

MEDICARE/MEDICAID CERTIFICATION AND TRANSMITTAL
PART I - TO BE COMPLETED BY THE STATE SURVEY AGENCY

ID: CQGF

Facility ID: 00763

1. MEDICARE/MEDICAID PROVIDER NO. (L1) 245524	3. NAME AND ADDRESS OF FACILITY (L3) LITTLE SISTERS OF THE POOR (L4) 330 EXCHANGE STREET SOUTH (L5) SAINT PAUL, MN (L6) 55102	4. TYPE OF ACTION: <u>7</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
2.STATE VENDOR OR MEDICAID NO. (L2) 825540700	5. EFFECTIVE DATE CHANGE OF OWNERSHIP (L9)	FISCAL YEAR ENDING DATE: (L35) 12/31
6. DATE OF SURVEY 03/18/2015 (L34)	7. PROVIDER/SUPPLIER CATEGORY <u>03</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	
8. ACCREDITATION STATUS: <u> </u> (L10) 0 Unaccredited 1 TJC 2 AOA 3 Other		

11. LTC PERIOD OF CERTIFICATION From (a) : To (b) :	10.THE FACILITY IS CERTIFIED AS: X A. In Compliance With Program Requirements Compliance Based On: <u> </u> 1. Acceptable POC B. Not in Compliance with Program Requirements and/or Applied Waivers: * Code: A* (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
12.Total Facility Beds 73 (L18)		
13.Total Certified Beds 73 (L17)		

14. LTC CERTIFIED BED BREAKDOWN	15. FACILITY MEETS
18 SNF 18/19 SNF 19 SNF ICF IID 40 33 (L37) (L38) (L39) (L42) (L43)	1861 (e) (1) or 1861 (j) (1): (L15)

16. STATE SURVEY AGENCY REMARKS (IF APPLICABLE SHOW LTC CANCELLATION DATE):

17. SURVEYOR SIGNATURE <u>Sue Reuss, Supervisor</u> Date : 03/23/2015 (L19)	18. STATE SURVEY AGENCY APPROVAL <u>Anne Kleppe, Enforcement Specialist</u> Date: 03/25/2015 (L20)
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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 : <u> </u>
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22. ORIGINAL DATE OF PARTICIPATION 02/01/1988 (L24)	23. LTC AGREEMENT BEGINNING DATE (L41)	24. LTC AGREEMENT ENDING DATE (L25)	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
25. LTC EXTENSION DATE: (L27)	27. ALTERNATIVE SANCTIONS A. Suspension of Admissions: (L44) B. Rescind Suspension Date: (L45)		

28. TERMINATION DATE: (L28)	29. INTERMEDIARY/CARRIER NO. 03001 (L31)	30. REMARKS
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31. RO RECEIPT OF CMS-1539 (L32)	32. DETERMINATION OF APPROVAL DATE 03/12/2015 (L33)	DETERMINATION APPROVAL
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Protecting, Maintaining and Improving the Health of Minnesotans

CMS Certification Number (CCN): 24-5524

Electronically Delivered: March 25, 2015

Sister Mary Elizabeth Anderson, Administrator
Little Sisters of the Poor
330 Exchange Street South
Saint Paul, Minnesota 55102

Dear Sister Anderson:

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. Furthermore, we are recommending to the Minnesota Department of Human Services that your facility is recertified in the Medicaid program.

Effective March 6, 2015 the above facility is certified for:

40 - Skilled Nursing Facility/Nursing Facility Beds
33 - Nursing Facility II Beds

Your facility's Medicare approved area consists of 40 skilled nursing facility beds. Your facility's Medicaid approved area consists of all 33 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. Please contact me if you have any questions about this electronic notice.

Sincerely,

A handwritten signature in cursive script, appearing to read "Anne Kleppe".

Anne Kleppe, Enforcement Specialist
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Email: anne.kleppe@state.mn.us
Telephone: (651) 201-4124 Fax: (651) 215-9697



Protecting, Maintaining and Improving the Health of Minnesotans

Electronically Delivered: March 26, 2015

Sister Mary Elizabeth Anderson, Administrator
Little Sisters of the Poor
330 Exchange Street South
Saint Paul, Minnesota 55102

RE: Project Number S5524024

Dear Sister Anderson:

On February 17, 2015, we informed you that we would recommend enforcement remedies based on the deficiencies cited by this Department for a standard survey, completed on February 5, 2015. 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 March 18, 2015, the Minnesota Department of Health completed a Post Certification Revisit (PCR) by review of your plan of correction and on March 17, 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 February 5, 2015. We presumed, based on your plan of correction, that your facility had corrected these deficiencies as of February 27, 2015. Based on our PCR, we have determined that your facility has corrected the deficiencies issued pursuant to our standard survey, completed on February 5, 2015, effective March 6, 2015 and therefore remedies outlined in our letter to you dated February 17, 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.

Please contact me if you have any questions about this electronic notice.

Sincerely,

A handwritten signature in cursive script that reads "Anne Kleppe".

Anne Kleppe, Enforcement Specialist
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Email: anne.kleppe@state.mn.us
Telephone: (651) 201-4124 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 245524	(Y2) Multiple Construction A. Building _____ B. Wing _____	(Y3) Date of Revisit 3/18/2015
Name of Facility LITTLE SISTERS OF THE POOR	Street Address, City, State, Zip Code 330 EXCHANGE STREET SOUTH SAINT PAUL, MN 55102	

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 F0431	Correction Completed 03/06/2015	ID Prefix _____	Correction Completed	ID Prefix _____	Correction Completed
Reg. # 483.60(b), (d), (e)		Reg. # _____		Reg. # _____	
LSC _____		LSC _____		LSC _____	
ID Prefix _____	Correction Completed	ID Prefix _____	Correction Completed	ID Prefix _____	Correction Completed
Reg. # _____		Reg. # _____		Reg. # _____	
LSC _____		LSC _____		LSC _____	
ID Prefix _____	Correction Completed	ID Prefix _____	Correction Completed	ID Prefix _____	Correction Completed
Reg. # _____		Reg. # _____		Reg. # _____	
LSC _____		LSC _____		LSC _____	
ID Prefix _____	Correction Completed	ID Prefix _____	Correction Completed	ID Prefix _____	Correction Completed
Reg. # _____		Reg. # _____		Reg. # _____	
LSC _____		LSC _____		LSC _____	
ID Prefix _____	Correction Completed	ID Prefix _____	Correction Completed	ID Prefix _____	Correction Completed
Reg. # _____		Reg. # _____		Reg. # _____	
LSC _____		LSC _____		LSC _____	

Reviewed By _____	Reviewed By SR/AK	Date: 03/23/2015	Signature of Surveyor: _____ 16022	Date: 03/18/2015
Reviewed By _____	Reviewed By	Date:	Signature of Surveyor:	Date:

Followup to Survey Completed on: 2/5/2015	Check for any Uncorrected Deficiencies. Was a Summary of Uncorrected Deficiencies (CMS-2567) Sent to the Facility? <table style="float: right;"> <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 245524	(Y2) Multiple Construction A. Building 01 - MAIN BUILDING 01 B. Wing	(Y3) Date of Revisit 3/17/2015
Name of Facility LITTLE SISTERS OF THE POOR	Street Address, City, State, Zip Code 330 EXCHANGE STREET SOUTH SAINT PAUL, MN 55102	

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 K0050	Correction Completed 02/23/2015	ID Prefix _____ Reg. # NFPA 101 LSC K0062	Correction Completed 02/23/2015	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 PS/AK	Date: 03/23/2015	Signature of Surveyor: 12424	Date: 03/17/2015
Reviewed By _____ CMS RO	Reviewed By	Date:	Signature of Surveyor:	Date:

Followup to Survey Completed on: 2/3/2015	Check for any Uncorrected Deficiencies. Was a Summary of Uncorrected Deficiencies (CMS-2567) Sent to the Facility? <table style="float: right;"> <tr> <td>YES</td> <td>NO</td> </tr> </table>	YES	NO
YES	NO		

MEDICARE/MEDICAID CERTIFICATION AND TRANSMITTAL
PART I - TO BE COMPLETED BY THE STATE SURVEY AGENCY

ID: CQGF

Facility ID: 00763

<p>1. MEDICARE/MEDICAID PROVIDER NO. (L1) 245524</p> <p>2. STATE VENDOR OR MEDICAID NO. (L2) 825540700</p>	<p>3. NAME AND ADDRESS OF FACILITY (L3) LITTLE SISTERS OF THE POOR (L4) 330 EXCHANGE STREET SOUTH (L5) SAINT PAUL, MN (L6) 55102</p>	<p>4. TYPE OF ACTION: 2 (L8)</p> <p>1. Initial 2. Recertification 3. Termination 4. CHOW 5. Validation 6. Complaint 7. On-Site Visit 9. Other 8. Full Survey After Complaint</p>															
<p>5. EFFECTIVE DATE CHANGE OF OWNERSHIP (L9)</p> <p>6. DATE OF SURVEY 02/05/2015 (L34)</p> <p>8. ACCREDITATION STATUS: ___ (L10) 0 Unaccredited 1 TJC 2 AOA 3 Other</p>	<p>7. PROVIDER/SUPPLIER CATEGORY <u>03</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</p>	<p>FISCAL YEAR ENDING DATE: (L35) 12/31</p>															
<p>11. LTC PERIOD OF CERTIFICATION From (a): To (b):</p> <p>12. Total Facility Beds 73 (L18)</p> <p>13. Total Certified Beds 73 (L17)</p>	<p>10. THE FACILITY IS CERTIFIED AS:</p> <p><input checked="" type="checkbox"/> Compliance With <u>And/Or Approved Waivers Of The Following Requirements:</u></p> <p style="margin-left: 20px;">Program Requirements ___ 2. Technical Personnel ___ 6. Scope of Services Limit Compliance Based On: ___ 3. 24 Hour RN ___ 7. Medical Director 1. Acceptable POC ___ 4. 7-Day RN (Rural SNF) ___ 8. Patient Room Size ___ 5. Life Safety Code ___ 9. Beds/Room</p> <p>B. Not in Compliance with Program <input checked="" type="checkbox"/> Requirements and/or Applied Waivers: * Code: B (L12)</p>																
<p>14. LTC CERTIFIED BED BREAKDOWN</p> <table style="width:100%; border-collapse: collapse;"> <tr> <td style="width:20%;">18 SNF</td> <td style="width:20%;">18/19 SNF</td> <td style="width:20%;">19 SNF</td> <td style="width:20%;">ICF</td> <td style="width:20%;">IID</td> </tr> <tr> <td></td> <td style="text-align: center;">40</td> <td style="text-align: center;">33</td> <td></td> <td></td> </tr> <tr> <td>(L37)</td> <td>(L38)</td> <td>(L39)</td> <td>(L42)</td> <td>(L43)</td> </tr> </table>		18 SNF	18/19 SNF	19 SNF	ICF	IID		40	33			(L37)	(L38)	(L39)	(L42)	(L43)	<p>15. FACILITY MEETS</p> <p>1861 (e) (1) or 1861 (j) (1): (L15)</p>
18 SNF	18/19 SNF	19 SNF	ICF	IID													
	40	33															
(L37)	(L38)	(L39)	(L42)	(L43)													
<p>16. STATE SURVEY AGENCY REMARKS (IF APPLICABLE SHOW LTC CANCELLATION DATE):</p>																	
<p>17. SURVEYOR SIGNATURE <u>Vidya Tomar, HFE NE II</u></p>	<p>Date: 03/02/2015 (L19)</p>	<p>18. STATE SURVEY AGENCY APPROVAL <u>Anne Kleppe, Enforcement Specialist</u> 03/11/2015 (L20)</p>															
<p>PART II - TO BE COMPLETED BY HCFA REGIONAL OFFICE OR SINGLE STATE AGENCY</p>																	
<p>19. DETERMINATION OF ELIGIBILITY</p> <p>___ 1. Facility is Eligible to Participate ___ 2. Facility is not Eligible (L21)</p>	<p>20. COMPLIANCE WITH CIVIL RIGHTS ACT:</p>	<p>21. 1. Statement of Financial Solvency (HCFA-2572) 2. Ownership/Control Interest Disclosure Stmt (HCFA-1513) 3. Both of the Above: _____</p>															
<p>22. ORIGINAL DATE OF PARTICIPATION 02/01/1988 (L24)</p>	<p>23. LTC AGREEMENT BEGINNING DATE (L41)</p>	<p>24. LTC AGREEMENT ENDING DATE (L25)</p>															
<p>25. LTC EXTENSION DATE: (L27)</p>	<p>27. ALTERNATIVE SANCTIONS</p> <p>A. Suspension of Admissions: (L44)</p> <p>B. Rescind Suspension Date: (L45)</p>																
<p>28. TERMINATION DATE:</p>	<p>29. INTERMEDIARY/CARRIER NO. 03001 (L28)</p>	<p>30. REMARKS</p>															
<p>31. RO RECEIPT OF CMS-1539 (L32)</p>	<p>32. DETERMINATION OF APPROVAL DATE (L33)</p>	<p>DETERMINATION APPROVAL</p>															



Protecting, Maintaining and Improving the Health of Minnesotans

Certified Mail # 7010 1670 0000 8044 5766

February 17, 2015

Sister Mary Elizabeth Anderson, Administrator
Little Sisters of the Poor
330 Exchange Street South
Saint Paul, MN 55102

RE: Project Number S5524024

Dear Sister Anderson:

On February 5, 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;

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:

Susanne Reuss, Unit Supervisor
Minnesota Department of Health
P.O. Box 64900
St. Paul, Minnesota 55164-0900

Email: susanne.reuss@state.mn.us
Telephone: (651) 201-3793
Fax: (651) 201-3790

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 March 17, 2015, the Department of Health will impose the following remedy:

- State Monitoring. (42 CFR 488.422)

PLAN OF CORRECTION (PoC)

A PoC for the deficiencies must be submitted within **ten calendar days** of your receipt of this letter. Your PoC 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,

- Include signature of provider and date.

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

If an acceptable PoC 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 PoC 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 PoC 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 PoC 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 PoC for the respective deficiencies (if any) is acceptable.

VERIFICATION OF SUBSTANTIAL COMPLIANCE

Upon receipt of an acceptable PoC, 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 PoC, unless it is determined that either correction actually occurred between the latest correction date on the PoC and the date of the first revisit, or correction occurred sooner than the latest correction date on the PoC.

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 May 5, 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 August 5, 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 a PoC 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 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:

Little Sisters of the Poor
February 17, 2015
Page 5

Mr. Patrick Sheehan, Supervisor
Health Care Fire Inspections
State Fire Marshal Division
444 Minnesota Street, Suite 145
St. Paul, Minnesota 55101-5145

Email: pat.sheehan@state.mn.us
Telephone: (651) 201-7205
Fax: (651) 215-0525

Feel free to contact me if you have questions.

Sincerely,



Anne Kleppe, Enforcement Specialist
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Email: anne.kleppe@state.mn.us
Telephone: (651) 201-4124 Fax: (651) 215-9697

Enclosure

cc: Licensing and Certification File

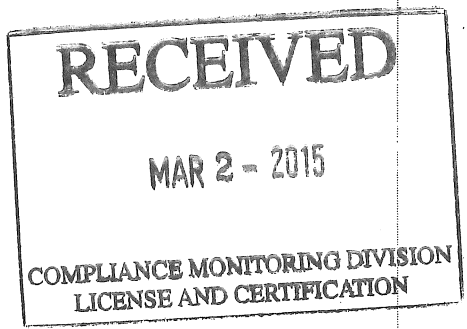
DEPARTMENT OF HEALTH AND HUMAN SERVICES
CENTERS FOR MEDICARE & MEDICAID SERVICES

PRINTED: 02/17/2015
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245524	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____	(X3) DATE SURVEY COMPLETED 02/05/2015
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NAME OF PROVIDER OR SUPPLIER LITTLE SISTERS OF THE POOR	STREET ADDRESS, CITY, STATE, ZIP CODE 330 EXCHANGE STREET SOUTH SAINT PAUL, MN 55102
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(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
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F 000	INITIAL COMMENTS The facility's plan of correction (POC) 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. Upon receipt of an acceptable 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 431 SS=E	483.60(b), (d), (e) DRUG RECORDS, LABEL/STORE DRUGS & BIOLOGICALS The facility must employ or obtain the services of a licensed pharmacist who establishes a system of records of receipt and disposition of all controlled drugs in sufficient detail to enable an accurate reconciliation; and determines that drug records are in order and that an account of all controlled drugs is maintained and periodically reconciled. Drugs and biologicals used in the facility must be labeled in accordance with currently accepted professional principles, and include the appropriate accessory and cautionary instructions, and the expiration date when applicable. In accordance with State and Federal laws, the facility must store all drugs and biologicals in locked compartments under proper temperature controls, and permit only authorized personnel to have access to the keys. The facility must provide separately locked,	F 431		All expired medications , unlabeled and undated meds were disposed and medications were reordered on 2/3/15. for R3,R37,R38,R2,R10,R39,R32,R35,R40 , R46, R62,R73,R77 Consult pharmacy nurse came to facility to complete medication cart and med room review on 2/11/15. There were no other outdated, unlabeled medications in the med cart. Stock medications reviewed and removed and replaced. On 2/25/15 nurses meeting was held and review of deficiency was discussed. Nurses were educated on plan of correction. Policy was re-

3/2/15
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LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE <i>Mary Elizabeth Anderson, Adm, RN</i>	TITLE	(X6) DATE 2/27/15
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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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CENTERS FOR MEDICARE & MEDICAID SERVICES

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F 431	<p>Continued From page 1</p> <p>permanently affixed compartments for storage of controlled drugs listed in Schedule II of the Comprehensive Drug Abuse Prevention and Control Act of 1976 and other drugs subject to abuse, except when the facility uses single unit package drug distribution systems in which the quantity stored is minimal and a missing dose can be readily detected.</p> <p>This REQUIREMENT is not met as evidenced by: Based on observation, interview and document review, the facility failed to ensure medications were stored and labeled properly for 13 of 21 residents (R3, R37, R38, R2, R10, R39, R32, R35, R40, R46, R62, R73 and R77) whose medications were observed for medication storage.</p> <p>Findings include:</p> <p>During observations of multiple medication storage areas throughout the facility, medications for R3, R37, R38, R2, R10, R39, R32, R35, R40, R46, R62, R73 and R77, which included eye drops, nasal sprays and inhaler, lacked dates to indicate when they were opened, unlabeled or the medications were expired.</p> <p>During the medication storage tour on 2/3/15, at 10:12 a.m. with licensed practical nurse (LPN)-A, in the 2nd floor medication storage area, one unlabeled bottle of Dorzolamide hcl-Timolol maleate ophthalmic (eye drops used to decrease pressure in eyes) solution dated 1/17/15.</p> <p>On 2/3/15, at 10:18 a.m. LPN-A and trained</p>	F 431	<p>viewed . All TMA's informed and educated of new practice by 3/6/15.</p> <p>An auditing form was developed for nursing staff to monitor med carts and med rooms on a monthly basis. In addition will continue with consult pharmacy nurse to complete quarterly audits.</p> <p>DON will track med cart/room audit forms on a monthly basis and follow up as needed for any system errors or problems.</p> <p>The POC will be integrated into the QA program and reviewed quarterly for effectiveness.</p> <p>The completion dates for our new POC will be totally complete by 3/6/15 at which time all nursing staff will have been educated and new audit form in place and ready for use.</p>
			3/6/15

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F 431	<p>Continued From page 2</p> <p>medication aide (TMA)-A verified the medications needed to be stored properly, with proper labels and a date when opened. At 10:21 a.m. LPN-A stated, the eye drop should have been labeled and would inform her supervisor to take corrective actions for storing medications properly.</p> <p>During the medication storage tour on 2/3/15, at 11:22 a.m. with TMA-B, in the 5th floor medication storage area, and the medication carts, multiple opened, undated and unlabeled medication bottles were stored. Observations included the following:</p> <ul style="list-style-type: none"> . R3's azelastine (anti-allergy) eye drop bottle was opened and undated. . R37's liquitears (dry eyes) eye drop bottle was opened, used and was undated. . R38's Erythromycin eye ointment (antibiotic/lubricant) was opened, undated and was expired. <p>During interview on 2/3/15 at 11:26 a.m. registered nurse (RN)-A verified the medications needed to be stored properly, with correct open date. The expired medications needed to be discarded from the medication room and the medication carts. Further, RN-A stated she would notify RN-B (the infection control nurse) as to what was observed and take the steps needed to correct the issue because eye drops should be dated when opened.</p>	F 431		
	<p>During the medication storage tour on 2/3/15, at 2:25 p.m. the 3rd floor medication storage cart was reviewed. The following observations were made:</p> <ul style="list-style-type: none"> . R2's fluticasone (Allergic Rhinitis) nasal spray bottle was opened, used and undated. 			

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F 431	<p>Continued From page 3</p> <ul style="list-style-type: none"> · R10's nasal saline (Dry nose) nasal spray bottle was opened and was expired. · R39's Ipratropium (Allergic Rhinitis) nasal spray bottle was opened, used and undated. <p>During interview on 2/3/15, at 2:35 p.m. LPN-B verified the medications needed to be labeled and stored properly. LPN-B added that opened medications needed to be dated when opened and expired medications needed to be removed from the storage area. LPN-B further stated, she was going to order some new medications from the pharmacy to replace the undated, expired medications.</p> <p>During the medication storage tour on 2/3/15 at 2:40 p.m. with RN-A, on the 4th floor, multiple opened, undated, expired and unlabeled medication bottles/inhalers were observed stored in the medication carts. Observations included the following:</p> <ul style="list-style-type: none"> · R32's flovent (Asthma) inhaler was expired with a date of 9/12/14. · R35's Fluticasone (Non-Allergic Rhinitis) nasal spray bottle was opened, used and undated. · R40's artifi tears eye ointment (eye moisture) was opened and undated. In addition, proair oral inhaler (to treat SOB/Wheezing symptoms) was expired with a date of 12/8/14. 	F 431		
	<ul style="list-style-type: none"> · R46's erythromycin eye ointment (Dry eyes) was opened, used and undated. · R62's nevanac (Macular Edema) eye drop bottle was opened, used and undated. · R73's patanol (Allergic Conjunctivitis) eye drop bottle was opened and undated. · R77's Travatan (Glaucoma) eye drop bottle was opened, used and undated. 			

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F 431	<p>Continued From page 4</p> <p>During interview on 2/3/15, at 3:05 p.m. RN-A verified the medications needed to be stored properly, with proper labels. the nursing staff needed to check expired medications and remove them from the medication room and carts. Further, RN-A stated she would notify RN-B as to what was observed and take the steps needed to correct the issue because eye drops should be dated when opened.</p> <p>During interview on 2/4/15, at 8:01 a.m. the director of nursing (DON) indicated, staff were supposed to date medication bottles when opened, check for expired medications, remove expired medications and re-order them from the pharmacy. DON explained, all medications removed from the storage area had been re-ordered from the pharmacy. DON added, the nurse consultant from the pharmacy comes quarterly to check medication cart and the medication rooms to make sure the medication labels are accurate and current. DON explained all residents' medications needed to be dated and labeled accurately for safe medication administration.</p> <p>During interview on 2/4/15, at 8:30 a.m. the facility's consultant pharmacist (CP) stated her expectation was for facility staff to date each medication bottle when opened and discard expired medications.</p>	F 431			
	The facility's undated. Medication Expiration Dates policy read, "Outdated, contaminated, or deteriorated medications and those in containers that are cracked, soiled, unlabeled, or without secure closures are immediately removed from stock, disposed of according to facility procedures for medication destruction, and				

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F 431	Continued From page 5 reordered from the pharmacy if a current order exists."	F 431			

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K 000	<p>INITIAL COMMENTS</p> <p>FIRE SAFETY</p> <p>THE FACILITY'S POC 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.</p> <p>UPON RECEIPT OF AN ACCEPTABLE POC, 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.</p> <p>At the time of this survey, LITTLE SISTERS OF THE POOR was found not in substantial compliance with the requirements for participation in Medicare/Medicaid, 42 CFR, Subpart 483.70(a), Life Safety from Fire, and National Fire Protection Association (NFPA) Standard 101 - 2000 edition.</p> <p>PLEASE RETURN THE PLAN OF CORRECTION FOR THE FIRE SAFETY DEFICIENCIES (K-TAGS) TO:</p> <p>HEALTHCARE FIRE INSPECTIONS STATE FIRE MARSHAL DIVISION 445 MINNESOTA STREET, SUITE 145 ST. PAUL, MN 55101-5145</p> <p>Or by email to: Angela.Kappenman@state.mn.us and Marian.Whitney@state.mn.us</p>	K 000	<p>POC ok</p> <p>ES 3-2-15</p> <div style="border: 2px solid red; padding: 5px; text-align: center;"> <p>RECEIVED</p> <p>FEB 27 2015</p> <p>MN DEPT. OF PUBLIC SAFETY STATE FIRE MARSHAL DIVISION</p> </div>	

DC: 3-17-15
 Exit: 2-5-15

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE: Mary Elizabeth Anderson TITLE: Adm., R (X6) DATE: 2/27/15

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 THE PLAN OF CORRECTION FOR EACH DEFICIENCY MUST INCLUDE ALL OF THE FOLLOWING INFORMATION: 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 prevent a reoccurrence of the deficiency. This 5-story building constructed in 1977 was determined to be of Type II(222) construction. It has no basement and is fully fire sprinklered throughout. The facility has a capacity of 73 beds. At the time of survey the census was 69. The requirement at 42 CFR Subpart 483.70(a) is NOT MET as evidenced by:	K 000		
K 050 SS=F	NFPA 101 LIFE SAFETY CODE STANDARD Fire drills are held at unexpected times under varying conditions, at least quarterly on each shift. The staff is familiar with procedures and is aware that drills are part of established routine. Responsibility for planning and conducting drills is assigned only to competent persons who are qualified to exercise leadership. Where drills are conducted between 9 PM and 6 AM a coded announcement may be used instead of audible alarms. 19.7.1.2 This STANDARD is not met as evidenced by: Based on review of reports and interview, it was	K 050	A 12 month calendar has been created, starting January 2015 and ongoing. This computer program will send a reminder to the task log indicating time for Fire Drill.	2/23/15

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K 050	Continued From page 2 determined that the facility failed to conduct the required number of fire drills for each shift in the last 12-month period in accordance with NFPA 101 LSC (00) Section 19.7.1.2. This deficient practice could affect how staff react in the event of a fire. Findings include: During the facility tour between 09:00 AM and 1:00 PM on 02/03/2015, based on review of available documentation it was reveled that fire drills have not been conducted on a one per shift per quarter basis based by the following: 1) Fire drills were not conducted on a quarterly Basis. No fire drills were conducted on the evening shift during the 4th quarter of 2015. 2) Fire drills were not conducted on a quarterly Basis. No fire drills were conducted on the night shift during the 3rd quarter of 2015. This deficiency was varified by the Facilities Director (RS). NFPA 101 LIFE SAFETY CODE STANDARD	K 050		
K 062 SS=F	Required automatic sprinkler systems are continuously maintained in reliable operating condition and are inspected and tested periodically. 19.7.6, 4.6.12, NFPA 13, NFPA 25, 9.7.5 This STANDARD is not met as evidenced by: Based on record review and interview the complete automatic fire sprinkler system is not	K 062	A licensed independent contractor will conduct Quarterly and Annual Flow test. Records for Flow Tests will be recorded in the Life Safety Book with Fire Drill records.	2/23/15

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K 062	Continued From page 3 being maintained in accordance with NFPA 25(99) Section 9.2.7. This deficient practice could effect all occupants of the building if the system were to fail under fire conditions. Findings include: On facility tour between 09:00 AM and 01:00 PM on 02/03/2015, it was revealed during review of available fire sprinkler records that there was no documentation of quarterly sprinkler flow testing testing in the last 12 months. During an interview with facilites Director (RS), stated he was rescently hired and was unaware of quarterly testing.	K 062			



Protecting, Maintaining and Improving the Health of Minnesotans

Certified Mail # 7010 1670 0000 8044 5766

February 17, 2015

Sister Mary Elizabeth Anderson, Administrator
Little Sisters of the Poor
330 Exchange Street South
Saint Paul, Minnesota 55102

Re: Enclosed State Nursing Home Licensing Orders - Project Number S5524024

Dear Sister Anderson:

The above facility was surveyed on February 2, 2015 through February 5, 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, Compliance Monitoring 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.

The State licensing orders are delineated on the attached Minnesota Department of Health order form (attached). 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.

When all orders are corrected, the order form should be signed and returned to:

Susanne Reuss, Unit Supervisor
Minnesota Department of Health
P.O. Box 64900
St. Paul, Minnesota 55164-0900

Email: susanne.reuss@state.mn.us
Telephone: (651) 201-3793
Fax: (651) 201-3790

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 immediately contact me.

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,



Anne Kleppe, Enforcement Specialist
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
Email: anne.kleppe@state.mn.us
Telephone: (651) 201-4124 Fax: (651) 215-9697

Enclosures

cc: Original - Facility
Licensing and Certification File

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00763	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED 02/05/2015
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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: On February 2nd through February 5th 2015, surveyors of this Department's staff, visited the above provider and the following correction orders are issued. When corrections are completed, please sign and date, make a copy of these orders and return the original to the Minnesota Department of Health, Division of</p>	2 000	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.	
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Minnesota Department of Health
LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE

TITLE

(X6) DATE

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00763	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED 02/05/2015
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NAME OF PROVIDER OR SUPPLIER LITTLE SISTERS OF THE POOR	STREET ADDRESS, CITY, STATE, ZIP CODE 330 EXCHANGE STREET SOUTH SAINT PAUL, MN 55102
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2 000	Continued From page 1 Compliance Monitoring, Licensing and Certification Programs; P.O. Box 64900, St. Paul, Minnesota 55164-0900.	2 000	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 which 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.	
21620	MN Rule 4658.1345 Labeling of Drugs Drugs used in the nursing home must be labeled in accordance with part 6800.6300. This MN Requirement is not met as evidenced by: Based on observation, interview and document review, the facility failed to ensure medications were stored and labeled properly for 13 of 21	21620		

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21620	<p>Continued From page 2</p> <p>residents (R3, R37, R38, R2, R10, R39, R32, R35, R40, R46, R62, R73 and R77) whose medications were observed for medication storage.</p> <p>Findings include:</p> <p>During observations of multiple medication storage areas throughout the facility, medications for R3, R37, R38, R2, R10, R39, R32, R35, R40, R46, R62, R73 and R77, which included eye drops, nasal sprays and inhaler, lacked dates to indicate when they were opened, when they expired and some medications were unlabeled.</p> <p>During the medication storage tour on 2/3/15, at 10:12 a.m. with licensed practical nurse (LPN)-A, in the 2nd floor medication storage area, one unlabeled bottle of Dorzolamide hcl-Timolol maleate ophthalmic (eye drops used to decrease pressure in eyes) solution dated 1/17/15.</p> <p>On 2/3/15, at 10:18 a.m. LPN-A and trained medication aide (TMA)-A verified the medications needed to be stored properly, with proper labels and a date when opened. At 10:21 a.m. LPN-A stated, the eye drops should have been labeled and would inform her supervisor to take corrective actions for storing medications properly.</p> <p>During the medication storage tour on 2/3/15, at 11:22 a.m. with TMA-B, in the 5th floor medication storage area, and the medication carts, multiple opened, undated and unlabeled medication bottles were stored. Observations included the following:</p> <ul style="list-style-type: none"> . R3's azelastine (anti-allergy) eye drop bottle was opened and undated. . R37's liquitears (dry eyes) eye drop bottle 	21620		

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21620	<p>Continued From page 3</p> <p>was opened, used and was undated.</p> <ul style="list-style-type: none"> . R38's Erythromycin eye ointment (antibiotic/lubricant) was opened, undated and was expired. <p>During interview on 2/3/15 at 11:26 a.m. registered nurse (RN)-A verified the medications needed to be stored properly, with correct open date. The expired medications needed to be discarded from the medication room and the medication carts. Further, RN-A stated she would notify RN-B (the infection control nurse) as to what was observed and take the steps needed to correct the issue because eye drops should be dated when opened.</p> <p>During the medication storage tour on 2/3/15, at 2:25 p.m. the 3rd floor medication storage cart was reviewed. The following observations were made:</p> <ul style="list-style-type: none"> . R2's fluticasone (Allergic Rhinitis) nasal spray bottle was opened, used and undated. . R10's nasal saline (Dry nose) nasal spray bottle was opened and was expired. . R39's Ipratropium (Allergic Rhinitis) nasal spray bottle was opened, used and undated. <p>During interview on 2/3/15, at 2:35 p.m. LPN-B verified the medications needed to be labeled and stored properly. LPN-B added that opened medications needed to be dated when opened and expired medications needed to be removed from the storage area. LPN-B further stated, she was going to order some new medications from the pharmacy to replace the undated, expired medications.</p> <p>During the medication storage tour on 2/3/15 at 2:40 p.m. with RN-A, on the 4th floor, multiple opened, undated, expired and unlabeled</p>	21620		

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21620	<p>Continued From page 4</p> <p>medication bottles/inhalers were observed stored in the medication carts. Observations included the following:</p> <ul style="list-style-type: none"> . R32's flovent (Asthma) inhaler was expired with a date of 9/12/14. . R35's Fluticasone (Non-Allergic Rhinitis) nasal spray bottle was opened, used and undated. . R40's artifi tears eye ointment (eye moisture) was opened and undated. In addition, proair oral inhaler (to treat SOB/Wheezing symptoms) was expired with a date of 12/8/14. . R46's erythromycin eye ointment (Dry eyes) was opened, used and undated. . R62's nevanac (Macular Edema) eye drop bottle was opened, used and undated. . R73's patanol (Allergic Conjunctivitis) eye drop bottle was opened and undated. . R77's Travatan (Glaucoma) eye drop bottle was opened, used and undated. <p>During interview on 2/3/15, at 3:05 p.m. RN-A verified the medications needed to be stored properly, with proper labels, the nursing staff needed to check expired medications and remove them from the medication room and carts. Further, RN-A stated she would notify RN-B as to what was observed and take the steps needed to correct the issue because eye drops should be dated when opened.</p> <p>During interview on 2/4/15, at 8:01 a.m. the director of nursing (DON) indicated, staff were supposed to date medication bottles when opened, check for expired medications, remove expired medications and re-order them from the pharmacy. DON explained, all medications removed from the storage area had been re-ordered from the pharmacy. DON added, the nurse consultant from the pharmacy comes</p>	21620		

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21620	<p>Continued From page 5</p> <p>quarterly to check medication cart and the medication rooms to make sure the medication labels are accurate and current. DON explained all residents' medications needed to be dated and labeled accurately for safe medication administration.</p> <p>During interview on 2/4/15, at 8:30 a.m. the facility's consultant pharmacist (CP) stated her expectation was for facility staff to date each medication bottle when opened and discard expired medications.</p> <p>The facility's undated, Medication Expiration Dates policy read, "Outdated, contaminated, or deteriorated medications and those in containers that are cracked, soiled, unlabeled, or without secure closures are immediately removed from stock, disposed of according to facility procedures for medication destruction, and reordered from the pharmacy if a current order exists."</p> <p>SUGGESTED METHOD OF CORRECTION: The administrator, director of nursing (DON) and consulting pharmacist could review and revise policies and procedures for proper storage of medications. Nursing staff could be educated as necessary to the importance of labeling medications properly and discarding expired medications. The DON or designee, along with the pharmacist, could audit medications on a regular basis to ensure compliance.</p> <p>TIME PERIOD FOR CORRECTION: Twenty-one (21) days.</p>	21620		