

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:  075387	(X2) MULTIPLE CONSTRUCTION A. Building B. Wing	(X3) DATE SURVEY COMPLETED  03/17/2026
NAME OF PROVIDER OR SUPPLIER  Autumn Lake Healthcare at Norwalk		STREET ADDRESS, CITY, STATE, ZIP CODE  34 Midrocks Drive Norwalk, CT 06851	

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 0684</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Few</p>	<p>Provide appropriate treatment and care according to orders, resident's preferences and goals.</p> <p><b>**NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY**</b> Based on review of the clinical record reviews, observations, facility policy and staff interviews for 1 of 7 residents ( Resident #1 ) reviewed for choices, the facility failed to ensure that a physician's order regarding leave of absence (LOA) was followed, and for 2 of 4 residents reviewed for accidents (Residents #17 and #130), the facility failed to ensure a nursing assessments were completed following a near-fall event and a change of condition and for 1 of 1 resident (Resident # 38) reviewed for communication, the facility failed to ensure staff appropriately identified a venous access site, obtain the correct physician orders for its use. The findings included:</p> <p>Resident #17's diagnoses included orthopedic conditions, phantom limb syndrome with pain, cervical disc (discs that provide space between the bones of the neck) degeneration, chronic pain syndrome, cervical radiculopathy (neck compression that causes pain to radiate to the arms or changes feeling in the arms), heart failure, asthma, bipolar disorder, depression, anxiety, and schizoaffective disorder (a chronic mental health condition characterized by a combination of schizophrenia symptoms, such as hallucinations, delusions, disorganized speech, and mood disorders, such as mania or depression).</p> <p>A Medicare 5-Day Minimum Data Set (MDS) assessment dated [DATE] indicated Resident #17 was cognitively intact. The resident used no assistive device for mobility, had a reduced range of motion on one side of the upper extremities, required set-up assistance for eating and showering, moderate assistance for toilet hygiene and repositioning, and was independent with oral hygiene, dressing, and transfers. The assessment also identified that the resident was prescribed multiple psychoactive (medications that alter brain function and can change perception, mood, behavior, and consciousness) medications, opioids (strong pain medications), and an antipsychotic medication.</p> <p>On 11/12/25, Resident #17 experienced a witnessed fall, with subsequent identification of non-displaced fractures of several toes on the left foot.</p> <p>Nursing assessments were completed on 11/12/25, along with a change-in-condition nursing note describing the incident and assessment findings. A follow-up evaluation was conducted on 11/13/25 by an advanced practice registered nurse (APRN) who ordered a left foot x-ray to rule out fracture due to focal (specific spot or area) pain in the foot.</p> <p>A Resident Care Plan (RCP) update, dated 11/12/25, included interventions such as orthostatic blood pressure monitoring (checking blood pressure with changes in position) each shift for three days, a medical work-up, and monitoring for signs and symptoms of latent (present but not yet showing signs or symptoms) injury or pain.</p> <p>A change-in-condition note dated 11/14/25 at 4:52 PM indicated the resident was evaluated by orthopedics, and a repeat x-ray confirmed fractures of the third, fourth, and fifth toes of the left foot. (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
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<p>F 0684</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Few</p>	<p>change the IV administration set (tubing)every 96 hours.</p> <p>The admission Minimum Data Set (MDS) assessment dated [DATE] indicated Resident #38 was cognitively intact had an intravenous (IV) device and was receiving intravenous(IV)medication.</p> <p>Resident #38's care plan dated 2/23/2026 indicated Resident #38 was receiving IV medications for an infection with interventions including documenting and to report to the physician any signs of redness at the IV insertion site.</p> <p>On 3/10/2026 at 10:40 AM an interview and record review with charge nurse LPN #1 identified Resident #38 had orders for a central line catheter for administration of antibiotic medication intravenously ( a central line is the longest IV catheter that reaches large veins near the heart, designed for long-term use, high-volume fluid resuscitation, and administration of caustic medications). While reviewing the paper clinical record with LPN #1, a loose paper document labeled PICC/Midline Nursing Documentation from an outside IV company, placement company dated 3/03/2026 indicated Resident #38 had a midline catheter(an intermediate-length venous access line inserted into an upper arm vein with its tip stopping near the axilla, used for 1&amp;ndash;4 weeks of intravenous therapies such as antibiotics) placed in the hospital but the line was pulled out by the Resident # 38 and the IV Nurse was there to replace the midline catheter. A 12 centimeter (cm) long midline catheter (one valve) was placed in a vein in the middle of the inside of the left arm with notes indicating to provide midline catheter care per protocol and post procedure instructions were provided to a facility nurse(not named). LPN # 1 reviewed the hospital discharge summary and the Intra-Agency Report upon surveyor request, and it indicated Resident #38 was discharged to the facility with a midline catheter (not a central line). LPN #1 indicated not knowing why the admitting nurse obtained physician orders for a Central Line Catheter. LPN #1 was unable to locate any physician's orders in the clinical record for dressing changes to the IV site and called for the nursing supervisor.</p> <p>On 3/10/26 at 10:42 AM a clinical record review, interview and observation with the nurse manager/Supervisor RN #1 indicated Resident #38 physician's orders were for a central line and the resident did not receive anything s/he should not have. Review of the physician's orders indicated not only flush the catheter with normal saline but Heparin as well. RN #1 indicated the facility does not use Heparin in any IVs as the practice ended years ago. She/he further indicated that when the physician's orders are electronically by the nurse the prepopulated orders for heparin need to be removed. RN #1 pulled up the standing physician's orders for a midline catheter and the same normal saline flush as the central line was what would have been ordered.</p> <p>An observation of the resident room found a labeled bag of saline syringes for IV-line flushing, and the observations of the medication rooms by surveyors identified only normal saline syringes containing 10ml of solution for IV flushing, no heparin syringes were found. Further review of the orders identified no documentation for a dressing change was completed since admission; no documentation of the IV line being flushed, or consistent documentation every shift as there were no orders on the Medication/Treatment Administration records( MAR/TAR) indicating how to take care of the IV devise and document that care was completed per the physician's orders and facility protocol. Review of the admission orders for the central line catheter identified the scheduling of the dressing changes and flushes were not completed. An observation at 11:10 AM with RN #1 in the resident's room identified Resident #38 had a clear dressing over the insertion site of the midline catheter in the inner left elbow dated 3/9/2026 and the catheter had a manufactural label that stated MIDLINE. RN #1 indicated s/he would inform the APRN and obtain new physician's orders.</p> <p>(continued on next page)</p>		

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<p>F 0684</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Few</p>	<p>On 3/10/2026 ( no time indicated) subject to surveyor inquiry, physician orders were obtained for a Midline catheter(long peripheral catheter, not a central catheter) directed to flush the catheter with 10ml of normal saline solution, to change the catheter dressing 24 hours after insertion; on admission, and weekly using a transparent dressing , to observe the IV site every 2 hours during continuous therapy, every shift during intermittent therapy or when not in use.</p> <p>The facility policy labeled( Midline/Extended Dwell Catheter Flushing given onsite indicated a physician's order is required for vascular access devices with orders specific to the flushing solution and documentation of the procedures in the resident's clinical record will include date and time, site assessment flushing agent and volume amount used, any difficulty flushing and intervention, the resident's response to the procedure and any resident/caregiver education provided.</p> <p>The facility policy labeled Midline/Extended Dwell Catheter Dressing Change given onsite indicated the physician IV therapy order for care and maintenance is required, a transparent dressing is preferred referring to the IV order form for the frequency of the change utilizing sterile technique once the soiled dressing is removed. Assessment of the vascular access device included checking the site at least every 2 hours during a continuous infusion, before during and after medication administration, during dressing changes at a minimum of once per shift when not in use and at prescribed intervals if complications are observed.</p> <p>Resident #1's diagnoses included type 2 diabetes mellitus, depressive episodes, schizophrenia, anxiety disorder, chronic kidney disease, end-stage renal disease (ESRD) requiring hemodialysis, weakness, chronic pulmonary edema, mild cognitive impairment of unknown etiology, and dementia with psychotic disturbance.</p> <p>A physician's order dated 11/15/25 directed Resident #1 may leave the facility on leave of absence (LOA) with a responsible party and medications.</p> <p>A quarterly Minimum Data Set (MDS) assessment dated [DATE] indicated Resident #1 was cognitively intact with a Brief Inventory of Mental Status (BIMS) Summary Score of 15. The resident used a walker and wheelchair for mobility, required varying levels of assistance with activities of daily living, and was receiving specialized treatment for ESRD.</p> <p>A psychiatric advanced practice registered nurse (APRN #1) evaluation dated 12/8/25 indicated the resident was psychiatrically stable with no acute concerns and no risk to self or others.</p> <p>A nursing note dated 12/11/25 indicated Resident #1 informed staff he/she was going to the lobby but subsequently left the facility via a ride-share without notifying staff or formally signing out on LOA, which was inconsistent with the physician's order requiring a responsible party accompaniment.</p> <p>Facility staff identified that the resident had left when he/she was no longer present in the lobby and initiated efforts to determine the resident's whereabouts. Staff contacted the resident by phone, and the resident reported that he/she had gone to a local hospital to see his/her nephrologist.</p> <p>Resident #1's conservator and co-conservator were notified, and upon the resident's return, an alert bracelet was placed on the resident's ankle as a reminder to notify staff prior to leaving the facility. Education was provided to the resident and conservators regarding the importance of notifying staff for safety and medication management. All parties expressed understanding. (continued on next page)</p>		

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<p>F 0842</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Few</p>	<p>Safeguard resident-identifiable information and/or maintain medical records on each resident that are in accordance with accepted professional standards.</p> <p>Based on review of the clinical records, review of facility policy and staff interview for 1 of 4 residents (Resident # 130) reviewed for accidents, the facility failed to ensure the clinical record was complete and accurate. The findings include: Resident #130's diagnosis included dementia and cerebral infarction. The 06/13/2025 quarterly Minimum Data Set (MDS) Assessment indicate Resident #130 had moderate cognitive decline. The care plan dated 06/18/2025 indicated Resident #130 had altered cardiovascular status with interventions to assess for chest pain, shortness of breath and cyanosis (when oxygen moves away from the lips and nail beds, leaving the skin/nailbeds a blue/gray color versus pink), and to report changes to the physician. A progress note labeled Change in Condition dated 08/15/2025 at 05:06 PM indicated resident #130 was sitting in a wheelchair and noted to be pale and unresponsive the physician was notified and an order obtained to be sent to the emergency room. 911 was called while awaiting their arrival Resident #130 had become alert and was at his/her baseline and was sent to the emergency room. An assessment labeled Change in Condition/Concurrent Review completed by LPN # 3 indicated Resident #130's blood pressure was 142/65, pulse 52 and regular rhythm, and temperature was 97.0 degrees F. and the physician and responsible party were notified of Resident #130's transfer to the hospital for evaluation. An interview and clinical record review with the current Director of Nursing Services (DNS) on 03/11/2026 at 01:21 PM indicated there was no documented Registered Nurse (RN) assessment of Resident #130 prior to the transfer to the emergency room and s/he would have expected an RN assessment of Resident #130 written in the progress notes if the LPN documented the assessment the RN would then sign the assessment the LPN had documented. The DNS further indicated not finding Documentation of the exact time of the incident, when The Emergency Medical Services (EMS) arrived and left the facility with the resident, the state of alertness at the time of the transfer, sending a copy of the resident face sheet including all diagnosis medications and other pertinent data with the resident to the hospital and calling the hospital emergency room with expectation of the resident arriving to them along with the reason for the transfer and resident condition. The DNS further indicated not finding any documentation in Resident #130's clinical record. On 03/12/2026 at 2:45 PM an interview with LPN #4 working the 3-11 shift on 08/15/2025 indicated s/he arrived late to his/her shift and while counting medication at the medication cart with LPN #3 a nurse aide rolled Resident #130 in her w/c to the nurses as s/he was not responding. LPN#4 indicated having immediately checked the blood sugar level of Resident #130, unable to remember the exact reading but within normal range (70-110) and having minimally responded to a sternal rub (a painful stimulus technique used by professionals to check for responsiveness in unconscious patients, involving firm, grinding knuckle pressure on the breastbone LPN #4 and the nurse aide brought Resident #130 to her room and transferred him/her to the bed and Resident #130 had started to come around further indicating the 3-11 nursing supervisor, RN #5 was at the nurses station making the calls and preparing the transfer paperwork and had not been in the room. No documentation of the medical evaluation and care provided by LPN #4 as described within his/her interview we found within the clinical record. An interview on 03/12/2026 at 02:40 PM with RN #5 indicated she had seen Resident #130 in the wheelchair and started to come to but was unable to remember any details. The facility policy labeled Charting and Documentation indicated documentation should include medications administered, treatments or services performed, changes in the resident's condition, events, incidents or accidents involving the resident.</p>		