

DEPARTMENT OF HEALTH AND HUMAN SERVICES

CENTERS FOR MEDICARE & MEDICAID SERVICES

MEDICARE/MEDICAID CERTIFICATION AND TRANSMITTAL

ID: 2GV2

PART I - TO BE COMPLETED BY THE STATE SURVEY AGENCY

Facility ID: 00236

1. MEDICARE/MEDICAID PROVIDER NO. (L1) 24E102 2. STATE VENDOR OR MEDICAID NO. (L2) 411742500	3. NAME AND ADDRESS OF FACILITY (L3) MOUNT OLIVET HOME (L4) 5517 LYNDAL AVE SOUTH (L5) MINNEAPOLIS, MN (L6) 55419	4. TYPE OF ACTION: <u>7</u> (L8) <div style="display: flex; justify-content: space-between;"> <div> 1. Initial 3. Termination 5. Validation 7. On-Site Visit </div> <div> 2. Recertification 4. CHOW 6. Complaint 9. Other </div> </div> 8. Full Survey After Complaint
5. EFFECTIVE DATE CHANGE OF OWNERSHIP (L9) 6. DATE OF SURVEY 05/30/2014 (L34) 8. ACCREDITATION STATUS: <u> </u> (L10) 0 Unaccredited 1 TJC 2 AOA 3 Other	7. PROVIDER/SUPPLIER CATEGORY <u>10</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) 09/30
11. LTC PERIOD OF CERTIFICATION From (a) : To (b) : 12. Total Facility Beds 94 (L18) 13. Total Certified Beds 94 (L17)	10. THE FACILITY IS CERTIFIED AS: <div style="display: flex;"> <div style="flex: 1;"> X A. In Compliance With Program Requirements Compliance Based On: <u> </u> 1. Acceptable POC </div> <div style="flex: 1;"> <u>And/Or Approved Waivers Of The Following Requirements:</u> <div style="display: flex; justify-content: space-between;"> <div> 2. Technical Personnel 3. 24 Hour RN 4. 7-Day RN (Rural SNF) 5. Life Safety Code </div> <div> 6. Scope of Services Limit 7. Medical Director 8. Patient Room Size 9. Beds/Room </div> </div> </div> </div> B. Not in Compliance with Program Requirements and/or Applied Waivers: * Code: A* (L12)	
14. LTC CERTIFIED BED BREAKDOWN <div style="display: flex; justify-content: space-around;"> <div>18 SNF (L37)</div> <div>18/19 SNF (L38)</div> <div>19 SNF 94 (L39)</div> <div>ICF (L42)</div> <div>IID (L43)</div> </div>		15. FACILITY MEETS 1861 (e) (1) or 1861 (j) (1): (L15)
16. STATE SURVEY AGENCY REMARKS (IF APPLICABLE SHOW LTC CANCELLATION DATE):		
17. SURVEYOR SIGNATURE <u>Gayle Lantto, Supervisor</u>	Date : 07/23/2014 (L19)	18. STATE SURVEY AGENCY APPROVAL <u>Anne Kleppe, Enforcement Specialist</u>
Date: 08/28/2014 (L20)		

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

19. DETERMINATION OF ELIGIBILITY <input checked="" type="checkbox"/> 1. Facility is Eligible to Participate <input type="checkbox"/> 2. Facility is not Eligible (L21)	20. COMPLIANCE WITH CIVIL RIGHTS ACT:	21. 1. Statement of Financial Solvency (HCFA-2572) 2. Ownership/Control Interest Disclosure Stmt (HCFA-1513) 3. Both of the Above : <u> </u>
22. ORIGINAL DATE OF PARTICIPATION 01/01/1975 (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. (L31)	
31. RO RECEIPT OF CMS-1539 (L32)	32. DETERMINATION OF APPROVAL DATE 07/14/2014 (L33)	
30. REMARKS DETERMINATION APPROVAL		



Protecting, Maintaining and Improving the Health of Minnesotans

CMS Certification Number (CCN): 24-E102

August 28, 2014

Mr. Timothy Hokanson, Administrator
Mount Olivet Home
5517 Lyndale Avenue South
Minneapolis, Minnesota 55419

Dear Mr. Hokanson:

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 June 27, 2014, the above facility is certified for:

94 - Nursing Facility II Beds

Your facility's Medicare approved area consists of all 94 nursing facility II beds.

You should advise our office of any changes in staffing, services, or organization, which might affect your certification status. Please note, it is your responsibility to share the information contained in this letter and the results of this PCR with the President of your facility's Governing Body.

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.

Sincerely,

A handwritten signature in cursive script, appearing to read "Anne Kleppe", is located below the "Sincerely," text.

Anne Kleppe, Enforcement Specialist
Licensing and Certification Program
Division of Compliance Monitoring
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

July 23, 2014

Mr. Timothy Hokanson, Administrator
Mount Olivet Home
5517 Lyndale Avenue South
Minneapolis, Minnesota 55419

RE: Project Number SE102025

Dear Mr. Hokanson:

On June 18, 2014, we informed you that we would recommend enforcement remedies based on the deficiencies cited by this Department for a standard survey, completed on May 30, 2014. This survey found the most serious deficiencies to be isolated deficiencies that constituted no actual harm with potential for more than minimal harm that was not immediate jeopardy (Level D) whereby corrections were required.

On July 14, 2014, the Minnesota Department of Health completed a Post Certification Revisit (PCR) by review of your plan of correction to verify that your facility had achieved and maintained compliance with federal certification deficiencies issued pursuant to a standard survey, completed on May 30, 2014. We presumed, based on your plan of correction, that your facility had corrected these deficiencies as of June 27, 2014. Based on our PCR, we have determined that your facility has corrected the deficiencies issued pursuant to our standard survey, completed on May 30, 2014, effective June 27, 2014 and therefore remedies outlined in our letter to you dated June 18, 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. Enclosed is a copy of the Post Certification Revisit Form, (CMS-2567B) from this visit.

Feel free to contact me if you have questions.

Sincerely,

A handwritten signature in cursive script, appearing to read "Anne Kleppe", is located below the "Sincerely," text.

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

Enclosure

cc: Licensing and Certification File

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 24E102	(Y2) Multiple Construction A. Building B. Wing	(Y3) Date of Revisit 7/14/2014
Name of Facility MOUNT OLIVET HOME		Street Address, City, State, Zip Code 5517 LYNDALE AVENUE SOUTH MINNEAPOLIS, MN 55419

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 F0282 Reg. # 483.20(k)(3)(ii) LSC	Correction Completed 06/17/2014	ID Prefix F0329 Reg. # 483.25(l) LSC	Correction Completed 06/27/2014	ID Prefix F0441 Reg. # 483.65 LSC	Correction Completed 06/06/2014
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

Reviewed By State Agency	Reviewed By GL/AK	Date: 07/23/2014	Signature of Surveyor: 15507	Date: 07/14/2014
Reviewed By CMS RO	Reviewed By	Date:	Signature of Surveyor:	Date:
Followup to Survey Completed on: 5/30/2014		Check for any Uncorrected Deficiencies. Was a Summary of Uncorrected Deficiencies (CMS-2567) Sent to the Facility? YES NO		

SURVEY TEAM COMPOSITION AND WORKLOAD REPORT

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

Provider/Supplier Number 24E102	Provider/Supplier Name MOUNT OLIVET HOME
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Type of Survey (select all that apply):

M					
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A Complaint Investigation E Initial Certification I Recertification
B Dumping Investigation F Inspection of Care J Sanction/Hearing
C Federal Monitoring G Validation K State License
D Follow-up Visit H Life safety Code L Chow

Extent of Survey (Select all that apply):

A					
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A Routine/Standard (all providers/suppliers)
B Extended Survey (HHA or long term care facility)
C Partial Extended Survey (HHA)
D Other Survey

SURVEY TEAM AND WORKLOAD DATA

Please enter the workload information for each surveyor. Use the surveyor's information number.

Surveyor Id Number (A)	First Date Arrived (B)	Last Date Departed (C)	Pre-Survey Preparation Hours (D)	On-Site Hours 12am-8am (E)	On-Site Hours 8am-6pm (F)	On-Site Hours 6pm-12am (G)	Travel Hours (H)	Off-Site Report Preparation Hours (I)
Team Leader 1. 15507	7/14/2014	7/14/2014	0.25	0.00	0.00	0.00	0.00	0.25
2.								
3.								
4.								
5.								
6.								
7.								
8.								
9.								
10.								

Total Supervisory Review Hours 0.25

Total Clerical/Data Entry Hours..... 3.25

Was Statement of Deficiencies given to the provider on-site at completion of the survey? N

DEPARTMENT OF HEALTH AND HUMAN SERVICES

CENTERS FOR MEDICARE & MEDICAID SERVICES

MEDICARE/MEDICAID CERTIFICATION AND TRANSMITTAL

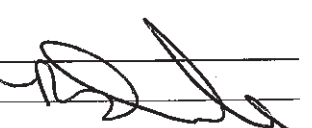
ID: 2GV2

PART I - TO BE COMPLETED BY THE STATE SURVEY AGENCY

Facility ID: 00236

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2. STATE VENDOR OR MEDICAID NO. (L2) 411742500		5. EFFECTIVE DATE CHANGE OF OWNERSHIP (L9)		6. DATE OF SURVEY 05/30/2014 (L34)	
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12. Total Facility Beds 94 (L18)		13. Total Certified Beds (L17)			
14. LTC CERTIFIED BED BREAKDOWN 18 SNF 18/19 SNF 19 SNF ICF IID (L37) (L38) (L39) (L42) (L43)		15. FACILITY MEETS 1861 (e) (1) or 1861 (j) (1): (L15)			
16. STATE SURVEY AGENCY REMARKS (IF APPLICABLE SHOW LTC CANCELLATION DATE):					
17. SURVEYOR SIGNATURE Shawn Soucek, HPR Social Work Specialist Date: 06/30/2014 (L19)			18. STATE SURVEY AGENCY APPROVAL Anne Kleppe, Enforcement Specialist Date: 07/11/2014 (L20)		

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

19. DETERMINATION OF ELIGIBILITY <input checked="" type="checkbox"/> 1. Facility is Eligible to Participate ___ 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 01/01/1975 (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) <u>VOLUNTARY</u> <u>00</u> <u>INVOLUNTARY</u> 01-Merger, Closure 05-Fail to Meet Health/Safety 02-Dissatisfaction W/ Reimbursement 06-Fail to Meet Agreement 03-Risk of Involuntary Termination <u>OTHER</u> 04-Other Reason for Withdrawal 07-Provider Status Change 00-Active	
28. TERMINATION DATE: (L28)		29. INTERMEDIARY/CARRIER NO. (L31)		30. REMARKS	
31. RO RECEIPT OF CMS-1539 (L32)		32. DETERMINATION OF APPROVAL DATE 7-14-14 (L33)		DETERMINATION APPROVAL 	



Protecting, Maintaining and Improving the Health of Minnesotans

Certified Mail # 7013 2250 0001 6356 5132

June 18, 2014

Mr. Timothy Hokanson, Administrator
Mount Olivet Home
5517 Lyndale Avenue South
Minneapolis, Minnesota 55419

RE: Project Number SE102025

Dear Mr. Hokanson:

On May 30, 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 **isolated deficiencies that constitute no actual harm with potential for more than minimal harm that is not immediate jeopardy (Level D)**, 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:

Gayle Lantto, Unit Supervisor
Minnesota Department of Health
P.O. Box 64900
St. Paul, Minnesota 55164-0900
Telephone: (651) 201-3794
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 July 9, 2014, 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 August 30, 2014 (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 November 30, 2014 (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

Mount Olivet Home

June 18, 2014

Page 5

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:

Mr. Patrick Sheehan, Supervisor
Health Care Fire Inspections
State Fire Marshal Division
444 Cedar Street, Suite 145
St. Paul, Minnesota 55101-5145
Telephone: (651) 201-7205
Fax: (651) 215-0541

Feel free to contact me if you have questions.

Sincerely,



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

Enclosure

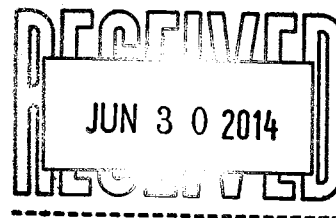
cc: Licensing and Certification File

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

PRINTED: 06/18/2014
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 24E102	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____	(X3) DATE SURVEY COMPLETED 05/30/2014
NAME OF PROVIDER OR SUPPLIER MOUNT OLIVET HOME			STREET ADDRESS, CITY, STATE, ZIP CODE 5517 LYNDAL AVENUE SOUTH MINNEAPOLIS, MN 55419	
(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. 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 282 SS=D	483.20(k)(3)(ii) SERVICES BY QUALIFIED PERSONS/PER CARE PLAN The services provided or arranged by the facility must be provided by qualified persons in accordance with each resident's written plan of care. This REQUIREMENT is not met as evidenced by: Based on interview and document review, the facility failed to follow the mood and behavioral care plan for 1 of 3 residents (R30) who were prescribed mood and behavioral altering medication and were reviewed for unnecessary medications. Findings include: R30's care plan dated 9/30/13, described the resident as having the potential for displaying verbal and physical aggression towards others. In addition, it was noted R30 was prescribed Celexa for depression and anxiety, and Depakote for mood disorder. The care plan directed staff to	F 282		

*POC accepted
bottom to
6/30/14*



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

TITLE

(X6) DATE

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

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

PRINTED: 06/18/2014
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 24E102	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED 05/30/2014
NAME OF PROVIDER OR SUPPLIER MOUNT OLIVET HOME			STREET ADDRESS, CITY, STATE, ZIP CODE 5517 LYNDAL AVENUE SOUTH MINNEAPOLIS, MN 55419		
(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 282	Continued From page 1 monitor the resident's behavior daily, and watch for adverse medication side effects. R30's behavior and potential medication side effects were not being monitored related to the use of medication intended to alter mood and behavior. R30 had diagnoses including dementia with behavior disturbances, anxiety and depression. At the time of a quarterly Minimum Data Set (MDS) dated 3/19/14, the resident did not exhibit behaviors such as verbal or physical directed towards others. The current physician's orders included Celexa 20 mg tablet daily for depression and Depakote 250 mg tablet twice a day for mood disorder.	F 282	The Behavior monitoring through electronic health records was re- instated with appropriate behaviors. The side effect monitoring was re- instated. An all house audit was done on residents with antipsychotic medications to see if the appropriate behavior monitoring was in place End of Month audits will be done on admits and re-admits to be sure monitoring is in place	5-30-14 5-30-14 6/3/14 On going	
F 329 SS=D	A registered nurse (RN)-A was interviewed on 5/30/14, at 9:00 a.m. and verified the staff had not been monitoring R30's behavior and potential side effects as directed on the care plan. 483.25(l) DRUG REGIMEN IS FREE FROM UNNECESSARY DRUGS Each resident's drug regimen must be free from unnecessary drugs. An unnecessary drug is any drug when used in excessive dose (including duplicate therapy); or for excessive duration; or without adequate monitoring; or without adequate indications for its use; or in the presence of adverse consequences which indicate the dose should be reduced or discontinued; or any combinations of the reasons above. Based on a comprehensive assessment of a resident, the facility must ensure that residents who have not used antipsychotic drugs are not given these drugs unless antipsychotic drug therapy is necessary to treat a specific condition	F 329	Behavior & Mood documentation procedure , Psychoactive Drug Procedure and Facility Process in gradual Dose Reduction procedure reviewed The Nurse Manager & DON will monitor for compliance	6-3-14 6-17-14 6-5-14 On going	

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F 329	<p>Continued From page 2</p> <p>as diagnosed and documented in the clinical record; and residents who use antipsychotic drugs receive gradual dose reductions, and behavioral interventions, unless clinically contraindicated, in an effort to discontinue these drugs.</p> <p>This REQUIREMENT is not met as evidenced by: Based on observation, interview and document review, the facility failed to identify adequate indications for use of mood and behavioral altering medications to support the continued need for the medication for 2 of 3 residents (R30, R35) reviewed for unnecessary medications.</p> <p>Findings include:</p> <p>R30's behavior and potential medication side effects were not being monitored related to the use of medication intended to alter mood and behavior. R30 had diagnoses including dementia with behavior disturbances, anxiety and depression. At the time of a quarterly Minimum Data Set (MDS) dated 3/19/14, the resident did not exhibit behaviors such as verbal or physical directed towards others. The current physician's orders included Celexa 20 mg tablet daily for depression and Depakote 250 mg tablet twice a day for mood disorder.</p> <p>Observations revealed the resident did not display any outward signs of behavioral or mood issues on 5/28/14, at 10:00 a.m. while seated in his wheelchair, on 5/29/14, at 7:00 a.m. while</p>			F 329	<p>Behavior monitoring and side effect monitoring through electronic health records was re-instated with appropriate behaviors.</p>		5-30-14

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F 329	<p>Continued From page 3</p> <p>getting ready for the day, and again on 5/30/14, at 7:30 a.m. when visiting with staff on the unit.</p> <p>R30's care plan dated 9/30/13, described the resident as having the potential for displaying verbal and physical aggression towards others. In addition, it was noted R30 was prescribed Celexa for depression and anxiety, and Depakote for mood disorder. The care plan directed staff to monitor the resident's behavior daily, and watch for adverse medication side effects.</p> <p>A registered nurse (RN)-A was interviewed on 5/30/14, at 9:00 a.m. and verified the staff had not been monitoring R30's behavior and potential side effects as directed on the care plan.</p> <p>A facility policy regarding the use of psychotropic medications was requested, but was not provided.</p> <p>R35's history included a copy of a physician clinic progress note near the time of the resident's admission dated 6/13/13, that revealed a long standing history of anxiety and depression with previous suicide attempts.</p> <p>The resident's care plan revised on 3/17/14, identified the following psychological changes and feelings: loss of independent living in the community with family and a desire to return to that setting, anxiety, feeling badly about self, and major depressive disorder with history of suicidal ideation. The care plan directed staff to monitor R35 every shift for signs of anxiety related to life situations, weepiness/crying, as well as delusions and hallucinations. However, facility staff were not monitoring the resident for the assessed and identified behavioral/mood issues. Instead,</p>	F 329	<p>Behavior & Mood Documentation procedure, Psychoactive Drug Procedure, and Facility Process in Gradual Dose Reduction procedure reviewed</p>	<p>6-3-14 6/17/14 6/5/14</p>	

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F 329	<p>Continued From page 4</p> <p>according to the Medication Administration Record (MAR) staff were monitoring R35 for physical/verbal aggression, hallucinations and delusions.</p> <p>On 5/28/14, at 4:24 p.m. a registered nurse (RN)-C stated R35 had bad feelings about not being able to babysit the grandchildren and to live with family. RN-C stated the facility tried to personalize the anxiety individuals may be feeling, which for R35 was related to loss of a previous living situation. RN-C noted staff was monitoring for physical and verbal aggression, hallucinations, and delusions, which was inconsistent with the resident's care plan. RN-C explained that the monitoring was not because the resident currently was exhibiting those behavioral issues, rather that there was potential for exhibiting them, and added, R35 "doesn't have any of those."</p> <p>R35's physician orders as of 5/28/14, for "major depression disorder single episode" included: Klonopin 1 milligram (mg) twice daily, (anticonvulsant commonly used for anxiety), Remeron 30 mg daily (antidepressant that had been increased from 7.5 mg on 2/9/14), Zoloft 100 mg daily (antidepressant) and Abilify 2 mg daily (antipsychotic medication). The orders also revealed that in 6/13 the anti-depressant medication Effexor had been decreased and then discontinued.</p> <p>A referral document to R35's psychiatrist dated 5/7/14, contained documentation by an unknown staff member, noting the resident stable and doing well. The documentation requested the psychiatrist complete a side-effect monitoring form as well as providing a rationale as to why a</p>	F 329	<p>Re-education of staff regarding appropriate target behaviors and documentation completed</p> <p>Re-education of Nurse Manager regarding reviewing the Care Plan, target behaviors, and side effects with quarterly assessment completed</p> <p>House audit completed regarding antipsychotic medication monitoring along with side effect monitoring.</p> <p>End of Month audits will be done on admits and re-admits to be sure monitoring is in place</p>	<p>6/23/14 - 6/27/14</p> <p>6/6/14</p> <p>6/3/14</p> <p>ongoing</p>	

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F 329	Continued From page 5 dose reduction was contraindicated if that was the case. The resident's medication was not reduced, and a rationale by the psychiatrist was not located in the resident's medical record as requested on the referral. Periodic documentation of the resident's mood symptoms was also lacking in the record to show the continued need for the use of Klonopin, Remeron, Zoloft and Abilify. In addition, R35 was prescribed Synthroid 125 micrograms (mcg) daily for thyroid function, but the thyroid stimulating hormone (TSH) testing had not been completed as requested by the primary physician. On 10/30/13, a nurse requested the physician order a TSH, however, laboratory testing results in R35's record did not show evidence a TSH had ever been completed since the resident's admission in 6/13. On 5/28/14 at 4:00 p.m. RN-C explained that she had thought the laboratory order had been discontinued, as R35 saw a physician off-site. At 4:03 p.m. resident care coordinator (RCC)-A stated the laboratory calendar had been reviewed and it was reported a TSH had been completed on 12/10/13. RN-C stated she would contact the physician's clinic to get the laboratory report. As of 4:30 p.m. on 5/28/14, RN-C had not been unable to obtain documentation showing the TSH had been completed.	F 329	Attempts were made to obtain the Psychiatric notes prior to the resident being discharged from the facility with no results. The individual resident stated discharged to home on 6/3/14 so not able to adjust the specific behavioral monitoring. A TSH was completed off site 9/12/13 with results placed in the medical record. The MD ordered to d/c the 12/10/13 TSH and all labs to be done at his office. Nurse Manager and DON will monitor compliance	6/19/14	Ongoing
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	F 441			

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F 441	<p>Continued From page 6 of disease and infection.</p> <p>(a) Infection Control Program The facility must establish an Infection Control Program under which it - (1) Investigates, controls, and prevents infections in the facility; (2) Decides what procedures, such as isolation, should be applied to an individual resident; and (3) Maintains a record of incidents and corrective actions related to infections.</p> <p>(b) Preventing Spread of Infection (1) When the Infection Control Program determines that a resident needs isolation to prevent the spread of infection, the facility must isolate the resident. (2) The facility must prohibit employees with a communicable disease or infected skin lesions from direct contact with residents or their food, if direct contact will transmit the disease. (3) The facility must require staff to wash their hands after each direct resident contact for which hand washing is indicated by accepted professional practice.</p> <p>(c) Linens Personnel must handle, store, process and transport linens so as to prevent the spread of infection.</p> <p>This REQUIREMENT is not met as evidenced by: Based on observation, interview and document review the facility failed to proper infection control procedures were followed for 1 of 1 resident (R102) whose personal cares were observed.</p>	F 441			

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F 441	Continued From page 7 Findings include: R102 was assisted to use the toilet by a nursing assistant (NA)-A on 5/29/14, at 7:32 a.m. NA-A donned gloves and assisted the resident with grooming and dressing while the resident was seated on the toilet. NA-A then placed walker in front of R102 and assisted him to stand. While R102 was standing, NA-A washed the resident's buttocks with a washcloth and then pulled up the resident's underwear and pants. While still wearing the same gloves, NA-A opened a cabinet in the bathroom, removed oral care supplies, applied toothpaste to a toothbrush, and handed it to the resident. NA-A then filled a glass of water, and held it by the rim of the glass as it was handed to the resident. After R102 was finished brushing his teeth, NA-A ran gloved finger over the bristles of the brush to remove leftover toothpaste. NA-A then moistened a washcloth and washed R102's face and eyes, and then lotioned his face. At that point NA-A removed her gloves and without washing or sanitizing her hands, wet a comb and combed R102's hair. NA-A then proceeded to rinse and dry the resident's glasses and placed the on the resident. The garbage was then retrieved from bathroom. NA-A then donned gloves to retrieved the soiled laundry, and then removed the gloves but did not wash or sanitize hands. NA-A then proceeded to make the resident's bed. At 8:00 a.m. NA-A left R102's room to dispose of the garage and linens, but again did not wash or sanitize her hands. At 8:07 a.m. NA-A went in another resident's room and turned off the room light. NA-A then proceeded to another resident's room and knocked and entered the resident's room. At 8:05 a.m. NA-A entered yet another resident's room,	F 441	Infection Control Policy and Procedure reviewed The NA/R involved was re-educated Staff were in-serviced on proper hand washing Random audits are being completed Nurses, Nurse Manager and DON will monitor compliance	6/4/14 6/6/14 6/5/14 & prn 6-5-14 and on going On going	

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F 441	<p>Continued From page 8 shut the door, and then washed her hands.</p> <p>A registered nurse (RN)-B was interviewed regarding the infection control breaches on 5/30/14, at 9:20 a.m. RN-B stated the expectation was that the staff would remove soiled gloves after perineal care, and then perform hand washing prior to then initiating other tasks.</p> <p>A policy regarding hand hygiene was requested, but was not provided.</p>	F 441			

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

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FE107023

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 24E102	(X2) MULTIPLE CONSTRUCTION A. BUILDING 01 - MAIN BUILDING 01 B. WING _____		(X3) DATE SURVEY COMPLETED 06/03/2014
NAME OF PROVIDER OR SUPPLIER MOUNT OLIVET HOME			STREET ADDRESS, CITY, STATE, ZIP CODE 5517 LYNDALE AVENUE SOUTH MINNEAPOLIS, MN 55419		
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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, Mount Olivet Home was found in substantial compliance with the requirements for participation in Medicare/Medicaid at 42 CFR, Subpart 483.70(a), Life Safety from Fire, and the 2000 edition of National Fire Protection Association (NFPA) Standard 101, Life Safety Code (LSC), Chapter 19 Existing Health Care.</p> <p>Mount Olivet Home is a 4-story building with a no basement. The building was constructed at 2 different times. The original building was constructed in 1968 and was determined to be of Type II(222) construction. In 2003, an addition was constructed to the South side of the building that was determined to be of Type II(222) construction. Because the original building and the addition meet the construction type allowed for existing buildings, the facility was surveyed as one building.</p> <p>The building is fully fire sprinkler protected. The facility has a complete fire alarm system with smoke detection in the corridors and spaces open to the corridor, that is monitored for automatic fire department notification. The facility has a licensed capacity of 94 beds and had a census of 87 at the time of the survey.</p> <p>The requirement at 42 CFR Subpart 483.70(a) is MET.</p>	K 000			
LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE			TITLE		(X6) DATE

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



Protecting, Maintaining and Improving the Health of Minnesotans

Certified Mail # 7013 2250 0001 6356 5132

June 18, 2014

Mr. Timothy Hokanson, Administrator
Mount Olivet Home
5517 Lyndale Avenue South
Minneapolis, Minnesota 55419

Re: Enclosed State Boarding Care Home Licensing Orders - Project Number SE102025

Dear Mr. Hokanson:

The above facility survey was completed on May 30, 2014 for the purpose of assessing compliance with Minnesota Department of Health Boarding Care 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.

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 Boarding Care 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 is 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.

Mount Olivet Home

June 18, 2014

Page 2

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

Gayle Lantto, Unit Supervisor
Minnesota Department of Health
P.O. Box 64900
St. Paul, Minnesota 55164-0900
Telephone: (651) 201-3794
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.

Feel free to contact me with any questions related to this letter.

Sincerely,



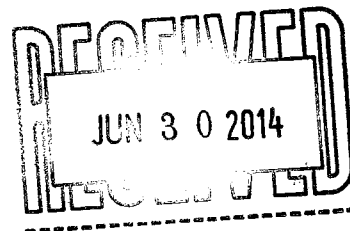
Anne Kleppe, Enforcement Specialist
Licensing and Certification Program
Division of Compliance Monitoring
Minnesota Department of Health
Email: anne.kleppe@state.mn.us
Telephone: (651) 201-4124
Fax: (651) 215-9697

Enclosure(s)

cc: Original - Facility
Licensing and Certification File

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00236	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____	(X3) DATE SURVEY COMPLETED 05/30/2014
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3 000	<p>INITIAL COMMENTS</p> <p>*****ATTENTION*****</p> <p>BOARDING CARE HOME 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 May 27, 28, 29, and 30, 2014, 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,</p>	3 000		



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

[Handwritten Signature]

TITLE

Administrator

(X6) DATE

05-27-14