



Protecting, Maintaining and Improving the Health of All Minnesotans

Electronically Delivered
June 30, 2026

Administrator
Tuff Memorial Home
505 EAST 4TH STREET
HILLS, MN 56138

RE: CCN: 245548
Cycle Start Date: May 26, 2026

Dear Administrator:

On May 26, 2026, the Minnesota Department of Public Safety, 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.

Sincerely,

A handwritten signature in black ink that reads 'Kamala Fiske-Downing'.

Kamala Fiske-Downing
Compliance Analyst | Federal Enforcement
Health Regulation Division
Minnesota Department of Health
Kamala.Fiske-Downing@state.mn.us
Office: 651-201-4112



Protecting, Maintaining and Improving the Health of All Minnesotans

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June 4, 2026

Administrator
Tuff Memorial Home
505 EAST 4TH STREET
HILLS, MN 56138

RE: CCN:245548

Cycle Start Date: May 27, 2026

Dear Administrator:

On May 27, 2026, 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" and/or an "E" tag), i.e., the plan of correction should be directed to:

Nicole Dahl, RN, Regional Operations Supervisor
Marshall District Office
Health Regulation Division
Minnesota Department of Health
1400 East Lyon Street, Suite 102
Marshall, Minnesota 56258-2504
Email: Nicole.Dahl@state.mn.us
Office: 507-476-4230

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 August 27, 2026 (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).

In addition, if substantial compliance with the regulations is not verified by November 27, 2026 (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)

In accordance with 42 CFR 488.331 and Minnesota Statute 144A.10 subd 15, 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: <https://forms.web.health.state.mn.us/form/NHDisputeResolution>

This request must be sent within the same ten calendar days you have for submitting an ePoC for the cited deficiencies. 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.

A copy of the Department's informal dispute resolution policies is posted on the MDH Information Bulletin website at:

https://www.health.state.mn.us/facilities/regulation/infobulletins/ib04_8.html

INDEPENDENT INFORMAL DISPUTE RESOLUTION (INDEPENDENT IDR)

In accordance with 42 CFR § 488.431 and Minnesota Statute 144A.10 subd 16, when a CMP subject to being collected and placed in an escrow account is imposed, you have one opportunity to question cited deficiencies through an Independent IDR process. You may also contest scope and severity assessments for deficiencies which resulted in a finding of SQC or immediate jeopardy. 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:

<https://forms.web.health.state.mn.us/form/NHDisputeResolution>

A facility may not use both IDR and independent IDR for the same deficiency citation(s) arising from the same survey unless the IDR process was completed prior to the imposition of the CMP. This request must be sent within ten calendar days of receipt of this offer. An incomplete Independent IDR process will not delay the effective date of any enforcement action.

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:

Travis Z. Ahrens

State Fire Safety Supervisor

Health Care & Correctional Facilities

MN Department of Public Safety-Fire Marshal Division

445 Minnesota St., Suite 145

St. Paul, MN 55101

Email: travis.ahrens@state.mn.us

Web: www.sfm.dps.mn.gov

Cell: 1-507-308-4189

Feel free to contact me if you have questions.

Sincerely,



Kamala Fiske-Downing
Compliance Analyst | Federal Enforcement
Health Regulation Division
Minnesota Department of Health
Kamala.Fiske-Downing@state.mn.us
Office: 651-201-4112

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTIONS		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245548		(X2) MULTIPLE CONSTRUCTION A. BUILDING B. WING		(X3) DATE SURVEY COMPLETED 05/26/2026	
NAME OF PROVIDER OR SUPPLIER Tuff Memorial Home				STREET ADDRESS, CITY, STATE, ZIP CODE 505 EAST 4TH STREET , HILLS, Minnesota, 56138			
(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
E0000	Initial Comments On 5/26/26, a survey for compliance with CFR §483.73, Appendix Z, Emergency Preparedness Requirements was conducted during a standard recertification survey. The facility was IN compliance. The facility is enrolled in ePOC and therefore a signature is not required at the bottom of the first page of the CMS-2567 form. Although no plan of correction is required, it is required that the facility acknowledge receipt of the electronic documents.			E0000			
F0000	INITIAL COMMENTS On 5/26/26, a federal recertification survey was conducted using the Risk-Based Survey (RBS) process at your facility. Your facility was IN compliance with §42 CFR 483, Subpart B, Requirements for Long Term Care Facilities. The facility is enrolled in ePOC and therefore a signature is not required at the bottom of the first page of the CMS-2567 form. Although no plan of correction is required, the facility must acknowledge receipt of the electronic documents.			F0000			

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 reverse for further 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.

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE	TITLE	(X6) DATE

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTIONS		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 245548		(X2) MULTIPLE CONSTRUCTION A. BUILDING 01 - MAIN BUILD... B. WING		(X3) DATE SURVEY COMPLETED 05/27/2026	
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K0000 Bldg. 01	INITIAL COMMENTS FIRE SAFETY An annual Life Safety Code survey was conducted on 05/27/2026 by the Minnesota Department of Public Safety, State Fire Marshal Division. At the time of this survey, Tuff Memorial Home was found not in compliance with the requirements for participation in Medicare/Medicaid at 42 CFR, Subpart 483.70(a), Life Safety from Fire, and the 2012 edition of National Fire Protection Association (NFPA) 101, Life Safety Code (LSC), Chapter 19 Existing Health Care and the 2012 edition of NFPA 99, Health Care Facilities Code. 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. 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. PLEASE RETURN THE PLAN OF CORRECTION FOR THE FIRE SAFETY DEFICIENCIES (K-TAGS) TO: If PARTICIPATING IN THE E-POC PROCESS, a paper copy of the plan of correction is not required. Healthcare Fire Inspections State Fire Marshal Division 445 Minnesota St., Suite 145			K0000			06/05/2026

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 reverse for further 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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K0000 Bldg. 01	Continued from page 1 St. Paul, MN 55101-5145, OR By email to: FM.HC.Inspections@state.mn.us THE PLAN OF CORRECTION FOR EACH DEFICIENCY MUST INCLUDE ALL OF THE FOLLOWING INFORMATION: A detailed description of the corrective action taken or planned to correct the deficiency. Address the measures that will be put in place to ensure the deficiency does not reoccur. Indicate how the facility plans to monitor future performance to ensure solutions are sustained. Identify who is responsible for the corrective actions and monitoring of compliance. The actual or proposed date for completion of the remedy. Tuff Memorial Home was constructed as follows: The original building was constructed in 1959, is one-story, has a partial basement, is fully fire sprinkler protected and is of Type II(111) construction; The 1st Addition was constructed in 1962, is one-story, has no basement, is fully fire sprinkler protected and is of Type II(111) construction; The 2nd Addition was constructed in 1975, is one-story, has no basement, is fully fire sprinkler protected and is of Type II(111) construction; The 3rd Addition was constructed in 1988, is one-story, has a full basement, is fully fire sprinkler protected and is of Type V(111) construction; The 4th Addition was constructed in 1998, is one-story, has no basement, is fully fire sprinkler protected and is of Type V(000) construction.			K0000			06/05/2026

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K0000 Bldg. 01	Continued from page 2 The facility has a capacity of 48 beds and had a census of 40 at the time of the survey.			K0000			06/05/2026
K0920 SS = E Bldg. 01	The requirement at 42 CFR, Subpart 483.70(a) is NOT MET as evidenced by: Electrical Equipment - Power Cords and Extens CFR(s): NFPA 101 Electrical Equipment - Power Cords and Extension Cords Power strips in a patient care vicinity are only used for components of movable patient-care-related electrical equipment (PCREE) assemblies that have been assembled by qualified personnel and meet the conditions of 10.2.3.6. Power strips in the patient care vicinity may not be used for non-PCREE (e.g., personal electronics), except in long-term care resident rooms that do not use PCREE. Power strips for PCREE meet UL 1363A or UL 60601-1. Power strips for non-PCREE in the patient care rooms (outside of vicinity) meet UL 1363. In non-patient care rooms, power strips meet other UL standards. All power strips are used with general precautions. Extension cords are not used as a substitute for fixed wiring of a structure. Extension cords used temporarily are removed immediately upon completion of the purpose for which it was installed and meets the conditions of 10.2.4. 10.2.3.6 (NFPA 99), 10.2.4 (NFPA 99), 400-8 (NFPA 70), 590.3(D) (NFPA 70), TIA 12-5 This STANDARD is NOT MET as evidenced by: Based on observation and staff interview, the facility failed to maintain the usage of electrical adaptive devices NFPA 99 (2012 edition), Health Care Facilities Code, sections 10.5.2.3.1 and 10.2.4.2.1, NFPA 101 (2012 edition), Life Safety Code, section 9.1.2, NFPA 70, (2011 edition), National Electrical Code, sections 400.8, and UL 1363. This deficient finding could have a patterned impact on the residents within the facility. Findings include: 1. On 05/27/2026 at 10:03 AM, it was revealed by observation that a dehumidifier was plugged in with an extension in the hvac room in the basement. 2. On 05/27/2026 at 10:19 AM, it was revealed by observation that there was a refrigerator plugged			K0920	K0920 Electrical Equipment - Power cords and extension cords Corrective Action for Residents Affected by the deficient practice- The deficient practices were corrected immediately upon discovery. The extension cord used to power the dehumidifier in the HVAC room was removed and the dehumidifier was connected to an approved permanent electrical outlet. In the Med room, the daisy chained power strips supplying power to the refrigerator were removed and the refrigerator was connected to an appropriate electrical receptacle in accordance with NFPA and manufacturer requirements. No residents were adversely affected by these deficient practices. How the facility will identify other residents who could be affected- All residents could potentially be affected by improper use of electrical adaptive devices. A facility wide inspection of electrical equipment, extension cords, and power strips was conducted to identify additional instances of improper use, including extension cords used in place of permanent wiring and daisy chained power strips. Measures put into place to ensure the deficient practice will not recur- The maintenance director and department heads were educated regarding NFPA 99, & 70, education included the prohibition of extension cords as a substitute for permanent wiring and the prohibition of daisy chaining power strips. The facilities environmental and life safety inspection checklist was updated to include the monitoring for improper use of extension cords, power strips, and patient care related electrical equipment (PCREE) How the facility will monitor its corrective action- The maintenance director or designee will conduct		06/05/2026

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K0920 SS = E Bldg. 01	Continued from page 3 into a power strip that was daisy chained into another power strip in the Med Room. An interview with the Maintenance Director verified these deficient findings at the time of discovery.			K0920	Continued from page 3 weekly audits of all departments for four weeks and monthly audits thereafter for two months to verify compliance with electrical safety requirements. Audit findings will be documented and reported to the QAPI committee. Any identified issues will be corrected immediately, and additional education will be provided as necessary.		06/05/2026
K0741 SS = D Bldg. 01	Smoking Regulations CFR(s): NFPA 101 Smoking Regulations Smoking regulations shall be adopted and shall include not less than the following provisions: (1) Smoking shall be prohibited in any room, ward, or compartment where flammable liquids, combustible gases, or oxygen is used or stored and in any other hazardous location, and such area shall be posted with signs that read NO SMOKING or shall be posted with the international symbol for no smoking. (2) In health care occupancies where smoking is prohibited and signs are prominently placed at all major entrances, secondary signs with language that prohibits smoking shall not be required. (3) Smoking by patients classified as not responsible shall be prohibited. (4) The requirement of 18.7.4(3) shall not apply where the patient is under direct supervision. (5) Ashtrays of noncombustible material and safe design shall be provided in all areas where smoking is permitted. (6) Metal containers with self-closing cover devices into which ashtrays can be emptied shall be readily available to all areas where smoking is permitted. 18.7.4, 19.7.4 This STANDARD is NOT MET as evidenced by: Based on document review, observation, and staff interview, the facility failed to enforce smoking policies per NFPA 101 (2012 edition), Life Safety Code, sections 19.7.4. This deficient finding could have an isolated impact on the occupants within the facility. Findings include:			K0741	KO741 Smoking Regulations Corrective actions The facility immediately obtained and placed a metal container with a self closing lid in the designated smoking area on 5/27/2026. The smoking area was inspected to ensure compliance with NFPA 101 section 19.7.4 No residents were adversely affected by the deficient practice. How the facility will identify residents who could be affected. No residents utilize the designated employee smoking area, The designated smoking area and related safety equipment were inspected to verify compliance with life safety code requirements. Measures put into place to ensure the deficient practice will not recur The facility maintenance supervisor was educated on NFPA 101 smoking regulation requirements, including the requirement that a metal container with a self-closing lid be readily available in all designated smoking areas. The facility smoking area checklist has been updated to include verification of the self-closing metal receptacle. How the facility will monitor corrective actions The maintenance director or designee will inspect the designated smoking area weekly for four weeks and monthly thereafter for two months to verify the required self-closing metal receptacle remains in place and functional. Findings will be documented and reported to the Quality Assurance and Performance Improvement (QAPI) Committee. Any identified concerns will be corrected immediately.		06/05/2026

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K0741 SS = D Bldg. 01	Continued from page 4 On 05/27/2026 at 10:10 AM, it was revealed by observation that there was not a metal container with a self-closing cover in the designated smoking area. An interview with the Maintenance Director verified this deficient finding at the time of discovery.			K0741			06/05/2026