

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

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STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 165446	(X2) MULTIPLE CONSTRUCTION A. Building B. Wing	(X3) DATE SURVEY COMPLETED 03/12/2026
NAME OF PROVIDER OR SUPPLIER Westbrook Acres		STREET ADDRESS, CITY, STATE, ZIP CODE 605 Garfield Street Gladbrook, IA 50635	
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 0552 Level of Harm - Minimal harm or potential for actual harm Residents Affected - Few	Ensure that residents are fully informed and understand their health status, care and treatments. Based on staff interview, clinical record review and facility policy review, the facility failed to obtain consent to administer psychotropic medications taken by 1 of 5 residents reviewed for unnecessary medications (Resident #9). The facility reported a census of 40 residents. Findings include: Review of the Minimum Data Set (MDS) assessment for Resident #9 dated 12/23/25 revealed a Brief Interview for Mental Status (BIMS) score of 10 out of 15, which indicated moderate cognitive impairment. The list of diagnoses included Alzheimer's Disease, Cerebrovascular Accident (CVA or stroke), seizure disorder, anxiety disorder, and psychotic disorder. Resident #9 used the following high-risk medications: antipsychotic, antianxiety, antidepressant, and anticonvulsant. Review of the Care Plan initiated 3/12/25 revealed a Focus area for psychotropic medication use related to diagnoses of anxiety, dementia, frontal lobe meningioma (brain tumor), and psychotic disorder. The Care Plan indicated that Resident #9 was currently took Buspirone (medication used to treat anxiety disorders). Interventions included, in part: a. Administer meds as per physician orders. Date initiated 3/12/25.b. Abnormal Involuntary Movement Scale (AIMS) test every 6 months. Date initiated: 9/25/25. c. Consult with Pharmacy for medication review as needed. Date initiated 3/12/25. d. Gradual Dose Reduction (GDR) as per physician when deemed appropriate. Date initiated 3/12/25. e. Monitor exacerbation of mood or behavior issues or acute changes and notify physician as needed. Date initiated 3/12/25. f. Monitor for potential significant side effects antianxiety and antipsychotic medications and notify physician of abnormal as needed. Date initiated: 3/12/25. Review of Resident #9's Medication Administration Records (MAR) between the dates of March 2025 and March 2026 revealed the following psychotropic medications and dose changes for the resident's medications: 1. Buspirone (Buspar) tablet 5 milligrams (mg), with instructions to give 1 tablet by mouth three times a day for Generalized Anxiety Disorder. Date initiated: 3/03/25a. Increase in Buspirone to 10 mg, with instructions to give 1 tablet three times a day. Start date: 5/09/25.b. Decrease in Buspirone to 7.5 mg, with instructions to give half a 15 mg tablet (7.5 mg) three times a day. Start date: 12/02/25. 2. Quetiapine (Seroquel) tablet 25 mg, with instructions to give one tablet by mouth at bedtime for dementia. Date initiated: 3/03/25a. Decrease in Quetiapine to 12.5 mg at bedtime, between the dates of 4/08/25 and 4/21/25, then discontinue (stop) medication on 4/22/25. b. Restart Quetiapine 12.5 mg, with instructions to give 12.5 mg daily. Start date: 9/26/25. 3. Citalopram (Celexa) tablet 10 mg, with instructions to give one tablet by mouth daily for anxiety/depression. Date initiated: 7/31/25. a. Increase Citalopram to 20 mg, with instructions to give 20 mg by mouth daily. Start date: 9/16/25. Review of the facility provided document titled Psychotropic Medication Therapy Informed Consent Form revealed the following:1. Resident #9 signed consent on 3/03/25 to receive the antianxiety medication Buspirone 5 mg three times a day for anxiety/irritability. The medical records lacked documentation of consent obtained when dosage was increased to 10 mg three times a day on 5/09/25. 2. Resident #9 signed consent on 3/03/25 to receive the antipsychotic medication Quetiapine 25 mg daily for dementia with behavioral disturbance. The medical records lacked documentation of consent obtained following restarted Quetiapine 12.5 mg daily on 9/26/25.3. Resident #9's medical records lacked documentation of consent obtained for the antidepressant medication Citalopram 10 mg daily, ordered for depression, or consent obtained when Citalopram (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: 165446 If continuation sheet Page 1 of 6

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F 0552 Level of Harm - Minimal harm or potential for actual harm Residents Affected - Few	increased to 20 mg daily. On 3/12/25 at 2:22 PM, the Director of Nursing (DON) stated the normal process for psychotropic medication consents included obtaining consent from a resident or resident representative upon admission, after discussing potential side effects. The DON explained also was done intermittently when there were changes in medication dosage or if side effects were observed. The DON denied having documentation of psychotropic medication consents obtained for Resident #9's Citalopram started, increased Buspirone dosage, or restarted Quetiapine. Review of the facility policy titled Residents' [NAME] of Rights revised 11/2016 revealed the following per Section C. Planning and Implementing Care: The Resident has the right to be informed of, and participate in, his or her treatment, including: (1) The right to be fully informed in language that he or she can understand of his or her total health status, including but not limited to, his or her medical condition. (5) The right to be informed in advance, by the physician or other practitioner, or professional, of the risks and benefits of proposed care, of treatment and treatment alternatives or treatment options and to choose the alternative or option he or she prefers.		

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F 0605 Level of Harm - Minimal harm or potential for actual harm Residents Affected - Few	Prevent the use of unnecessary psychotropic medications or use medications that may restrain a resident's ability to function. **NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY** Based on staff interview, clinical record review, and facility policy review, the facility failed to ensure an as needed psychotropic medication was limited to 14 days unless the prescribing practitioner documented a rationale to extend medication for 1 of 5 residents reviewed for unnecessary medications (Resident #34). The facility reported a census of 40 residents. Findings include: Review of the Minimum Data Set (MDS) assessment dated [DATE] revealed the resident had a Brief Interview for Mental Status (BIMS) score of 12 out of 15, which indicated moderate cognitive impairment. Review of the resident's diagnoses per the MDS included cancer, non-Alzheimer's dementia, Cerebrovascular Accident (CVA or stroke), anxiety disorder, and adult failure to thrive. Review of the Care Plan Focus area initiated on 12/30/25 for Resident #34's use of the psychotropic medication revealed the following: The resident used Lorazepam, related to anxiety disorder, with times of yelling out to peers/staff, insomnia, non-sensical speech, and fixation on health. The Care Plan identified the resident currently took Lorazepam 0.5 milligrams (mg) twice per day as needed. Interventions included the following, in part: a. Administer anti-anxiety medications as ordered by physician. Monitor for side effects and effectiveness after given. Date initiated: 1/15/26. b. Abnormal Involuntary Movement Scale (AIMS) test every 6 months. Date initiated: 1/23/26. c. Consult with Hospice as needed. Date initiated: 2/03/26. d. Educate the resident/family/caregivers about risks, benefits, and the side effects and/or toxic symptoms of Lorazepam. Date initiated: 1/15/26. e. Monitor the resident for significant adverse drug reactions with anti-anxiety medication. Date initiated: 1/15/26. Review of a Nurse's Note dated 12/30/25 at 1:40 PM revealed Resident #34 had agitation and aggression noted and new order received for 0.5 mg of Lorazepam twice a day as needed was processed. The Nurse's Note dated 2/12/26 at 11:13 AM revealed the Hospice nurse visited and the facility received a new order to increase Resident #34's Lorazepam to 0.5 mg every 4 hours as needed for increased unusual behaviors. Review of Resident #34's Medication Administration Record (MAR) dated March 2026 revealed a current order for Lorazepam (Ativan) 0.5 mg, with instructions to give 1 tablet every 4 hours as needed (order dated 2/16/26). Lorazepam 0.5 mg was given multiple times in the month for agitation, anxiety, and restlessness. Review of Resident #34's Electronic Health Record (EHR) lacked documentation of physician rationale to extend the order for Lorazepam 0.5 mg as needed for greater than 14 days. During an interview on 3/12/26 at 1:18 PM, Staff E, Licensed Practical Nurse (LPN) reported Resident #34 started the Lorazepam 0.5 mg on 12/30/25 and was increased by Hospice. Staff E stated the resident's as needed Lorazepam was minimally effective in treating Resident #34's symptoms of anxiety, restlessness, and agitation unless given early or as soon as symptoms started. On 3/12/26 at 3:30 PM, the Facility Administrator denied documentation related to physician indication to continue Resident #34's Lorazepam 0.5 mg as needed past 14 days. The facility policy titled Drug Regimen Review Policy, undated, revealed the objective of the drug regimen review policy was to try to minimize or prevent adverse consequences or to prevent residents from receiving unnecessary drugs.		

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F 0627 Level of Harm - Minimal harm or potential for actual harm Residents Affected - Few	Ensure the transfer/discharge meets the resident's needs/preferences and that the resident is prepared for a safe transfer/discharge. **NOTE- TERMS IN BRACKETS HAVE BEEN EDITED TO PROTECT CONFIDENTIALITY** Based on clinical record review, staff interview and policy review the facility failed to complete a comprehensive assessment and evaluation of a resident for readmission to the facility after hospitalization for 1 of 3 residents reviewed for discharge (Resident #48). The facility reported a census of 40 residents. Findings include: The Minimum Data Set (MDS) assessment dated [DATE] for Resident #48 revealed the resident admitted to the facility on [DATE]. The MDS assessment dated [DATE] for Resident #48 documented an unplanned discharge to a short term general hospital, with return not anticipated. Diagnoses included urinary tract infection over last 30 days, depression and vascular dementia. A Baseline Care plan initiated 1/9/26 documented Resident #48 admitted from the hospital with the goal to return home after therapy goals were met. Resident #48 was alert and oriented to self, and confused. The discharge goal revealed to remain at Long Term Care (LTC) setting. Resident #48 required assist of one with transfer, bathing, grooming, and locomotion on and off the unit. The resident used a walker. A Physical Therapy Evaluation and Plan of Treatment for Resident #48 beginning 1/9/26 documented the resident was referred by neurologist for skilled therapy after hospitalization for acute cystitis (bladder infection) and weakness. The resident was expected to transition to Long Term Care (LTC) after skilled therapy. Complexities included acute cystitis, legal blindness, heart disease, stroke and vascular dementia. The Transition/Discharge section of the form revealed the resident was to reside in the LTC facility. The admission Summary Note dated 1/9/26 at 12:20 PM revealed the resident admitted to the facility, was alert, and was pleasantly confused. The Nurse's Note dated 1/10/26 at 12:41 AM revealed the Resident #48 was able to make their needs known and denied pain. The Nurse's Note dated 1/10/26 at 12:55 PM revealed the resident required reminders to use call light and to not transfer independently. The Nurse's Note dated 1/10/26 at 3:58 PM revealed the resident's family visited, and the resident rested in a chair. The Nurse's Note dated 1/11/26 at 2:33 AM revealed Resident #48 rested quietly in bed, voiced no complaints, and was able to make their needs known. The Nurse's Note dated 1/11/26 at 9:50 AM revealed the resident required assist of one with cares and required cues and reminders from staff. The Nurse's Note dated 1/12/26 at 4:45 AM documented Resident #48 refused to get up to bathroom, and stated, I dont get up at night. Per the Nurse's Note, incontinence care was provided, and the resident was pleasant and cooperative. The Nurse's Note dated 1/12/26 at 8:00 AM revealed the resident was slightly non-compliant with morning cares and dressing, with assist of two (staff) used. The Nurse's Note further revealed the resident yelled in the main dining room that she did not belong [at facility] and insisted on going home right now. The resident mimicked staff, yelled at staff, and tried to kick and swing at staff. The resident attempted to push their chair into other residents, refused to eat breakfast, and was noncompliant with therapies. The Nurse's Note further revealed the resident was placed in her room and was offered to sit in her recliner. The resident refused, and yelled to get her out of here right now. The resident was left alone to calm, and was checked on frequently from the doorway. The Nurse's Note later in the day at 10:00 AM revealed the resident said no thank you to the offer of her recliner. The Nurse's Note dated 1/12/26 at 1:27 PM revealed the resident had a new order to obtain a urinalysis. Per the Nurse's Note dated 1/13/26 at 1:30 AM, the resident's urine specimen was obtained. The Nurse's Note dated 1/13/26 at 7:45 AM revealed the resident refused medication and spit them out two attempts. The resident demanded to go back to their room. The Nurse's Note dated 1/13/26 at 8:10 AM revealed the resident was observed in their room in their wheelchair with the wheelchair tipped backwards. The handle bars of the wheelchair prevented the resident's head from being on the floor. The Nurse's Note dated 1/13/26 at 8:45 AM revealed the resident wanted to leave, staff tried to talk with her, and the nurse instructed to go ahead and have 911 paged. The Nurse's Note dated 1/13/26 at 9:15 AM revealed notification to the (continued on next page)		

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