record revealed R66's second TST(tuberculin skin test) was given on 7/1/15, but no interpretation of the results for the second TST was documented.</p> <p>E1 was a newly hired employee. The second TST was given on 1/14/15, and was read on 1/16/15 with a result documented as negative. At the time of review, there was no documentation of induration. After interview on 7/10/15, 0 mm induration was added.</p> <p>During interview on 7/10/15, at 1:07 p.m. the director of nursing (DON) confirmed all residents and newly hired employees were required to have</p>	21426		

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21426	Continued From page 13 the TB baseline symptom screening completed. The DON stated the facility had thought their electronic medical record system had covered the symptom screening, but confirmed it had not. The DON also confirmed all TST results must have the induration documented with the result, not only negative. Review of the facility policy titled TB Control Plan, revised 5/15, revealed TB symptom screening and skin testing would be completed for all newly hired employees. SUGGESTED METHOD OF CORRECTION: The infection control nurse or designee could review the TB policies and procedures to ensure required information is included. Appropriate staff could be educated regarding requirements. Audits could be conducted and the results reviewed at the quality committee meetings. TIME PERIOD FOR CORRECTION: Twenty-one (21) days.	21426		
21565	MN Rule 4658.1325 Subp. 4 Administration of Medications Self Admin Subp. 4. Self-administration. A resident may self-administer medications if the comprehensive resident assessment and comprehensive plan of care as required in parts 4658.0400 and 4658.0405 indicate this practice is safe and there is a written order from the attending physician. This MN Requirement is not met as evidenced by: Based on observation, interview, and document	21565	Completed	8/14/15

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21565	<p>Continued From page 14</p> <p>review, the facility failed to ensure a compressive self administration medication assessment was completed to determine ability of safe self medication administration for 1 of 1 residents (R196) observed to self administer medications during the survey.</p> <p>Findings include:</p> <p>During a medication pass observation on 7/7/15, at 5:38 p.m. licensed practical nurse (LPN)-A dispensed (a blood pressure medication) metoprolol 100 mg (milligrams) into a white paper cup. LPN-A placed the white paper medication cup on the dining table next to R196, and walked back to medication cart. LPN- A signed the medication as given on the electronic medication administration form and then proceeded with her duties. LPN-A did not observe or return to R196 during the entire observation. At 5:53 p.m. LPN-A left the dining room and pushed the medication cart down a hall of the facility, out of view of R196.</p> <p>R196's admission Minimum Data Set (MDS) care area assessment (CAA), dated 6/30/15, identified R196 had severe cognitive impairment, with diagnoses which included dementia, had confusion, disorientation, forgetfulness, difficulty with recall and short and long term memory.</p> <p>During an interview on 7/07/2015, at 6:00 p.m. LPN-A stated, " [R196] can manage meds (medications) a little bit." LPN-A indicated she felt R196 would take the medications independently. LPN-A indicated a visitor was always with R196; therefore, no follow up was needed to ensure R196 received the medication.</p> <p>During an interview on 7/09/2015, at 8:56 a.m.</p>	21565		

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21565	<p>Continued From page 15</p> <p>LPN-B verified the facility practice of residents who self administer medications have initial assessments, a physician order, and were reviewed monthly for continued appropriateness of self medication administration.</p> <p>Review of R196's electronic and paper chart identified the following: The current physician's orders dated 6/10/15 - 7/10/15 identified an order for metoprolol 100 mg, oral twice a day. However, the record lacked a physician's order for the resident to self administer medications. The record lacked documentation a self administration of medication assessment had been completed, to determine if R196 was safe to self administer medications. R196s' care plan revised 7/8/15, did not identify R196 was safe to self administer medications.</p> <p>During an interview on 7/10/2015, 5:13 p.m. the director of nursing (DON) verified a resident self administration assessment was required to be completed in order for medications to be left at a residents side. The DON stated "the [facility] policy would prohibit" self administration without an assessment for safety completed.</p> <p>The undated facility policy titled Routine Orders identified, Self-Administration of Medications identified "May initiate after assessment and care plan.</p> <p>SUGGESTED METHOD OF CORRECTION: The director of nursing and/or their designee should review the facility's policy and procedures and educate the facility staff responsible for the provision of self administration of medications. Audits could be conducted and the</p>	21565		

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21565	Continued From page 16 results reviewed at the quality committee meetings. TIME PERIOD FOR CORRECTION: Twenty-one (21) days.	21565		