



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically Delivered
April 9, 2021

Administrator
Birchwood Health Care Center
604 - 1st Street Ne
Forest Lake, MN 55025

RE: CCN: 245200
Cycle Start Date: March 2, 2021

Dear Administrator:

On April 1, 2021, the Minnesota Department of Health completed a revisit to verify that your facility had achieved and maintained compliance. Based on our review, we have determined that your facility has achieved substantial compliance; therefore no remedies will be imposed.

Feel free to contact me if you have questions.

A handwritten signature in black ink, appearing to read 'Melissa Poepping'.

Melissa Poepping, Health Program Representative Senior
Program Assurance | Licensing and Certification
Minnesota Department of Health
P.O. Box 64970
Saint Paul, Minnesota 55164-0970
Phone: 651-201-4117
Email: melissa.poepping@state.mn.us



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Electronically delivered
March 11, 2021

Administrator
Birchwood Health Care Center
604 - 1st Street Ne
Forest Lake, MN 55025

RE: CCN: 245200
Cycle Start Date: March 2, 2021

Dear Administrator:

On March 2, 2021, a 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 constituted no actual harm with potential for more than minimal harm that was not immediate jeopardy (Level E), as evidenced by the electronically attached CMS-2567 whereby corrections are required.

ELECTRONIC PLAN OF CORRECTION (ePoC)

Within **ten (10) calendar days** after your receipt of this notice, you must submit an acceptable ePOC for the deficiencies cited. An acceptable ePOC will serve as your allegation of compliance. Upon receipt of an acceptable ePOC, we will authorize a revisit to your facility to determine if substantial compliance has been achieved.

To be acceptable, a provider's ePOC must include the following:

- How corrective action will be accomplished for those residents found to have been affected by the deficient practice.
- How the facility will identify other residents having the potential to be affected by the same deficient practice.
- What measures will be put into place, or systemic changes made, to ensure that the deficient practice will not recur.
- How the facility will monitor its corrective actions to ensure that the deficient practice is being corrected and will not recur.
- The date that each deficiency will be corrected.
- An electronic acknowledgement signature and date by an official facility representative.

The state agency may, in lieu of an onsite 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:

- Denial of payment for new Medicare and Medicaid admissions (42 CFR 488.417);
- Civil money penalty (42 CFR 488.430 through 488.444).
- Termination of your facility's Medicare and/or Medicaid agreement (488.456(b)).

DEPARTMENT CONTACT

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

Karen Aldinger, Unit Supervisor
Metro C District Office
Licensing and Certification Program
Health Regulation Division
Minnesota Department of Health
85 East Seventh Place, Suite 220
P.O. Box 64900
Saint Paul, Minnesota 55164-0900
Email: karen.aldinger@state.mn.us
Office: (651) 201-3794 Mobile: (320) 249-2805

PRESUMPTION OF COMPLIANCE - CREDIBLE ALLEGATION OF COMPLIANCE

The facility's ePoC will serve as your allegation of compliance upon the Department's acceptance. 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, a Post Certification Revisit (PCR), of your facility will be conducted to validate that substantial compliance with the regulations has been attained in accordance with your verification.

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.

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 June 2, 2021 (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).

Birchwood Health Care Center

March 11, 2021

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In addition, if substantial compliance with the regulations is not verified by September 2, 2021 (six months after the identification of noncompliance) your provider agreement will be terminated. 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.

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.

INFORMAL DISPUTE RESOLUTION (IDR) / INDEPENDENT INFORMAL DISPUTE RESOLUTION (IIDR)

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:
https://mdhprovidercontent.web.health.state.mn.us/ltr_idr.cfm

You must notify MDH at this website of your request for an IDR or IIDR within the 10 calendar day period allotted for submitting an acceptable electronic plan of correction. A copy of the Department's informal dispute resolution policies are posted on the MDH Information Bulletin website at:
https://www.health.state.mn.us/facilities/regulation/infobulletins/ib04_8.html

Please note that the failure to complete the informal dispute resolution process will not delay the dates specified for compliance or the imposition of remedies.

Feel free to contact me if you have questions.

Sincerely,



Melissa Poeping, Health Program Representative Senior
Program Assurance | Licensing and Certification
Minnesota Department of Health
P.O. Box 64970
Saint Paul, Minnesota 55164-0970
Phone: 651-201-4117
Email: melissa.poeping@state.mn.us

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

PRINTED: 04/14/2021
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245200	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED C 03/02/2021
NAME OF PROVIDER OR SUPPLIER BIRCHWOOD HEALTH CARE CENTER			STREET ADDRESS, CITY, STATE, ZIP CODE 604 - 1ST STREET NE FOREST LAKE, MN 55025		
(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 On 3/2/21, an abbreviated survey was completed at your facility to conduct a complaint investigation. Your facility was found NOT to be in compliance with 42 CFR Part 483, Requirements for Long Term Care Facilities. The following complaint was found to be SUBSTANTIATED: H5200049C (MN00068952). A deficiency is being cited at F760. 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 760 SS=E	Residents are Free of Significant Med Errors CFR(s): 483.45(f)(2) The facility must ensure that its- §483.45(f)(2) Residents are free of any significant medication errors. This REQUIREMENT is not met as evidenced by: Based on interview and document review, the facility failed to ensure 1 of 2 residents (R1) received the correct dose of insulin per physician orders, reviewed for insulin administration. In	F 760	F760 The preparation of the following plan of correction for this deficiency does not	4/1/21	

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

TITLE

(X6) DATE

Electronically Signed

03/16/2021

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

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F 760	<p>Continued From page 1</p> <p>addition, the facility failed to routinely follow their policy to ensure accurate insulin dosing, of having two licensed nurses checking the dose prior to administration. This practice had the potential to affect 9 residents currently receiving insulin.</p> <p>Findings include:</p> <p>R1's quarterly Minimum Data Set (MDS) dated 2/4/21, indicated R1 received insulin daily and had severely impaired cognition. R1's diagnosis included type 2 diabetes mellitus without complications.</p> <p>R1's February 2021 medication administration record (MAR) included, Insulin Glargine solution pen-injector 100 UNIT/ML inject 20 units subcutaneously (under the skin) one time a day at bedtime</p> <p>R1's Facility Reported Incident (FRI) dated 3/1/21, at 8:00 p.m. included, registered nurse (RN)-A had administered 100 units of Lantus insulin to R1 instead of the ordered 20 units. Upon discovering the error, R1's blood sugar was immediately checked. At 10:30 p.m. R1's blood sugar was found to be 66 (normal 60-100). The on-call physician was contacted and ordered R1 to be sent to the hospital for observation where R1 remained and returned to the facility on 2/29/21, evening shift.</p> <p>On 3/2/21, at 11:40 a.m. licensed practical nurse (LPN)-A assigned at the memory care unit stated nurses were supposed to compare the physician order in the computer with the label on the insulin pen to make sure they matched and would then dial the units the resident was to receive after priming the pen. LPN-A then stated nurses were</p>	F 760	<p>constitute and should not be interpreted as an admission nor an agreement by the facility of the truth of the facts alleged on conclusions set forth in the statement of deficiencies. The plan of correction prepared for this deficiency was executed solely because it is required by provisions of State and Federal law. Without waiving the foregoing statement, the facility states that:</p> <ol style="list-style-type: none"> 1. With respect to the identified resident, R1 was evaluated in the emergency room and returned to the facility the following day with no new orders. The identified nurse received education and competency regarding insulin order verification prior to administration. 2. All residents with insulin orders were identified and insulin orders reviewed to ensure accurate entry in EHR that clearly states the dosage to be administered. 3. The facility's medication administration policy was revised to include insulin order verification by either a second nurse or trained individual. All facility licensed staff and trained medication aides will receive education and competency on insulin administration that will include how to read insulin orders in EMAR and how to verify dose with the second nurse or trained individual. All trained medication aids will receive education and competency on insulin dose verification. All education and competencies will be completed by April 1, 2021. 		

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F 760	<p>Continued From page 2</p> <p>supposed to have another nurse double check the insulin and amount and the other nurse was to co-sign with their initials. LPN-A further stated if there was no other nurse in the memory care unit the nurse was supposed to go downstairs to have the insulin verified before administrating it.</p> <p>On 3/2/21, at 1:04 p.m. RN-B stated the facility policy was after dialing the amount of prescribed insulin, the nurse was responsible to have another nurse do a second check to make sure it was the right amount before administering it. RN-A also stated at times, when there was no licensed nurse in the unit, she had the TMA double check and she would put the TMA initials and she acknowledged TMA-A had double checked the insulin she had administered during the morning shift today.</p> <p>On 3/2/21, at 1:20 p.m. LPN-B stated she would read the order in the computer then she would dial the amount and before administering it, usually another nurse was supposed to double check the insulin. LPN-B also stated if there was no other nurse at the unit, she would have the TMA double check before giving it. LPN-B further stated, "It's our policy to double when a nurse is not there, a TMA will check it and we even show them the blood sugars. I will put the initial of the double checker in the computer that's what I do."</p> <p>When interviewed on 3/2/21, at 1:40 p.m. the director of nursing (DON) stated it was the facility policy for a nurse who was administering insulin to have another nurse double check to ensure the correct dose and correct type of insulin has being drawn up before it was administered to a resident. "If you need to administer insulin, you find another nurse. You have them double check</p>	F 760	<p>4. The Director of Nursing or Designee will audit insulin administration on various shifts twice per week for one month and once per week for two months. The data collected will be presented to the QAPI committee by the Executive Director or designee. The data will be reviewed/discussed at regular QAPI meetings. At this time the QAPI committee will make the decision/recommendation regarding any necessary follow-up studies.</p>		

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F 760	<p>Continued From page 3</p> <p>the dosage and [insulin] vial." The nurse who provides the second check then co-signs the medication administration in the electronic medical record. The DON added, "[Errors] can happen if you go too fast, didn't get someone to cosign, or you get distracted." The DON acknowledged RN-A knew what the issue was, but did not follow the policy for a co-signer.</p> <p>When interviewed on 3/2/21, at 2:20 p.m., RN-A stated, "In big bold letters it said Lantus 100 units per milliliter then under that it said to give 20 units. I just saw 100 and gave that to [R1]." Upon discovering the error RN-A stated she immediately checked R1's blood sugar, vitals signs, and contacted the on-call physician. "The doctor said to send her out [to the hospital]." RN-A added, "I know with insulin there should be two people verifying the dose. At Birchwood we don't. It's just me doing nurse and [TMA] trained medication aide work. It would mean I have to go down stairs, show another nurse, and come back up. Just to give insulin." RN-A went on to state, "We don't have time and don't do that. If there is a nurse and TMA the TMA would just look at it, but they can't sign it out, since they are not nurses. I never work when there are TMA's there. If there are two people you should have two nurses check insulin [dosing]."</p> <p>The facility Medication Administration policy revised 6/24/2018, directed nurse, "Insulin is to be double checked with another nurse. The 2nd nurse will verify that the amount drawn up is correct and initial as the co-signer on the eMAR."</p>	F 760			

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CENTERS FOR MEDICARE & MEDICAID SERVICES

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F 000	<p>INITIAL COMMENTS</p> <p>On 3/2/21, an abbreviated survey was completed at your facility to conduct a complaint investigation. Your facility was found NOT to be in compliance with 42 CFR Part 483, Requirements for Long Term Care Facilities.</p> <p>The following complaint was found to be SUBSTANTIATED: H5200049C (MN00068952). A deficiency is being cited at F684.</p> <p>The facility's plan of correction (POC) will serve as your allegation of compliance upon the Department's acceptance.</p> <p>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.</p> <p>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.</p>	F 000			
F 760 SS=E	<p>Residents are Free of Significant Med Errors CFR(s): 483.45(f)(2)</p> <p>The facility must ensure that its- §483.45(f)(2) Residents are free of any significant medication errors. This REQUIREMENT is not met as evidenced by: Based on interview and document review, the facility failed to ensure 1 of 2 residents (R1) received the correct dose of insulin per physician orders, reviewed for insulin administration. In</p>	F 760	<p>F760</p> <p>The preparation of the following plan of correction for this deficiency does not</p>	4/1/21	

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

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F 760	<p>Continued From page 1</p> <p>addition, the facility failed to routinely follow their policy to ensure accurate insulin dosing, of having two licensed nurses checking the dose prior to administration. This practice had the potential to affect 9 residents currently receiving insulin.</p> <p>Findings include:</p> <p>R1's quarterly Minimum Data Set (MDS) dated 2/4/21, indicated R1 received insulin daily and had severely impaired cognition. R1's diagnosis included type 2 diabetes mellitus without complications.</p> <p>R1's February 2021 medication administration record (MAR) included, Insulin Glargine solution pen-injector 100 UNIT/ML inject 20 units subcutaneously (under the skin) one time a day at bedtime</p> <p>R1's Facility Reported Incident (FRI) dated 3/1/21, at 8:00 p.m. included, registered nurse (RN)-A had administered 100 units of Lantus insulin to R1 instead of the ordered 20 units. Upon discovering the error, R1's blood sugar was immediately checked. At 10:30 p.m. R1's blood sugar was found to be 66 (normal 60-100). The on-call physician was contacted and ordered R1 to be sent to the hospital for observation where R1 remained and returned to the facility on 2/29/21, evening shift.</p> <p>On 3/2/21, at 11:40 a.m. licensed practical nurse (LPN)-A assigned at the memory care unit stated nurses were supposed to compare the physician order in the computer with the label on the insulin pen to make sure they matched and would then dial the units the resident was to receive after priming the pen. LPN-A then stated nurses were</p>	F 760	<p>constitute and should not be interpreted as an admission nor an agreement by the facility of the truth of the facts alleged on conclusions set forth in the statement of deficiencies. The plan of correction prepared for this deficiency was executed solely because it is required by provisions of State and Federal law. Without waiving the foregoing statement, the facility states that:</p> <ol style="list-style-type: none"> 1. With respect to the identified resident, R1 was evaluated in the emergency room and returned to the facility the following day with no new orders. The identified nurse received education and competency regarding insulin order verification prior to administration. 2. All residents with insulin orders were identified and insulin orders reviewed to ensure accurate entry in EHR that clearly states the dosage to be administered. 3. The facility's medication administration policy was revised to include insulin order verification by either a second nurse or trained individual. All facility licensed staff and trained medication aides will receive education and competency on insulin administration that will include how to read insulin orders in EMAR and how to verify dose with the second nurse or trained individual. All trained medication aids will receive education and competency on insulin dose verification. All education and competencies will be completed by April 1, 2021. 		

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NAME OF PROVIDER OR SUPPLIER BIRCHWOOD HEALTH CARE CENTER			STREET ADDRESS, CITY, STATE, ZIP CODE 604 - 1ST STREET NE FOREST LAKE, MN 55025		
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F 760	<p>Continued From page 3</p> <p>the dosage and [insulin] vial." The nurse who provides the second check then co-signs the medication administration in the electronic medical record. The DON added, "[Errors] can happen if you go too fast, didn't get someone to cosign, or you get distracted." The DON acknowledged RN-A knew what the issue was, but did not follow the policy for a co-signer.</p> <p>When interviewed on 3/2/21, at 2:20 p.m., RN-A stated, "In big bold letters it said Lantus 100 units per milliliter then under that it said to give 20 units. I just saw 100 and gave that to [R1]." Upon discovering the error RN-A stated she immediately checked R1's blood sugar, vitals signs, and contacted the on-call physician. "The doctor said to send her out [to the hospital]." RN-A added, "I know with insulin there should be two people verifying the dose. At Birchwood we don't. It's just me doing nurse and [TMA] trained medication aide work. It would mean I have to go down stairs, show another nurse, and come back up. Just to give insulin." RN-A went on to state, "We don't have time and don't do that. If there is a nurse and TMA the TMA would just look at it, but they can't sign it out, since they are not nurses. I never work when there are TMA's there. If there are two people you should have two nurses check insulin [dosing]."</p> <p>The facility Medication Administration policy revised 6/24/2018, directed nurse, "Insulin is to be double checked with another nurse. The 2nd nurse will verify that the amount drawn up is correct and initial as the co-signer on the eMAR."</p>	F 760			



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically delivered

March 11, 2021

Administrator
Birchwood Health Care Center
604 - 1st Street Ne
Forest Lake, MN 55025

Re: Event ID: 2RUP11

Dear Administrator:

The above facility survey was completed on March 2, 2021 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 no violations of these rules promulgated under Minnesota Stat. section 144.653 and/or Minnesota Stat. Section 144A.10.

Electronically posted is the Minnesota Department of Health order form stating that no violations were noted at the time of this survey. The Minnesota Department of Health is documenting the State Licensing Correction Orders using federal software. Please disregard the heading of the fourth column which states, "Provider's Plan of Correction." This applies to Federal deficiencies only. There is no requirement to submit a Plan of Correction.

Please feel free to call me with any questions.

Sincerely,

A handwritten signature in black ink, appearing to read 'Melissa Poepping'.

Melissa Poepping, Health Program Representative Senior
Program Assurance | Licensing and Certification
Minnesota Department of Health
P.O. Box 64970
Saint Paul, Minnesota 55164-0970
Phone: 651-201-4117
Email: melissa.poepping@state.mn.us

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00853	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 03/02/2021
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NAME OF PROVIDER OR SUPPLIER BIRCHWOOD HEALTH CARE CENTER	STREET ADDRESS, CITY, STATE, ZIP CODE 604 - 1ST STREET NE FOREST LAKE, MN 55025
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2 000	<p>Initial Comments</p> <p>*****ATTENTION*****</p> <p>NH LICENSING CORRECTION ORDER</p> <p>In accordance with Minnesota Statute, section 144A.10, this correction order has been issued pursuant to a survey. If, upon reinspection, it is found that the deficiency or deficiencies cited herein are not corrected, a fine for each violation not corrected shall be assessed in accordance with a schedule of fines promulgated by rule of the Minnesota Department of Health.</p> <p>Determination of whether a violation has been corrected requires compliance with all requirements of the rule provided at the tag number and MN Rule number indicated below. When a rule contains several items, failure to comply with any of the items will be considered lack of compliance. Lack of compliance upon re-inspection with any item of multi-part rule will result in the assessment of a fine even if the item that was violated during the initial inspection was corrected.</p> <p>You may request a hearing on any assessments that may result from non-compliance with these orders provided that a written request is made to the Department within 15 days of receipt of a notice of assessment for non-compliance.</p> <p>INITIAL COMMENTS: On 3/2/21, an abbreviated survey was conducted to determine compliance with State Licensure. Your facility was found to be IN compliance with the MN State Licensure.</p> <p>The following complaint was investigated: H5200049C (MN00068952)</p>	2 000		

Minnesota Department of Health LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE	TITLE	(X6) DATE
Electronically Signed		03/16/21

Minnesota Department of Health

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 00853	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING _____	(X3) DATE SURVEY COMPLETED C 03/02/2021
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NAME OF PROVIDER OR SUPPLIER BIRCHWOOD HEALTH CARE CENTER	STREET ADDRESS, CITY, STATE, ZIP CODE 604 - 1ST STREET NE FOREST LAKE, MN 55025
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2 000	Continued From page 1 NO licensing orders were issued. The facility is enrolled in ePOC and therefore a signature is not required at the bottom of the first page of state form. Although no plan of correction is required, it is required that the facility acknowledge receipt of the electronic documents.	2 000		