

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 465139	(X2) MULTIPLE CONSTRUCTION A. Building B. Wing	(X3) DATE SURVEY COMPLETED 01/21/2026
NAME OF PROVIDER OR SUPPLIER Monument Healthcare Millcreek		STREET ADDRESS, CITY, STATE, ZIP CODE 1201 East 4500 South Salt Lake City, UT 84117	
For information on the nursing home's plan to correct this deficiency, please contact the nursing home or the state survey agency.			
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (Each deficiency must be preceded by full regulatory or LSC identifying information)		
<p>F 0694</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Few</p>	<p>Provide for the safe, appropriate administration of IV fluids for a resident when needed.</p> <p>**NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY** Resident 2 was admitted to the facility on [DATE] with diagnoses which consisted of displaced bimalleolar fracture of right lower extremity, displaced avulsion fracture of right talus, chronic pulmonary edema, congestive heart failure, chronic kidney disease, kidney transplant status, unspecified protein-calorie malnutrition, fistula of the intestine, contusion of the abdominal wall, dyspnea, disorders of phosphorus metabolism, and hypokalemia. On 1/20/26 through 1/21/26 resident 2's medical records were reviewed. On 10/29/25, resident 2's admission Minimum Data Set (MDS) Assessment documented a Brief Interview for Mental Status (BIMS) score of 15, which indicated that the resident was cognitively intact. On 10/24/25, resident 2's physician ordered TPN 1600 milliliter (ml) to run continuously in the evening per the pharmacy instructions. The pharmacy order summary documented the following for resident 2's TPN:Start Volume - 100 mlStart Rate - 62.5 ml/hour (hr.)Start Period - 1.6 hrs.Run Volume - 1600 mlRun Rate - 125 ml/hr.Run Period - 12.8 hrs.End Volume - 100 mlEnd Rate - 62.5 ml/hr.End Period - 1.6 hrs.Overflow Volume - NoneContainer Type - Single ChamberCompounded Volume - 1800 mlThe order summary was last updated on 11/20/25. It should be noted that the total volume to be infused in the order summary was 1800 ml with a total run period of 16 hrs. On 12/29/25, resident 2's pharmacy label for the TPN documented Infuse intravenously 1800 ml continuously in the evening. Run at 125 ml/hr. over 12 hrs. It should be noted that the order summary start volume and start rate and the end volume and end rate did not match the pharmacy label nor did the combined run period times equaling 16 hrs. Additionally, it was unclear whether the pharmacy label had not matched the original physician orders since October 2025. Resident 2's progress notes revealed the following: a. On 12/28/25 at 11:01 PM, the Alert Charting note documented, Entered room to check on resident and found her connected to the TPN bag. The bag is empty. Bag is not on the pump. She has received her 1600 ml of TPN already although it is scheduled for 12 hours. VS [vital signs] assessed. Dr on call notified. BS [blood sugar] checked. Neuro check at her baseline. Lung sound clear x's [times] 4. Resident refuses to have her family called. States 'you can call them in the morning. Don't bother them this time of night.' Dr on call-DON [Director of Nursing]- and nurse manager notified. b. On 12/29/25 at 12:18 AM, the Order Note documented, .Received TPN over a short time period. New orders: STAT [immediate] CMP [complete metabolic panel] in the am, Head of bed elevated at all times, assess lungs every 2 hours, monitor every 2 hours for altered mental status, vital signs every 2 hours. Resident agrees with change in plan of care. New orders noted and implemented. c. On 12/29/25 at 9:26 AM, a physical medicine and rehabilitation follow up note documented, Patient assessed w/ [name of Medical Doctor omitted] this visit. [Resident 2] is observed in bed. Noted to be lethargic with altered mental status. Arousable to loud voice and tactile stimulation; however, patient closes eyes almost immediately upon opening. She reports constant, sharp chest pain, non radiating. Vitals stable on 3L [liters oxygen] NC [nasal cannula], denies SOB [shortness of breath]. d. On 12/29/25 at 6:01 PM, the Alert Charting note documented, Report</p> <p>(continued on next page)</p>		

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.

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE	TITLE	(X6) DATE
FORM CMS-2567 (02/99) Previous Versions Obsolete	Event ID: 465139	Facility ID: If continuation sheet Page 1 of 7

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<p>F 0694</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Few</p>	<p>RN 1 stated that resident 2 was alert and oriented, talking and appeared to be fine. RN 1 stated that the MD ordered to monitor resident 2 every 2 hours for lung sounds, orientation, and VS within normal range. RN 1 stated that complications with the bolus infusion of the TPN could include cardiac failure, lungs full of water, altered mental status, and that was what she was monitoring for. On 1/21/26 at 9:00 AM, a follow-up interview was conducted with the DON and ADM. The DON confirmed that per the TPN label the volume of 1600 ml infusing at a rate of 125 ml/hrs. should have taken 14.4 hrs. to complete and not 12 hours as documented on the label. The DON stated that the nurse should have clarified the order discrepancies with the pharmacy and then followed-up with the MD. The DON stated that she recalled the order being 125ml over 12 hrs. and was not sure if the order ever changed during the course of the treatment. The DON stated that she could see that there was some confusion with the order as it was written. On 1/21/26 at 1:17 PM, a telephone interview was conducted with Nurse Practitioner (NP) 1 and Medical Doctor (MD) 1. MD 1 stated that he and NP 1 were the consultant secondary team, and did not write any medication orders for resident 2. NP 1 stated that when she assessed resident 2 on 12/29/26, the resident complained of chest pain and appeared a bit altered from baseline. NP 1 stated that the nursing staff were aware of resident 2's condition. MD 1 stated that they spoke with the DON and the floor nurse caring for resident 2 and reported the new complaints of chest pain. MD1 stated that the DON informed him that resident 2 had received a bolus of TPN and they were doing an investigation into the situation. MD 1 stated that the nursing staff was aware of resident 2's change in condition and that it could possibly be caused by the TPN infusion. MD 1 stated that resident 2's primary team physician was aware of resident 2's condition. Review of the facility policy on Parenteral Nutrition (PN) documented the following under general guidelines: . 3. Guidelines for administering are as follows: a. The rate tapers upward for 1-2 hours when starting the infusion. b. Then the PN runs at a set rate for a determined time. c. The rate tapers downward for 1-2 hours before the infusion is stopped or discontinued. d. The time intervals and tapering rates will be determined by the physician or the pharmacist. e. The bag is then disconnected from the catheter and discarded. f. The catheter is flushed with saline/heparin per protocol. 4. Never stop or discontinue parenteral nutrition suddenly. a. The PN rate must be tapered downward over several hours to allow the pancreas to adjust to the decrease in glucose intake (and the subsequent decreased need for insulin). This will help prevent hypoglycemia. The policy was last updated 2/1/24.</p>		

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<p>F 0697</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Few</p>	<p>Provide safe, appropriate pain management for a resident who requires such services.</p> <p>**NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY** Based on interview and record review it was determined that the facility did not ensure that residents received the treatment and care in accordance with professional standards of practice, related to pain management. Specifically 1 of 4 residents sampled, was hospitalized due to opiate toxicity. Resident Identifier: 4 Findings Included: Resident 4 was admitted to the facility on [DATE] with diagnoses which included sepsis, obstructive and reflux uropathy, and calculus of kidney. Review of resident 4's medical record was completed on 1/21/26. On 1/1/26 at 10:47 AM, a Physician's Progress Admit Note for resident 4 revealed that a medication reconciliation was done with the patient's family and his primary care physician's records. Morphine tablets (immediate release, 7.5 milligrams (mg) every 6 hours as needed), were ordered for resident 4's dorsalgia. On 1/2/26 at 1:49 PM, a Medical Doctor (MD) History and Physical Note for resident 4 stated, Patient with large kidney stones. Patient complains of ongoing pain and wants more morphine than he is getting. On 1/3/26 at 8:28 AM, a Nursing Progress Order Note for resident 4 stated, .Give 1/2 tab PO [by mouth] every 3 hours for pain AND Give 1 tablet by mouth every 3 hours as needed for pain. On 1/3/26, a physician's order for resident 4 revealed the following Morphine Sulfate Oral Tablet 15 mg. Give 0.5 tablet by mouth every 3 hours as needed for pain. On 1/3/26, a physician's order for resident 4 revealed the following Morphine Sulfate Oral Tablet 15 MG. Give 1 tablet by mouth every 3 hours as needed for pain. It should be noted the new physician order resulted in an increase of 150 mg of Morphine every 24 hours as needed. Further review indicated that, based on the prescribed frequency and dosage, there was a potential for a total of 180 mg of Morphine to be administered within a 24-hour period. On 1/5/26 at 2:51 PM, a Physician's Follow Up Note for resident 4 revealed the following. Interval History: Patient seen in his room patient relays to me that his family felt like his [sic] he was slurring his words today. Patient has a new cough and feels congested in the lungs Pain management adequate with morphine.Details : Chief Complaint: slurred speech.Respiratory : Lungs CTA, Other: He has a congested cough.Assessment/Plan:.Ongoing management of nephrostomy, pain management, awaiting surgery. Pain management with morphine. I think there is some intermittent dysarthria caused by the morphine and we need to back off on this unless needed for pain levels higher than 4. It should be noted that an updated morphine was not located in the medical record. On 1/6/26 at 5:26 AM, a Nursing Progress Order Note for resident 4 stated, Resident is lethargic and wasn't speaking clearly. He slept all night but was smiling and compliant with medication and breathing treatment. BP [blood pressure] 167/95, HR [heart rate] 122, oxygen with CPAP [Continuous Positive Airway Pressure] is 88, and temp [temperature] is 97.7. Orders for chest x-ray are in but [name redacted] never arrived, resident complained yesterday of SOB [shortness of breath] and coughing. DON [Director of Nursing], ADON [Assistant Director of Nursing], and MD notified. On 1/15/26 at 4:16 PM, a Nursing Progress Note for resident 4 states, Pt with low O2 [oxygen] sat [saturation] at 83%; groggy and dry mouthed; not eating meals well; lungs with rhonchi bilaterally; nebulizer treatment given, O2 placed at 2L [liter]/NC [nasal cannula]; effective; o2 SATs increased to 93-94%; NP [Nurse Practitioner] notified or change in status. On 1/17/26 at 2:08 AM, a Nursing Progress Note for resident 4 states, Pt [patient] yelling for help, upon entering room pt found between the bathroom and his bed on his Lt [left] side. When asked what happened pt stated I rolled out of bed, after careful evaluation, no injuries noted. Assisted x [times] 3 staff back into bed. Bed in lowest position, call light in reach. Pt instructed to call for assistance before getting out of bed, Pt stated I didn't know I was paralyzed MD, Wife, Administrator, DON, Unit Manager notified. On 1/17/26 at 8:18 AM, resident 4's MAR revealed 1 tablet of 15 mg of Morphine Sulfate Oral Tablet was given. Pain Level</p> <p>(continued on next page)</p>		

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<p>F 0697</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Few</p>	<p>Noted as 10. On 1/17/26 at 10:30 PM, resident 4's MAR revealed 1 tablet of 15 mg of Morphine Sulfate Oral Tablet was given. Pain Level Noted as 7. On 1/18/26 at 7:38 AM, resident 4's MAR revealed 1 tablet of 15mg of Morphine Sulfate Oral Tablet was given. Pain Level Noted as 4. It should be noted that a total of 45 mg of Morphine Sulfate was administered in a 24 hour period. On 1/18/26 at 9:09 AM, a nursing progress note for resident 4 stated, Patient appears lethargic at this time. Oxygen saturation is unable to be obtained due to cold extremities; hands are cold to touch. Staff reported significant changes in patient condition compared to yesterday. Per report, patient experienced low oxygen saturation overnight. Oxygen has been titrated up to 5 L/min [minute] however, oxygen saturation remains unable to be obtained despite multiple attempts. Based on current assessment findings, including lethargy, inability to obtain oxygen saturation, cold extremities, and reported decline since yesterday. Patient is being transferred to [name redacted] via [name redacted] transportation. Dr. [name redacted] and management were notified of patient condition and transfer. Spouse, wife [name redacted], was notified; no answer-voicemail left requesting a call back. Patient's son, [name redacted], was also notified; no answer-voicemail left requesting call back for notification. Medication list, transfer records, and discharge documentation were sent with the patient. At time of transfer, vital signs were as follows: blood pressure 112/81 mmHg, temperature 97.1 F [Fahrenheit], pulse 78 bpm [beats per minute] , respirations 24 breaths per minute. Oxygen saturation remains unable to be obtained at this time. Blood sugar is 104 mg/dL [milligrams per deciliter]. Patient transferred in stable condition with ongoing concerns requiring further evaluation in the emergency department [ED]. The hospital ED provider notes dated 1/18/26 at 11:58 am, revealed the following. The patient arrives critically ill with hypoxia down to 68%, slow respiratory rate, confusion he will open his eyes. He is otherwise not very responsive. The patient is in critical condition, this patient's condition is life-threatening due to hypoxia, with ongoing risk of organ failure. He has been getting extra doses above the 0.5 tablets of morphine he supposed to be getting at least once a day maybe twice a day he has Narcan responsive and wakes up and his respiratory rate improved but his oxygenation does not, his blood pressure also improved with Narcan. the patient will be admitted to the intensive care unit with multiple diagnoses which include hypoxia, sepsis, and opiate toxicity. The following medications were administered: naloxone (NARCAN) 2mg in dextrose 5% (D%W) 500mL DRIP (has no administration in time range) .naloxone (NARCAN)0.4 mg/mL injection 0.2 mg (0.2 mg intravenously Given 1/18/26 1055) .naloxone (NARCAN)0.4 mg/mL injection 0.2 mg (0.2 mg intravenously Given 1/18/26 1230) On 1/21/26 at 9:19 AM, an interview with Licensed Practical Nurse (LPN) 1. revealed the following. LPN 1 stated when the physician gives an order, the nurse should verbally repeat the order back to the physician for verification. Once verification is completed the order is entered into the resident's medical record. LPN 1 stated he typically followed up with a provider if he encountered a morphine order with a frequency exceeding every four to six hours. LPN 1 stated that when administering pain medication, he evaluated parameters including pain levels, timing, the previous dose, and the resident's response. LPN 1 stated the resident should be able to verbalize the request for medication. LPN 1 stated that pain levels were subjective and that residents occasionally requested medication despite low pain levels. LPN 1 stated he would attempt to offer alternative interventions, but if the resident insisted, he administered the medication as prescribed. LPN 1 stated he had been resident 4's nurse previously when he returned back to work on 1/18/26, he was informed during shift report resident 4 had declined. LPN 1 stated that during the shift's initial assessment, resident 4 was alert but lethargic and requested a higher dose of morphine despite the LPN 1 offering a lower dose. LPN stated that his change of condition came within 3 hours. LPN 1 stated he did not follow up with the provider</p> <p>(continued on next page)</p>		

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