



Protecting, Maintaining and Improving the Health of Minnesotans

CMS Certification Number (CCN): 245370

February 9, 2015

Mr. Nathan Johnson, Administrator
Ecumen North Branch
5379 -383rd Street
North Branch, Minnesota 55056

Dear Mr. Johnson:

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 December 30, 2014 the above facility is certified for:

67 Skilled Nursing Facility/Nursing Facility Beds

Your facility's Medicare approved area consists of all 67 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
mark.meath@state.mn.us
Telephone: (651) 201-4118
Fax: (651) 215-9697



Protecting, Maintaining and Improving the Health of Minnesotans

Electronically delivered

February 9, 2015

Mr. Nathan Johnson, Administrator
Ecumen North Branch
5379 -383rd Street
North Branch, Minnesota 55056

RE: Project Number S5370030

Dear Mr. Johnson:

On November 25, 2014, we informed you that we would recommend enforcement remedies based on the deficiencies cited by this Department for a standard survey, completed on November 20, 2014. This survey found the most serious deficiencies to be widespread deficiencies that constituted no actual harm with potential for more than minimal harm that was not immediate jeopardy (Level F), whereby corrections were required.

On January 5, 2015, the Minnesota Department of Health completed a Post Certification Revisit (PCR) by review of your plan of correction and on January 6, 2015 the Minnesota Department of Public Safety completed a PCR to verify that your facility had achieved and maintained compliance with federal certification deficiencies issued pursuant to a standard survey, completed on November 20, 2014. We presumed, based on your plan of correction, that your facility had corrected these deficiencies as of December 30, 2014. Based on our PCR, we have determined that your facility has corrected the deficiencies issued pursuant to our standard survey, completed on November 20, 2014, effective January 5, 2015 and therefore remedies outlined in our letter to you dated November 25, 2014, 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
mark.meath@state.mn.us

Telephone: (651) 201-4118

Fax: (651) 215-9697

5370r15

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 245370	(Y2) Multiple Construction A. Building B. Wing	(Y3) Date of Revisit 1/5/2015
Name of Facility ECUMEN NORTH BRANCH	Street Address, City, State, Zip Code 5379 -383RD STREET NORTH BRANCH, MN 55056	

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 F0322 Reg. # 483.25(a)(2) LSC _____	Correction Completed 01/05/2015	ID Prefix F0441 Reg. # 483.65 LSC _____	Correction Completed 01/05/2015	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
ID Prefix _____ Reg. # _____ LSC _____	Correction Completed	ID Prefix _____ Reg. # _____ LSC _____	Correction Completed	ID Prefix _____ Reg. # _____ LSC _____	Correction Completed

Reviewed By _____ State Agency	Reviewed By CC/mm	Date: 02/09/2015	Signature of Surveyor: 13922	Date: 01/05/2015
Reviewed By _____ CMS RO	Reviewed By	Date:	Signature of Surveyor:	Date:

Followup to Survey Completed on: 11/20/2014	Check for any Uncorrected Deficiencies. Was a Summary of Uncorrected Deficiencies (CMS-2567) Sent to the Facility? <table style="float: right; margin-left: 20px;"> <tr> <td>YES</td> <td>NO</td> </tr> </table>	YES	NO
YES	NO		

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 245370	(Y2) Multiple Construction A. Building 02 - BLDG 2 B. Wing	(Y3) Date of Revisit 1/6/2015
Name of Facility ECUMEN NORTH BRANCH	Street Address, City, State, Zip Code 5379 -383RD STREET NORTH BRANCH, MN 55056	

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 _____ Reg. # NFPA 101 LSC K0062	Correction Completed 12/30/2014	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	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 _____	Reviewed By PS/mm	Date: 02/09/2015	Signature of Surveyor: 03005	Date: 01/06/2015
Reviewed By _____	Reviewed By	Date:	Signature of Surveyor:	Date:

Followup to Survey Completed on: 11/18/2014	Check for any Uncorrected Deficiencies. Was a Summary of Uncorrected Deficiencies (CMS-2567) Sent to the Facility? YES NO
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MEDICARE/MEDICAID CERTIFICATION AND TRANSMITTAL
PART I - TO BE COMPLETED BY THE STATE SURVEY AGENCY

ID: KC1L
Facility ID: 00066

1. MEDICARE/MEDICAID PROVIDER NO. (L1) 245370 2. STATE VENDOR OR MEDICAID NO. (L2) 533840900	3. NAME AND ADDRESS OF FACILITY (L3) ECUMEN NORTH BRANCH (L4) 5379 -383RD STREET (L5) NORTH BRANCH, MN (L6) 55056	4. TYPE OF ACTION: <u>2</u> (L8) 1. Initial 2. Recertification 3. Termination 4. CHOW 5. Validation 6. Complaint 7. On-Site Visit 9. Other 8. Full Survey After Complaint															
5. EFFECTIVE DATE CHANGE OF OWNERSHIP (L9) 6. DATE OF SURVEY 11/20/2014 (L34) 8. ACCREDITATION STATUS: <u> </u> (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	FISCAL YEAR ENDING DATE: (L35) 12/31															
11. LTC PERIOD OF CERTIFICATION From (a) : To (b) : 12. Total Facility Beds 67 (L18) 13. Total Certified Beds 67 (L17)	10. THE FACILITY IS CERTIFIED AS: A. In Compliance With Program Requirements Compliance Based On: <u> </u> 1. Acceptable POC X B. Not in Compliance with Program Requirements and/or Applied Waivers: * Code: B* (L12) And/Or Approved Waivers Of The Following Requirements: <u> </u> 2. Technical Personnel <u> </u> 6. Scope of Services Limit <u> </u> 3. 24 Hour RN <u> </u> 7. Medical Director <u> </u> 4. 7-Day RN (Rural SNF) <u> </u> 8. Patient Room Size <u> </u> 5. Life Safety Code <u> </u> 9. Beds/Room																
14. LTC CERTIFIED BED BREAKDOWN <table style="width:100%; border: none;"> <tr> <td style="text-align: center;">18 SNF</td> <td style="text-align: center;">18/19 SNF</td> <td style="text-align: center;">19 SNF</td> <td style="text-align: center;">ICF</td> <td style="text-align: center;">IID</td> </tr> <tr> <td style="text-align: center;">67</td> <td></td> <td></td> <td></td> <td></td> </tr> <tr> <td style="text-align: center;">(L37)</td> <td style="text-align: center;">(L38)</td> <td style="text-align: center;">(L39)</td> <td style="text-align: center;">(L42)</td> <td style="text-align: center;">(L43)</td> </tr> </table>		18 SNF	18/19 SNF	19 SNF	ICF	IID	67					(L37)	(L38)	(L39)	(L42)	(L43)	15. FACILITY MEETS 1861 (e) (1) or 1861 (j) (1): (L15)
18 SNF	18/19 SNF	19 SNF	ICF	IID													
67																	
(L37)	(L38)	(L39)	(L42)	(L43)													
16. STATE SURVEY AGENCY REMARKS (IF APPLICABLE SHOW LTC CANCELLATION DATE):																	
17. SURVEYOR SIGNATURE <u>Cheryl J Johnson, HFE NE II</u> Date : 12/30/2014 (L19)	18. STATE SURVEY AGENCY APPROVAL <u>Anne Kleppe, Enforcement Specialist</u> Date: 01/20/2015 (L20)																

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

19. DETERMINATION OF ELIGIBILITY <u> </u> 1. Facility is Eligible to Participate <u> </u> 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 12/01/1986 (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 <u>00</u> 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: (L28)	29. INTERMEDIARY/CARRIER NO. 03001 (L31)	
31. RO RECEIPT OF CMS-1539 (L32)	30. REMARKS Posted 01/20/2015 Co. DETERMINATION APPROVAL	
32. DETERMINATION OF APPROVAL DATE (L33)		



Protecting, Maintaining and Improving the Health of Minnesotans

Electronically delivered
November 25, 2014

Mr. Nathan Johnson, Administrator
Ecumen North Branch
5379 -383rd Street
North Branch, Minnesota 55056

RE: Project Number S5370030

Dear Mr. Johnson:

On November 20, 2014, 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 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 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, Supervisor
Fergus Falls Survey Team
Licensing and Certification Program
Division of Compliance Monitoring
Minnesota Department of Health
Gail.anderson@state..mn.us**

Phone: (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 December 30, 2014, the Department of Health will impose the following remedy:

- State Monitoring. (42 CFR 488.422)

In addition, the Department of Health is recommending to the CMS Region V Office that if your facility has not achieved substantial compliance by December 30, 2014 the following remedy will be imposed:

- Per instance civil money penalties. (42 CFR 488.430 through 488.444)

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.

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 February 20, 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 May 20, 2015 (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
Division of Compliance Monitoring
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/ltc/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: <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.

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. Patrick Sheehan, Supervisor
Health Care Fire Inspections
State Fire Marshal Division
pat.sheehan@state.mn.us
Telephone: (651) 201-7205
Fax: (651) 215-0525

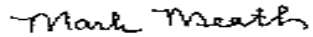
Ecumen North Branch

November 25, 2014

Page 6

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

Sincerely,

A handwritten signature in black ink that reads "Mark Meath". The signature is written in a cursive style.

Mark Meath, Enforcement Specialist
Program Assurance Unit
Licensing and Certification Program
Division of Compliance Monitoring
mark.meath@state.mn.us

Telephone: (651) 201-4118

Fax: (651) 215-9697

cc: Licensing and Certification File

5370s15

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

PRINTED: 12/30/2014
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245370	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED 11/20/2014
NAME OF PROVIDER OR SUPPLIER ECUMEN NORTH BRANCH			STREET ADDRESS, CITY, STATE, ZIP CODE 5379 -383RD STREET NORTH BRANCH, MN 55056		
(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 322 SS=D	CENSUS: 64 483.25(g)(2) NG TREATMENT/SERVICES - RESTORE EATING SKILLS Based on the comprehensive assessment of a resident, the facility must ensure that -- (1) A resident who has been able to eat enough alone or with assistance is not fed by naso gastric tube unless the resident ' s clinical condition demonstrates that use of a naso gastric tube was unavoidable; and (2) A resident who is fed by a naso-gastric or gastrostomy tube receives the appropriate treatment and services to prevent aspiration pneumonia, diarrhea, vomiting, dehydration, metabolic abnormalities, and nasal-pharyngeal ulcers and to restore, if possible, normal eating skills.	F 322		12/30/14	

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

TITLE

(X6) DATE

Electronically Signed

12/29/2014

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.

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245370	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED 11/20/2014
NAME OF PROVIDER OR SUPPLIER ECUMEN NORTH BRANCH			STREET ADDRESS, CITY, STATE, ZIP CODE 5379 -383RD STREET NORTH BRANCH, MN 55056		
(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 322	Continued From page 1 This REQUIREMENT is not met as evidenced by: Based on observation, interview and document review, the facility failed to check placement of the jejunostomy/gastrostomy tube (JG-tube, one tube to the jejunum [small intestine] for feeding and one tube into the stomach for medications) prior to medication administration for 1 of 1 residents (R94) observed for medication administration through a G-tube. Findings include: R94's Admission Record dated 11/14, indicated R94's diagnoses included pneumonia due to the inhalation of food or vomit, sepsis, adult failure to thrive, malnutrition, weakness, dysphagia (difficulty swallowing), nausea with vomiting, gastrostomy and esophageal reflux. The Physician's Orders dated 11/7/14, directed all medications via G-tube. Licensed practical nurse (LPN)-A prepared three separate cups of liquid medication for administration via G-tube for R94 on 11/18/14, at 9:40 a.m.. LPN-A flushed the G-tube with water, administered each medication separately into the G-tube, and flushed the tubing with approximately 30 milliliters (ml) of water between each medication. LPN-A did not check for placement of the G-tube prior to administering the water and medications. LPN-A stated she did not check placement prior to giving the medication because she checked placement when she flushed the	F 322	1. Corrective Action: a. Review J/G-Tube medication administration policy/procedure with all licensed nurses with emphasis of checking for placement. 2. Corrective Action as it applies to Other Residents: a. Audit all current residents with J/G Tube medication administration 3. Reoccurrence will be Prevented by: a. Staff education occurred on November 24 and 25th but it is also set to re-occur on January 14 and 15th of 2015. Educate on the Policy/Procedure of J/G-Tube Medication Administration. b. The DON or designee will complete random weekly audits 3 times a week for 1 month, once weekly x2 months and once monthly x3 months. 4. The Correction will be Monitored by: a. DON or Designee 5. Date of Completion: December 30, 2014		

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NAME OF PROVIDER OR SUPPLIER ECUMEN NORTH BRANCH			STREET ADDRESS, CITY, STATE, ZIP CODE 5379 -383RD STREET NORTH BRANCH, MN 55056		
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F 322	Continued From page 2 tube with water after the feeding was finished, "At about 7:30 a.m." The director of nurses, interviewed on 11/20/14, at 8:40 a.m., stated R94 had a G/J-tube. The medications were administered through the G-tube and feedings through the J-tube. The DON verified placement of the G-tube should be checked prior to medication administration. The undated procedure for Administering Medications Per NG (nasal gastric) or Etenal Tube procedure directed insertion of a 50 ml syringe into the NG tube and aspirate to check residual volume. Return residual and flush NG tube. The rational was to validate gastric capacity for receiving medication and flush solution. A policy for administering medication through a JG-tube was requested but not provided.	F 322			
F 441 SS=D	483.65 INFECTION CONTROL, PREVENT SPREAD, LINENS 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. (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.	F 441		12/30/14	

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F 441	<p>Continued From page 3</p> <p>(b) Preventing Spread of Infection</p> <p>(1) When the Infection Control Program determines that a resident needs isolation to prevent the spread of infection, the facility must isolate the resident.</p> <p>(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.</p> <p>(3) The facility must require staff to wash their hands after each direct resident contact for which 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 hand washing was not completed during cares per facility policy for 1 of 4 residents (R73) observed for infection control. Findings include: The quarterly Minimum Data Set dated 10/1/14, indicated R73 needed the extensive assistance of one staff with transfers, personal hygiene, dressing and toilet use. On 11/19/14, at 8:15 a.m. R73's cares were observed with nursing assistant (NA)-A. NA-A sanitized her hands and donned gloves before gathering R73's supplies and clothing. NA-A transferred R73 with the mechanical stand aid from the bed to the bathroom, removed R73's incontinent brief, and lowered R73 on to the toilet.</p>	F 441	<p>1. Corrective Action: a.CNA who failed to wash hands after removing gloves during care has been educated.</p> <p>2. Corrective Action as it applies to Other Residents: a.All residents have the potential to be effected by these deficient practices. b.Staff members are being monitored routinely and educated on the spot when they fail to perform hand washing appropriately.</p> <p>3. Reoccurrence will be Prevented by: a.Staff education occurred on</p>	

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F 441	<p>Continued From page 4</p> <p>NA-A removed R73's gown, washed and dried R73's face and upper body. NA-A assisted R73 into a shirt, applied the lift sling and the incontinent brief. NA-A stood R73 with the stand aid and washed and dried R73's groin and buttocks. NA-A threaded the urinary drainage bag through the leg of the pants, pulled up the pants and transferred R73 into the wheelchair. Without hand sanitization or glove change, NA-A placed the urinary drainage bag into a pillow case and hung it under the chair. NA-A tied the trash and linen bags and then removed the gloves without washing hands. NA-A placed a new bag in the trash can, raised the window shade, applied a blanket to R73's shoulders, brushed R73's hair, picked up the linen and trash bags and exited the room, still without hand sanitization. NA-A moved the stand aid into the hall, walked to the soiled utility room and opened the door with the handle. NA-A, interviewed on 11/19/14, at 8:45 a.m., stated she she usually washes or sanitizes her hands before leaving the room but had the bags of soiled linen and trash in her hands. NA-A verified she did not remove the gloves and wash or sanitize hands after providing peri-care.</p> <p>On 11/20/14, at 8:40 a.m. the director of nursing (DON) stated, "Hands should be washed or sanitized after removing gloves, when going from dirty to clean and before leaving the room." The Infection Control Guidelines for All Nursing Procedures policy revised on 4/13 indicated employees must wash their hands for 10 to 15 seconds using antimicrobial or non-antimicrobial soap and water under the following conditions: after contact with blood, body fluids, secretions, mucous membranes or non intact skin. After removing gloves. In most situations the preferred method of hand</p>	F 441	<p>November 24 and 25th but is also set to re-occur on January 14 and 15th of 2015. Educate on the Policy/Procedure of Infection Control related to Handwashing and Glove Changing.</p> <p>b.The DON or designee will complete random weekly audits 3 times a week for 1 month, once weekly x2 months and once monthly x3 months.</p> <p>4. The Correction will be Monitored by: a.DON or Deisignee</p> <p>5. Date of Completion: December 30, 2014</p>		

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CENTERS FOR MEDICARE & MEDICAID SERVICES

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F 441	Continued From page 5 hygiene was with an alcohol based hand rub if the hands were not visibly soiled before and after direct contact with the residents; before moving from a contaminated body area to a clean body site during resident care; and after removing gloves.	F 441			

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NAME OF PROVIDER OR SUPPLIER ECUMEN NORTH BRANCH	STREET ADDRESS, CITY, STATE, ZIP CODE 5379 -383RD STREET NORTH BRANCH, MN 55056
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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 Ecumen Borth Branch was found not in substantial compliance with the requirements for participation in Medicare/Medicaid at 42 CFR, Subpart 483.70(a). Life Safety from Fire, and the 200 edition of National Fire Protection Association (NFPA) Standard 101, Life Safety Code (LSC) Chapter 18 New Health Care.</p> <p>PLEASE RETURN THE PLAN OF CORRECTION FOR THE FIRE SAFETY DEFICIENCIES TO:</p> <p>Health Care Fire Inspections STATE FIRE MARSHAL DIVISION 444 CEDAR ST., SUITE 145 ST. PAUL, MN 55101-514, or</p> <p>By E-Mail to: Marian.Whitney@state.mn.us</p> <p>THE PLAN OF CORRECTION FOR EACH DEFICIENCY MUST INCLUDE ALL OF THE FOLLOWING INFORMATION;</p> <ol style="list-style-type: none"> 1. A description of what has been, or will be, done to correct the deficiency. 2.. The actual, or proposed , completion date. 3. The name and/or title of the person responsible for correction and monitoring to 	K 000		
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LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE Electronically Signed	TITLE	(X6) DATE 12/29/2014
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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.

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K 000	Continued From page 1 prevent a reoccurrence of the deficiency. Ecumen North Branch was constructed in 2006 & 2007, with opening in 2007. It is a one story building with no basement. The construction type is determined to be type V(111). The building is separated from the rest of the facility by 2 hour fire rated construction , with a 1 & 1/2 hour rated fire doors. The building is fully sprinkler protected. The facility has a complete automatic sprinkler system, with smoke detection in the corridors and spaces open to the corridor, that is monitored for automatic fire department notification. All resident rooms have single station smoke detectors that transmit to the nurses station. The facility is licensed for 67 beds and 65 were occupied at the time of inspection.	K 000			
K 062 SS=F	The requirement at 42 CFR, Subpart 483.70(a)) is NOT met by evidenced by: NFPA 101 LIFE SAFETY CODE STANDARD Required automatic sprinkler systems are continuously maintained in reliable operating condition and are inspected and tested periodically. 18.7.6, 4.6.12, NFPA 13, NFPA 25, 9.7.5 This STANDARD is not met as evidenced by: This STANDARD is not met as evidenced by: Based on documentation review and interview with staff, the facility has failed to properly inspect and maintain the automatic sprinkler system in	K 062	1. Quarterly flow tests will be conducted by our contracted sprinkler system company. 2. On November 24th we signed a new contract to begin performing quarterly flow	12/30/14	

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K 062	<p>Continued From page 2</p> <p>accordance with NFPA 101 LSC (00) section 18.7.6, 4.6.12. This deficient practice does not ensure that the fire sprinkler system is functioning properly and is fully operational in the event of a fire and could negatively affect all of the builing occupants.</p> <p>Findings include:</p> <p>At the conclusion of the facility tour on 11-18-14 approximately 9:30AM, based on a review of the available fire sprinkler test and inspection documentation, observations revealed a), that the facility failed to provide documentation for any fire sprinkler flow tests of the facility's complete automatic fire sprinkler system.</p> <p>This deficient practice was verified by the Facility Administrator (NJ) and the Director of Facility Maintenance at the time of exit.</p>	K 062	<p>tests, as well as the annual.</p> <p>3. The Maintenance Director or desginee will be responsible. We will audit that the contracted sprinkler company did the inspections after their first two inspections since they are quarterly, as well as put the logs in our book.</p> <p>4. Completed 12/30/2014</p>		