

Department of Health & Human Services
Centers for Medicare & Medicaid Services

Printed: 06/25/2026
Form Approved OMB
No. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 345088	(X2) MULTIPLE CONSTRUCTION A. Building B. Wing	(X3) DATE SURVEY COMPLETED 04/02/2026
NAME OF PROVIDER OR SUPPLIER Trinity Glen		STREET ADDRESS, CITY, STATE, ZIP CODE 849 Waterworks Road Winston-Salem, NC 27101	
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)		
F 0689 Level of Harm - Minimal harm or potential for actual harm Residents Affected - Few	Ensure that a nursing home area is free from accident hazards and provides adequate supervision to prevent accidents. **NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY** Based on record review and resident and staff interviews, the facility failed to provide care in a safe manner when Resident #4 rolled off the left side of her bed during incontinence care and required transfer to the hospital for medical evaluation. This was for 1 of 4 residents reviewed for accidents (Resident #4). Findings included: Resident #4 was admitted to the facility on [DATE]. The resident's diagnoses included vascular dementia, left side paralysis following stroke, and muscle weakness. Resident #4's care plan last updated on [DATE] revealed a focus area of risk for falls related to left side weakness and paralysis. Interventions included two-person physical assistance with bed mobility and using a lift for transfers. Review of the care guide (a guide that explains to staff members what and how much assistance each resident requires) last reviewed on [DATE] showed Resident #4 was a two staff assist rolling left and right, bed to chair transfers and used an extra-large sling with the lift. A quarterly Minimum Data Set (MDS) dated [DATE] indicated Resident #4 was cognitively intact and had range of motion impairment on one side of her upper and lower extremities. The MDS also indicated Resident #4 was dependent rolling left and right. A review of Resident #4's quarterly Interdisciplinary Team (IDT) risk assessment dated [DATE] revealed she was at moderate risk for falls. A review of Resident #4's physical therapy evaluation dated [DATE] showed Resident #4 was dependent rolling left to right and was unable to sit on edge of bed unsupported. During an interview with the Rehabilitation Manager on [DATE] at 7:45am, she stated Resident #4 was recently reevaluated on [DATE] for a decline in strength and functional mobility. The Rehabilitation Manager stated Resident #4 was unable to support herself and was dependent on staff for bed mobility due to her left side paralysis and weakness. During an interview with Resident #4 on [DATE] at 10:34 am she stated on [DATE], she rolled off the left side of her bed onto the floor while Nurse Aide (NA) #1 was providing care. Resident #4 stated she usually had two nurse aides turning her in bed, but sometimes only one would come in. Resident #4 indicated that when there was only one nurse aide, she usually turned on her left side and reached over to the side chair with her right hand and steadied herself that way. Resident #4 stated, on the day she fell, she thought the chair may have moved causing her to lose her balance. Resident #4 reported she didn't hit her head on anything but did hurt her right knee when she landed on the floor. Resident #4 indicated she went to the hospital because her knee hurt but nothing was broken. Resident #4 reported she had taken Tylenol for discomfort but had not needed anything else. During an interview with Nurse Aide #1 (NA #1) on [DATE] at 6:45am she stated on [DATE] during 3rd shift (7:00pm to 7:00am) at approximately 6:00am, she was providing incontinent care for Resident #4. NA #1 stated she usually worked on a different hall and had worked with Resident #4 a couple times prior. NA #1 stated she was aware that Resident #4 was dependent on staff for rolling left and right but she had cared for her by herself before by rolling Resident #4 on her left side and letting her hold onto the side chair with her right hand for support. NA #1 reported she could have asked the other nurse aide on the hall for assistance but thought the other nurse aide was already assisting another resident at that time. NA #1 stated she thought the side chair moved causing Resident #4 to lose her balance which allowed her to continue rolling left and off (continued on next page)		

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:	Facility ID: 345088
		If continuation sheet Page 1 of 4

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F 0756 Level of Harm - Minimal harm or potential for actual harm Residents Affected - Few	Ensure a licensed pharmacist perform a monthly drug regimen review, including the medical chart, following irregularity reporting guidelines in developed policies and procedures. **NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY** Based on record review and Nurse Practitioner, Pharmacist and Medical Director interviews, the facility failed to act on a pharmacy medication regimen recommendation for a Resident taking aripiprazole; aripiprazole is an atypical antipsychotic medication that can be used to treat depression and for which possible side effects include tardive dyskinesia (uncontrollable abnormal movements of the face, tongue or other body parts and requires frequent assessment and monitoring). The deficient practice occurred in 1 of 5 residents reviewed for unnecessary medications (Resident #14). Findings included: Resident #14 was admitted to the facility on [DATE] with diagnoses including depression. Review of records revealed a 1/5/26 physician's order for aripiprazole 5 mg (milligrams) by mouth once a day for depression. Review of the Pharmacist's monthly Medication Regimen Review (MRR) dated 1/7/26 revealed an abnormal involuntary movement scale (AIMS) assessment which revealed no identified involuntary movements or medication issues and there were no new pharmacy recommendations. The Pharmacist's monthly MRR dated 2/2/26 included a recommendation to the provider to consider tapering Resident #14 off of aripiprazole and introducing a different medication to treat her depression if appropriate. The Medical Director's response on the recommendation was noted as in agreement with a handwritten note dated 2/24/26 that the Nurse Practitioner would follow up with regards to the medication taper or to initiate a psychiatric services consult for the recommended tapering of aripiprazole. An AIMS assessment dated [DATE] revealed a score of zero, which indicated no abnormal involuntary body movements. Review of Resident #14's medication administration records (MAR) for February 2026 and March 2026 revealed Resident #14 had continued to receive 5mg of aripiprazole daily per the 1/6/26 order with no taper attempts noted. Review of the April 2026 MAR revealed the aripiprazole was documented as administered on 4/1/26 and 4/2/26. Further review of Resident #14's medical record revealed no documentation by the provider regarding follow up of the pharmacist's recommendation on 2/2/26, no dose taper attempts or initiation of a psychiatry services referral. On 4/2/26 at 11:00 AM, an interview with the Primary Nurse Practitioner was conducted. The Primary Nurse Practitioner stated she was not the one responsible for performing follow-up on medication taper recommendations for antipsychotic medications as she only did Primary Care services. The Primary Nurse Practitioner indicated the Psychiatry Nurse Practitioner was the person to speak to for all psychiatric related questions. When asked if she was aware of the 2/2/26 pharmacy recommendation and the Medical Director's note of acknowledgement, the Primary Nurse Practitioner repeated that she was not the person to speak to about anything related to psychiatry or the medication taper. A telephone interview with the Psychiatric Nurse Practitioner was conducted on 4/2/26 11:20 AM. The Psychiatric Nurse Practitioner indicated she did not know Resident #14. The Psychiatric Nurse Practitioner stated she had never received a referral for Resident #14 and Resident #14 was not on her list of new residents to see or evaluate so she could not answer any questions. On 4/2/26 11:54 AM, a telephone interview with the Medical Director was conducted. The Medical Director stated she remembered the pharmacist's recommendation dated 2/2/26 but did not realize the follow up had not been completed. The Medical Director indicated she had delegated this task to the NP and did not know why the follow-up had not been completed but that it should have occurred sooner rather than later. The Medical Director said she had not been informed of any new issues with Resident #14's condition or her medications and wasn't sure where the breakdown had occurred. The Medical Director further stated that either herself or the Primary Nurse Practitioner could perform the recommended follow-up and then decide if a taper versus a psychiatric consult was appropriate. The Medical Director said she would ensure this would be done. An interview with the Director of Nursing (DON) was conducted on 4/2/26 at 12:00 PM. The DON stated anytime a pharmacist made a recommendation for the provider, the recommendations were emailed to (continued on next page)		

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F 0756 Level of Harm - Minimal harm or potential for actual harm Residents Affected - Few	the Assistant Director of Nursing (ADON) who then printed off the recommendations and gave them to the provider for review. The DON said there was no box or bin where these forms were placed for the providers to pick up but rather the forms were actually handed to the providers in person for their review, then the provider would carry out the recommendations as they deemed appropriate. On 4/2/26 at 12:10 PM, a follow up interview with the Primary Nurse Practitioner was conducted and included a review of Resident #14's medical record and the pharmacy recommendation dated 2/2/26 the Medical Director had documented on. The Primary Nurse Practitioner stated she had never seen the 2/2/26 pharmacy recommendation and did not know about it. The Primary Nurse Practitioner indicated she did not know there was a directive for her to follow up on the pharmacy recommendation for Resident #14 to ascertain if a medication taper versus a psychiatric referral was appropriate. The Primary Nurse Practitioner said that all referrals and/or pharmacy recommendations were given to either the Medical Director or herself, but she did not know about this particular recommendation. An interview was conducted with the ADON on 4/2/26 at 12:41 PM. The ADON said when the pharmacist made a medication regimen recommendation, the pharmacist emailed all their recommendations to her in a batch email. The ADON said she then reviewed every recommendation and printed them and hand delivered them to the providers, to either the Primary Nurse Practitioner or to the Medical Director, whoever was on site on the given day. The ADON said the provider then reviewed the recommendations and signed off on them and then returned them to her for scanning into the electronic medical record. The ADON said she monitored the recommendations to ensure the recommendations were actually acted upon. The ADON said she was aware of the recommendation dated 2/2/26 for Resident #14, had noticed no follow up had been conducted and re-delivered the recommendation in person to the Primary Nurse Practitioner for review and follow up. The ADON said she could not remember when she re-submitted the recommendation to the Primary Nurse Practitioner, but it was after she noticed that no apparent action had been taken. On 4/2/26 at 1:20 PM, a telephone interview with the Pharmacist was conducted. The Pharmacist stated that anytime she made a medication regimen recommendation, she would follow up on her recommendations to see if they were carried out. The Pharmacist indicated she was able to see all of her outstanding recommendations in their computer system as soon as she logged in. The Pharmacist noted follow ups were usually conducted by the providers within 30 to 45 days from the date of the recommendation but she sometimes allowed for a maximum of 60 days due to the schedules of the providers and not knowing the exact days the providers were onsite. The Pharmacist said if a recommendation was not acted upon within that timeframe, she would initiate a new recommendation to the provider. The Pharmacist further stated she would be coming to the facility next week for her next monthly visit and would follow up on the 2/2/26 recommendation for Resident #14 at that time. In an interview with the Administrator on 4/2/26 at 2:20 PM, the Administrator stated she did not know exactly where the breakdown had occurred with the pharmacy recommendation not yet being acted upon and her expectation was if a pharmacist made a medication regimen recommendation that the recommendation would be acted upon as soon as possible when the provider became aware of it to ensure the residents' medication needs were being addressed and met.		