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

MEDICARE/MEDICAID CERTIFICATION AND TRANSMITTAL — ID: QIYI

PART I - TO BE COMPLETED BY THE STATE SURVEY AGENCY — Facility ID: 27189

1. MEDICARE/MEDICAID PROVIDER NO.
(L1) **245617**

2.STATE VENDOR OR MEDICAID NO.
(L2) **550012400**

3. NAME AND ADDRESS OF FACILITY
(L3) **CARONDELET VILLAGE CARE CENTER**
(L4) **525 FAIRVIEW AVENUE SOUTH**
(L5) **SAINT PAUL, MN** (L6) **55116**

4. TYPE OF ACTION: **7** (L8)

1. Initial	**2. Recertification**
3. Termination	**4. CHOW**
5. Validation	**6. Complaint**
7. On-Site Visit	**9. Other**
8. Full Survey After Complaint	

5. EFFECTIVE DATE CHANGE OF OWNERSHIP
(L9)

6. DATE OF SURVEY **09/28/2016** (L34)

8. ACCREDITATION STATUS: ___ (L10)
0 Unaccredited 1 TJC
2 AOA 3 Other

7. PROVIDER/SUPPLIER CATEGORY **02** (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 **45** (L18)

13.Total Certified Beds **45** (L17)

10.THE FACILITY IS CERTIFIED AS:

X A. In Compliance With Program Requirements
Compliance Based On:
___1. Acceptable POC

B. Not in Compliance with Program Requirements and/or Applied Waivers:

And/Or Approved Waivers Of The Following Requirements:
___ 2. Technical Personnel
___ 3. 24 Hour RN
___ 4. 7-Day RN (Rural SNF)
___ 5. Life Safety Code
__ 6. Scope of Services Limit
__ 7. Medical Director
__ 8. Patient Room Size
__ 9. Beds/Room

* Code: **A*** (L12)

14. LTC CERTIFIED BED BREAKDOWN

18 SNF	18/19 SNF	19 SNF	ICF	IID
	45			
(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):

See Attached Remarks

17. SURVEYOR SIGNATURE — Date :
Susanne Reuss, Unit Supervisor — 09/28/2016 (L19)

18. STATE SURVEY AGENCY APPROVAL — Date:
Kate JohnsTon, Program Specialist — 10/13/2016 (L20)

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

19. DETERMINATION OF ELIGIBILITY
X 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
08/27/2012
(L24)

23. LTC AGREEMENT BEGINNING DATE
(L41)

24. LTC AGREEMENT ENDING DATE
(L25)

25. LTC EXTENSION DATE:
(L27)

27. ALTERNATIVE SANCTIONS
A. Suspension of Admissions:
(L44)
B. Rescind Suspension Date:
(L45)

26. TERMINATION ACTION: (L30)
VOLUNTARY **00**
01-Merger, Closure
02-Dissatisfaction W/ Reimbursement
03-Risk of Involuntary Termination
04-Other Reason for Withdrawal

INVOLUNTARY
05-Fail to Meet Health/Safety
06-Fail to Meet Agreement

OTHER
07-Provider Status Change
00-Active

28. TERMINATION DATE:
(L28)

29. INTERMEDIARY/CARRIER NO.
03001
(L31)

30. REMARKS

31. RO RECEIPT OF CMS-1539
(L32)

32. DETERMINATION OF APPROVAL DATE
09/22/2016
(L33)

DETERMINATION APPROVAL

MDH
Minnesota Department of Health

PROTECTING, MAINTAINING AND IMPROVING THE HEALTH OF ALL MINNESOTANS

CMS Certification Number (CCN): 245617
October 13, 2016

Mr. Gavin Middleton, Administrator
Carondelet Village Care Center
525 Fairview Avenue South
Saint Paul, MN 55116

Dear Mr. Middleton:

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 October 13, 2016 the above facility is certified for or recommended for:

45 Skilled Nursing Facility/Nursing Facility Beds

Your facility's Medicare approved area consists of all 45 skilled nursing facility beds.

You should advise our office of any changes in staffing, services, or organization, which might affect your certification status.

If, at the time of your next survey, we find your facility to not be in substantial compliance your Medicare and Medicaid provider agreement may be subject to non-renewal or termination.

Please contact me if you have any questions.

Sincerely,

Kate JohnsTon, Program Specialist
Program Assurance Unit
Licensing and Certification Program
Health Regulation Division
85 East Seventh Place, Suite 220
P.O. Box 64900
St. Paul, Minnesota 55164-0900
kate.johnston@state.mn.us
Telephone: (651) 201-3992 Fax: (651) 215-9697

cc: Licensing and Certification File

MDH
Minnesota
Department
of Health

PROTECTING, MAINTAINING AND IMPROVING THE HEALTH OF ALL MINNESOTANS

Electronically delivered
October 13, 2016

Mr.. Gavin Middleton, Administrator
Carondelet Village Care Center
525 Fairview Avenue South
Saint Paul, MN 55116

RE: Project Number S5617004

Dear Mr.. Middleton:

On August 30, 2016, we informed you that we would recommend enforcement remedies based on the deficiencies cited by this Department for a standard survey, completed on August 18, 2016. This survey found the most serious deficiencies to be a pattern of deficiencies that constituted no actual harm with potential for more than minimal harm that was not immediate jeopardy (Level E) whereby corrections were required.

On September 28, 2016, the Minnesota Department of Health completed a Post Certification Revisit (PCR) by review of your plan of correction and on August 22, 2016 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 August 18, 2016. We presumed, based on your plan of correction, that your facility had corrected these deficiencies as of September 19, 2016. Based on our PCR, we have determined that your facility has corrected the deficiencies issued pursuant to our standard survey, completed on August 18, 2016, effective September 19, 2016 and therefore remedies outlined in our letter to you dated August 30, 2016, will not be imposed.

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

Feel free to contact me if you have questions.

Sincerely,

Kate Johnston

Kate JohnsTon, Program Specialist
Program Assurance Unit
Licensing and Certification Program

Health Regulation Division
85 East Seventh Place, Suite 220
P.O. Box 64900
St. Paul, Minnesota 55164-0900
kate.johnston@state.mn.us
Telephone: (651) 201-3992 Fax: (651) 215-9697

Enclosure
cc: Licensing and Certification File

POST-CERTIFICATION REVISIT REPORT

PROVIDER / SUPPLIER / CLIA / IDENTIFICATION NUMBER (Y1)	MULTIPLE CONSTRUCTION (Y2)	DATE OF REVISIT (Y3)
245617	A. Building B. Wing	9/28/2016

NAME OF FACILITY	STREET ADDRESS, CITY, STATE, ZIP CODE
CARONDELET VILLAGE CARE CENTER	525 FAIRVIEW AVENUE SOUTH SAINT PAUL, MN 55116

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).

ITEM Y4	DATE Y5	ITEM Y4	DATE Y5	ITEM Y4	DATE Y5
ID Prefix F0431 Reg. # 483.60(b), (d), (e) LSC	Correction Completed 09/19/2016	ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed
ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed
ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed
ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed
ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed	ID Prefix Reg. # LSC	Correction Completed

REVIEWED BY STATE AGENCY ☐	REVIEWED BY (INITIALS)	DATE	SIGNATURE OF SURVEYOR	DATE
	SR/KJ	10/13/2016	16022	09/28/2016
REVIEWED BY CMS RO ☐	**REVIEWED BY (INITIALS)**	**DATE**	**TITLE**	**DATE**

FOLLOWUP TO SURVEY COMPLETED ON
8/18/2016

☐ CHECK FOR ANY UNCORRECTED DEFICIENCIES. WAS A SUMMARY OF UNCORRECTED DEFICIENCIES (CMS-2567) SENT TO THE FACILITY? ☐ YES ☐ NO

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

MEDICARE/MEDICAID CERTIFICATION AND TRANSMITTAL — ID: QIYI

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(L1) **245617**

2.STATE VENDOR OR MEDICAID NO.
(L2) **550012400**

3. NAME AND ADDRESS OF FACILITY
(L3) **CARONDELET VILLAGE CARE CENTER**
(L4) **525 FAIRVIEW AVENUE SOUTH**
(L5) **SAINT PAUL, MN** (L6) **55116**

4. TYPE OF ACTION: **2** (L8)
1. Initial **2. Recertification**
3. Termination **4. CHOW**
5. Validation **6. Complaint**
7. On-Site Visit **9. Other**
8. Full Survey After Complaint

5. EFFECTIVE DATE CHANGE OF OWNERSHIP
(L9)

6. DATE OF SURVEY **08/18/2016** (L34)

8. ACCREDITATION STATUS: ___ (L10)
0 Unaccredited 1 TJC
2 AOA 3 Other

7. PROVIDER/SUPPLIER CATEGORY **02** (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 **45** (L18)

13.Total Certified Beds **45** (L17)

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A. In Compliance With Program Requirements
Compliance Based On:
___1. Acceptable POC

X B. Not in Compliance with Program Requirements and/or Applied Waivers:

And/Or Approved Waivers Of The Following Requirements:
___ 2. Technical Personnel
___ 3. 24 Hour RN
___ 4. 7-Day RN (Rural SNF)
___ 5. Life Safety Code
__ 6. Scope of Services Limit
__ 7. Medical Director
__ 8. Patient Room Size
__ 9. Beds/Room

* Code: **B*** (L12)

14. LTC CERTIFIED BED BREAKDOWN

18 SNF	18/19 SNF	19 SNF	ICF	IID
	45			
(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 — Date :
Cynthia Wentkiewicz, HFE NE II — 09/06/2016 (L19)

18. STATE SURVEY AGENCY APPROVAL — Date:
Kate JohnsTon, Program Specialist — 09/13/2016 (L20)

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

19. DETERMINATION OF ELIGIBILITY
_____ 1. Facility is Eligible to Participate
_____ 2. Facility is not Eligible
(L21)

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21. 1. Statement of Financial Solvency (HCFA-2572)
2. Ownership/Control Interest Disclosure Stmt (HCFA-1513)
3. Both of the Above :
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22. ORIGINAL DATE OF PARTICIPATION
08/27/2012
(L24)

23. LTC AGREEMENT BEGINNING DATE
(L41)

24. LTC AGREEMENT ENDING DATE
(L25)

25. LTC EXTENSION DATE:
(L27)

27. ALTERNATIVE SANCTIONS
A. Suspension of Admissions:
(L44)
B. Rescind Suspension Date:
(L45)

26. TERMINATION ACTION: (L30)
VOLUNTARY **00**
01-Merger, Closure
02-Dissatisfaction W/ Reimbursement
03-Risk of Involuntary Termination
04-Other Reason for Withdrawal
INVOLUNTARY
05-Fail to Meet Health/Safety
06-Fail to Meet Agreement
OTHER
07-Provider Status Change
00-Active

28. TERMINATION DATE:
(L28)

29. INTERMEDIARY/CARRIER NO.
03001
(L31)

30. REMARKS
Posted 09/22/2016 Co.

31. RO RECEIPT OF CMS-1539
(L32)

32. DETERMINATION OF APPROVAL DATE
(L33)

DETERMINATION APPROVAL

MDH
Minnesota
Department
of Health

PROTECTING, MAINTAINING AND IMPROVING THE HEALTH OF ALL MINNESOTANS

Electronically delivered
August 30, 2016

Mr. Gavin Middleton, Administrator
Carondelet Village Care Center
525 Fairview Avenue South
Saint Paul, MN 55116

RE: Project Number S5617004

Dear Mr. Middleton:

On August 22, 2016, a standard survey was completed at your facility by the Minnesota Departments of Health and Public Safety to determine if your facility was in compliance with Federal participation requirements for skilled nursing facilities and/or nursing facilities participating in the Medicare and/or Medicaid programs. This survey found the most serious deficiencies in your facility to be a pattern of deficiencies that constitute no actual harm with potential for more than minimal harm that is not immediate jeopardy (Level E), as evidenced by the attached CMS-2567 whereby corrections are required. A copy of the Statement of Deficiencies (CMS-2567) is enclosed.

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

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

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

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

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

the time of a revisit;

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

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

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

DEPARTMENT CONTACT

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

Brenda Fischer, Unit Supervisor
St. Cloud A Survey Team
Licensing & Certification
Health Regulation Division
Minnesota Department of Health
Midtown Square
3333 West Division, #212
St. Cloud, Minnesota 56301
Telephone: (320)223-7338
Fax: (320)223-7348

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 September 27, 2016, the Department of Health will impose the following remedy:

- State Monitoring. (42 CFR 488.422)

ELECTRONIC PLAN OF CORRECTION (ePoC)

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

- Address how corrective action will be accomplished for those residents found to have been affected by the deficient practice;

- Address how the facility will identify other residents having the potential to be affected by the same deficient practice;

- Address what measures will be put into place or systemic changes made to ensure that the deficient practice will not recur;

- Indicate how the facility plans to monitor its performance to make sure that solutions are sustained. The facility must develop a plan for ensuring that correction is achieved and sustained. This plan must be implemented, and the corrective action evaluated for its effectiveness. The plan of correction is integrated into the quality assurance system;

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

- Submit electronically to acknowledge your receipt of the electronic 2567, your review and your ePoC submission.

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

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

- Optional denial of payment for new Medicare and Medicaid admissions (42 CFR 488.417 (a));

- Per day civil money penalty (42 CFR 488.430 through 488.444).

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

PRESUMPTION OF COMPLIANCE - CREDIBLE ALLEGATION OF COMPLIANCE

The facility's ePoC will serve as your allegation of compliance upon the Department's acceptance. Your signature at the bottom of the first page of the CMS-2567 form will be used as verification of compliance. In order for your allegation of compliance to be acceptable to the Department, the ePoC

must meet the criteria listed in the plan of correction section above. You will be notified by the Minnesota Department of Health, Licensing and Certification Program staff and/or the Department of Public Safety, State Fire Marshal Division staff, if your ePoC for the respective deficiencies (if any) is acceptable.

VERIFICATION OF SUBSTANTIAL COMPLIANCE

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

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

Original deficiencies not corrected

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

Original deficiencies not corrected and new deficiencies found during the revisit

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

Original deficiencies corrected but new deficiencies found during the revisit

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

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

If substantial compliance with the regulations is not verified by November 18, 2016 (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 February 18, 2017 (six months after the identification of noncompliance) if your facility does not achieve substantial compliance. This action is mandated by the Social Security Act at Sections 1819(h)(2)(C) and 1919(h)(3)(D) and Federal regulations at 42 CFR Sections 488.412 and 488.456.

INFORMAL DISPUTE RESOLUTION

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

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

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

Feel free to contact me if you have questions.

Sincerely,

Kate JohnsTon, Program Specialist
Program Assurance Unit
Licensing and Certification Program
Health Regulation Division
85 East Seventh Place, Suite 220
P.O. Box 64900
St. Paul, Minnesota 55164-0900
kate.johnston@state.mn.us
Telephone: (651) 201-3992 Fax: (651) 215-9697

Enclosure
cc: Licensing and Certification File

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION	(X3) DATE SURVEY COMPLETED
	245617	A. BUILDING _____ B. WING _____	08/18/2016

NAME OF PROVIDER OR SUPPLIER	STREET ADDRESS, CITY, STATE, ZIP CODE
CARONDELET VILLAGE CARE CENTER	525 FAIRVIEW AVENUE SOUTH SAINT PAUL, MN 55116

(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)	(X5) COMPLETION DATE
F 000	INITIAL COMMENTS The facility's plan of correction (POC) will serve as your allegation of compliance upon the Department's acceptance. Because you are enrolled in ePOC, your signature is not required at the bottom of the first page of the CMS-2567 form. Your electronic submission of the POC will be used as verification of compliance. Upon receipt of an acceptable electronic POC, an on-site revisit of your facility may be conducted to validate that substantial compliance with the regulations has been attained in accordance with your verification.	F 000		
F 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.	F 431		9/19/16

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE	TITLE	(X6) DATE
Electronically Signed		09/01/2016

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

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION	(X3) DATE SURVEY COMPLETED
	245617	A. BUILDING _____ B. WING _____	08/18/2016

NAME OF PROVIDER OR SUPPLIER	STREET ADDRESS, CITY, STATE, ZIP CODE
CARONDELET VILLAGE CARE CENTER	525 FAIRVIEW AVENUE SOUTH SAINT PAUL, MN 55116

(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 431	Continued From page 1 The facility must provide separately locked, 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. This REQUIREMENT is not met as evidenced by: Based on observation, interview and document review, the facility failed to ensure medications were stored properly in three of three neighborhoods, involving 5 residents (R55, R8, R42, R7, R22) of 44 residents. Findings include: On 8/15/16, starting at 12:30 p.m. a tour of the facility's medication storage was conducted on all three neighborhoods. During the tour the facility failed to date eye medications when opened and failed to discard eye medications per facility policy and per manufacturer's recommendations. Observations included the following: On neighborhood one, a bottle of latanoprost (to treat glaucoma) for R55 was not dated when opened; and a bottle of Timolol (for glaucoma) for R8 was dated as opened on 7/14/16. Registered nurse (RN)-A stated R55 had received the Timolol on 8/15/16. On neighborhood two, two open bottles of Timolol	F 431	All Care Center medication carts and medication rooms were immediately audited for proper records, labeling, and storing of medications. This included the auditing of any unlabeled or expired medications. All expired medications were removed and replaced. Policy and procedure regarding Medication Storage and Expiration Guidelines has been reviewed and is current. Education on frequency and practice of checking medications for expiration date and Medication Storage and Expiration Guidelines has been completed for nursing staff. Audits regarding Medication Storage will be conducted on all medication carts weekly for 4 weeks with results reported to Quality Assurance for ongoing compliance and will determine the need	

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION	(X3) DATE SURVEY COMPLETED
	245617	A. BUILDING _____ B. WING _____	08/18/2016

NAME OF PROVIDER OR SUPPLIER	STREET ADDRESS, CITY, STATE, ZIP CODE
CARONDELET VILLAGE CARE CENTER	525 FAIRVIEW AVENUE SOUTH SAINT PAUL, MN 55116

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F 431	Continued From page 2 were observed for R42. One bottle was dated as having been opened on 2/22/16. The second bottle of Timolol contained a label indicating the eye medication was filled on 8/1/16, but the bottle was not dated when opened. Trained medication aide (TMA)-A verified the second eye medication bottle had not been dated when opened. TMA-A stated not knowing why there were two bottles of Timolol for R42 or if one bottle was being used before the other. On neighborhood three, an open and undated bottle of latanoprost was observed for R7. Also a bottle of the eye medication Azopt (to treat glaucoma) for R22 was dated as having been opened on 7/11/16. RN-B verified R22 had received the Azopt on this date and the latanoprost had not been dated when opened. The facility's Medication Storage and Expiration Guidelines policy dated 9/07, indicated latanoprost was to be discarded 45 days after it was first used; and all other unspecified eye drops were to be discarded according to the manufacturer's labeled date. The policy indicated unspecified eye drops did not need to be dated when opened. On 8/17/16, at 1:59 p.m. registered pharmacist (RPH-B) was interviewed regarding the dating and discarding of open eye medications. RPH-A stated the expectation was that facility staff would date the eye medications when opened. RPH-A stated the typical recommendation for Timolol was to discard the medication 28 days after opening. RPH-A looked up the manufacturer's recommendations for the Azopt and stated the product insert did not indicate a discard date after opening. RPH-A stated if there were no	F 431	for further auditing. The Clinical Administrator or designee is responsible for ongoing compliance. Date certain for the purposes of ongoing compliance is 09/19/2016.	

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	245617	A. BUILDING _____ B. WING _____	08/18/2016

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CARONDELET VILLAGE CARE CENTER	**525 FAIRVIEW AVENUE SOUTH SAINT PAUL, MN 55116**

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F 431	Continued From page 3 manufacturer's recommendations the consulting pharmacy recommended discarding the eye medication 30-days after first opened.	F 431		

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION	(X3) DATE SURVEY COMPLETED
	245617	A. BUILDING **01 - CARONDELET VILLAGE CARE CENTER** B. WING _____	**08/22/2016**

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K 000	INITIAL COMMENTS FIRE SAFETY A Life Safety Code Survey was conducted by the Minnesota Department of Public Safety. At the time of this survey, CARONDELET VILLAGE CARE CENTER was found to be in substantial compliance with the requirements for participation in Medicare/Medicaid at 42 CFR, Subpart 483.70(a), Life Safety from Fire, and the 2000 edition of National Fire Protection Association (NFPA) Standard 101, Life Safety Code (LSC), Chapter 18 New health Care. Carondelet Village Care Center is located on the first floor of a 4-story building with a full basement. The building was constructed in 2011, and was determined to be of Type II(222) construction. The building is fully fire sprinklered throughout. The facility has a fire alarm system with smoke detection in the corridors, spaces open to the corridors and all resident rooms that is monitored for automatic fire department notification. The facility has a capacity of 45 beds and had a census of 41 at the time of the survey. The requirement at 42 CFR Subpart 483.70(a) is MET	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.

MDH
Minnesota
Department
of Health

PROTECTING, MAINTAINING AND IMPROVING THE HEALTH OF ALL MINNESOTANS

Electronically submitted
August 30, 2016

Mr. Gavin Middleton, Administrator
Carondelet Village Care Center
525 Fairview Avenue South
Saint Paul, MN 55116

Re: Enclosed State Nursing Home Licensing Orders - Project Number S5617004

Dear Mr. Middleton:

The above facility was surveyed on August 22, 2016 through August 22, 2016 for the purpose of assessing compliance with Minnesota Department of Health Nursing Home Rules. At the time of the survey, the survey team from the Minnesota Department of Health, Health Regulation Division, noted one or more violations of these rules that are issued in accordance with Minnesota Stat. section 144.653 and/or Minnesota Stat. Section 144A.10. If, upon reinspection, it is found that the deficiency or deficiencies cited herein are not corrected, a civil fine for each deficiency not corrected shall be assessed in accordance with a schedule of fines promulgated by rule of the Minnesota Department of Health.

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

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

The assigned tag number appears in the far left column entitled "ID Prefix Tag." The state statute/rule number and the corresponding text of the state statute/rule out of compliance is listed in the "Summary Statement of Deficiencies" column and replaces the "To Comply" portion of the correction

order. This column also includes the findings that are in violation of the state statute after the statement, "This Rule is not met as evidenced by." Following the surveyors findings are the Suggested Method of Correction and the Time Period For Correction.

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

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

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

Kate JohnsTon, Program Specialist
Program Assurance Unit
Licensing and Certification Program
Health Regulation Division
85 East Seventh Place, Suite 220
P.O. Box 64900
St. Paul, Minnesota 55164-0900
kate.johnston@state.mn.us
Telephone: (651) 201-3992 Fax: (651) 215-9697

Enclosure(s)
cc: Original - Facility
Licensing and Certification File

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION	(X3) DATE SURVEY COMPLETED
	27189	A. BUILDING: _____ B. WING _____	08/18/2016

NAME OF PROVIDER OR SUPPLIER	STREET ADDRESS, CITY, STATE, ZIP CODE
CARONDELET VILLAGE CARE CENTER	525 FAIRVIEW AVENUE SOUTH SAINT PAUL, MN 55116

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2 000	Initial Comments *****ATTENTION****** NH LICENSING CORRECTION ORDER 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. 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. 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. INITIAL COMMENTS: On August 15, 16, 17, and 18th 2016, surveyors of this Department's staff, visited the above provider and the following correction orders are issued. Please indicate in your electronic plan of correction that you have reviewed these orders, and identify the date when they will be completed.	2 000		

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21620	Continued From page 1	21620		
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 properly in three of three neighborhoods, involving 5 residents (R55, R8, R42, R7, R22) of 44 residents. Findings include: On 8/15/16, starting at 12:30 p.m. a tour of the facility's medication storage was conducted on all three neighborhoods. During the tour the facility failed to date eye medications when opened and failed to discard eye medications per facility policy and per manufacturer's recommendations. Observations included the following: On neighborhood one, a bottle of latanoprost (to treat glaucoma) for R55 was not dated when opened; and a bottle of Timolol (for glaucoma) for R8 was dated as opened on 7/14/16. Registered nurse (RN)-A stated R55 had received the Timolol on 8/15/16. On neighborhood two, two open bottles of Timolol were observed for R42. One bottle was dated as having been opened on 2/22/16. The second bottle of Timolol contained a label indicating the eye medication was filled on 8/1/16, but the bottle was not dated when opened. Trained medication aide (TMA)-A verified the second eye medication bottle had not been dated when opened. TMA-A stated not knowing why there were two bottles of	21620	Corrected 09/19/16	9/19/16

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21620	Continued From page 2 Timolol for R42 or if one bottle was being used before the other. On neighborhood three, an open and undated bottle of latanoprost was observed for R7. Also a bottle of the eye medication Azopt (to treat glaucoma) for R22 was dated as having been opened on 7/11/16. RN-B verified R22 had received the Azopt on this date and the latanoprost had not been dated when opened. The facility's Medication Storage and Expiration Guidelines policy dated 9/07, indicated latanoprost was to be discarded 45 days after it was first used; and all other unspecified eye drops were to be discarded according to the manufacturer's labeled date. The policy indicated unspecified eye drops did not need to be dated when opened. On 8/17/16, at 1:59 p.m. registered pharmacist (RPH-B) was interviewed regarding the dating and discarding of open eye medications. RPH-A stated the expectation was that facility staff would date the eye medications when opened. RPH-A stated the typical recommendation for Timolol was to discard the medication 28 days after opening. RPH-A looked up the manufacturer's recommendations for the Azopt and stated the product insert did not indicate a discard date after opening. RPH-A stated if there were no manufacturer's recommendations the consulting pharmacy recommended discarding the eye medication 30-days after first opened. SUGGESTED METHOD OF CORRECTION: The administrator, director of nursing (DON) and consulting pharmacist could review and revise policies and procedures for proper storage of	21620		

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21620	Continued From page 3 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. TIME PERIOD FOR CORRECTION: Twenty one (21) days.	21620		