

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 395592	(X2) MULTIPLE CONSTRUCTION A. Building B. Wing	(X3) DATE SURVEY COMPLETED 09/04/2025
NAME OF PROVIDER OR SUPPLIER Haida Nursing and Rehab		STREET ADDRESS, CITY, STATE, ZIP CODE 397 Third Avenue Extension Hastings, PA 16646	

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 0552</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Few</p>	<p>Ensure that residents are fully informed and understand their health status, care and treatments.</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

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 395592	(X2) MULTIPLE CONSTRUCTION A. Building B. Wing	(X3) DATE SURVEY COMPLETED 09/04/2025
NAME OF PROVIDER OR SUPPLIER Haida Nursing and Rehab		STREET ADDRESS, CITY, STATE, ZIP CODE 397 Third Avenue Extension Hastings, PA 16646	
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 0552</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Few</p>	<p>Based on review of facility policies and clinical records, as well as staff interviews, it was determined that the facility failed to inform the resident representative in advance of the risks and benefits of a psychotropic medication (medications that affect the persons mental state, emotions and behavior) use and the treatment alternatives prior to initiating the administration of the medication for two of 5 residents reviewed (Resident 1 and 5). Findings Include: The facility's policy related to the use of psychotropic medications, dated April 23, 2025, indicated that prior to initiating or increasing a psychotropic medication, the resident, family, and/or resident representative must be informed of the benefits, risks, and alternatives for the medications, including any black box warnings for antipsychotic medications, in advance of such initiation or increase. The facility will document that the resident or resident representative was informed in advance of the risks and benefits of the proposed care, the treatment alternatives or other options and the preferred option to accept or decline in a format the facility deems to use (e.g., written consent form, narrative note, etc.). A quarterly Minimum Data Set (MDS) assessment (a mandated assessment of a resident's abilities and care needs) for Resident 1, dated May 30, 2025, revealed that the resident was severely cognitively impaired, received antipsychotic and antianxiety medications, and had diagnoses that included dementia and anxiety. A skilled nursing facility psychiatric visit dated March 25, 2025, indicated that Resident 1 was seen for ongoing agitation, was up yelling and was difficult to redirect at night. Start five milligrams (mg) of Buspar (medication to treat anxiety) three times a day. Patient consents for the use of current medications to be obtained with social services. Physician's orders for Resident 1, dated March 27 2025, revealed that the resident was to receive five mg of buspirone three times a day for dementia with agitation. There was no documented evidence in the resident's clinical record to indicate that the resident's representative was informed in advance of the risks and benefits and treatment alternatives prior to initiating the dose of Buspar. Interview with the Director of Nursing on September 4, 2025, at 2:54 p.m., confirmed that there was no documented evidence in the Resident' 1s clinical record that the resident's representative was informed in advance of the risks and benefits and treatment alternatives prior to initiating Buspar medication. An admission Minimum Data Set (MDS) assessment (a mandated assessment of a resident's abilities and care needs) for Resident 5, dated May 13, 2025, revealed that she was cognitively impaired, required assistance with care needs, received antipsychotic, antianxiety, and antidepressant medications (psychotropic medications) and had diagnoses that included dementia, depression, and anxiety. A physician's note for Resident 5, dated May 14, 2025, at 8:48 p.m. revealed that the resident experienced a sudden behavioral escalation. Psychiatry was consulted and increased her Seroquel (antipsychotic medication used to treat mental health disorders) dose to 250 milligrams (mg). Physician's orders for Resident 5, dated May 15, 2025, included an order for the resident to receive 250 mg of Seroquel in the evening related to dementia with agitation. There was no documented evidence in the resident's clinical record that the resident's representative was informed in advance of the risks and benefits and treatment alternatives prior to initiating the increased dose of Seroquel. A nursing note for Resident 5, dated May 27, 2025, at 2:11 p.m. revealed that Certified Physician's Assistant (CRNP-registered nurse with advanced education and training who can diagnose, treat and prescribe medication) with psychiatric services was called and updated with the resident's increased agitation and a medication change was ordered to add 50 mg of Seroquel twice daily in addition to the 250 mg of Seroquel daily in the evening. Physician's orders for Resident 5, dated May 27, 2025, included an order for the resident to receive two 25 mg tablets of Seroquel to equal 50 mg twice daily related to dementia with agitation. There was no documented evidence in the resident's clinical record that the resident's representative was informed in advance of the risks and benefits and treatment alternatives prior to initiating the increased dose of Seroquel. Interview with the Director of Nursing on September 4, 2025, at 3:25 p.m., confirmed that a nursing note, dated May 15, 2025, at 1:30 p.m. and a nursing note, dated May 27, 2025, at 2:13 p.m. indicated that Resident 5's representative was notified of the dosage changes to Seroquel; however, there was no documented evidence in the resident's clinical record that the resident's representative was informed in advance of the risks and benefits and treatment alternatives prior to initiating the increased doses of Seroquel.</p>		

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 395592	(X2) MULTIPLE CONSTRUCTION A. Building B. Wing	(X3) DATE SURVEY COMPLETED 09/04/2025
NAME OF PROVIDER OR SUPPLIER Haida Nursing and Rehab		STREET ADDRESS, CITY, STATE, ZIP CODE 397 Third Avenue Extension Hastings, PA 16646	

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 0605</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Some</p>	<p>Prevent the use of unnecessary psychotropic medications or use medications that may restrain a resident's ability to function.</p> <p>(continued on next page)</p>

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 395592	(X2) MULTIPLE CONSTRUCTION A. Building B. Wing	(X3) DATE SURVEY COMPLETED 09/04/2025
NAME OF PROVIDER OR SUPPLIER Haida Nursing and Rehab		STREET ADDRESS, CITY, STATE, ZIP CODE 397 Third Avenue Extension Hastings, PA 16646	
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 0605</p> <p>Level of Harm - Minimal harm or potential for actual harm</p> <p>Residents Affected - Some</p>	<p>Based on a review of facility policies and clinical records as well as staff interviews, it was determined that the facility failed to ensure that residents medication regime was free from unnecessary psychotropic medication (drugs that affect a person's mental state, emotions, and behavior) for two of 5 residents reviewed (Residents 1 and 5). Findings Include: The facility policy related to the use of psychotropic medications, dated April 23, 2025, indicated that residents will only receive psychotropic medications when other nonpharmacological interventions are clinically contraindicated. Additionally, these medications should only be used to treat the resident's medical symptoms and not used for discipline or staff convenience, which would deem it a chemical restraint. Nonpharmacological approaches must be attempted, unless clinically contraindicated, to minimize the need for psychotropic medications, use the lowest possible dose, or discontinue the medications. A quarterly Minimum Data Set (MDS) assessment (a mandated assessment of a resident's abilities and care needs) for Resident 1, dated May 30, 2025, revealed that the resident was severely cognitively impaired, was sometime understood, could sometimes understand, received antipsychotic and antianxiety medications, and had diagnoses that included dementia and anxiety. A skilled nursing facility psychiatric visit, dated March 25, 2025, indicated that Resident 1 was seen for ongoing agitation, was up yelling, and was difficult to redirect at night. Nonpharmacological interventions were to be used for this resident. Physician's orders for Resident 1, dated May 19, 2025 and June 11, 2025, revealed that the resident was to receive 0.25 milligrams (mg) of Alprazolam (a psychotropic medication) every eight hours as needed for anxiety. Review of Resident 1's Medication Administration Record (MAR) for May and June 2025, revealed that the resident received 0.25 mg of Alprazolam on May 19 at 2:41 p.m., on May 20 a. m at 10:17 a.m., on May 21 at 2:50 a.m., on June 11 at 1:50 p.m, on June 17 at 12:20 a.m., and on June 19 at 11:30 p.m. A review of Resident 1's clinical record revealed no documented evidence that non-pharmacological interventions were attempted prior to administering the as needed Alprazolam on the above-mentioned dated and times. Interview with the Director of Nursing on September 4, 2025, at 2:54 p.m. confirmed that there was no documented evidence that non-pharmacological interventions were attempted prior to administering as needed Alprazolam to Resident 1 on the above-mentioned dates/times. An admission MDS assessment for Resident 5, dated May 13, 2025, revealed that she was cognitively impaired, required assistance with care needs, received antipsychotic, antianxiety, and antidepressant medications (psychotropic medications) and had diagnoses that included dementia, depression, and anxiety. Physician's orders for Resident 5, dated May 9, 2025, revealed that the resident was to receive 0.5 milligrams (mg) of Lorazepam (a psychotropic medication) every eight hours, may use three times daily, as needed for anxiety. Physician's orders for Resident 5, dated May 24, 2025, included an order for the resident to receive 0.5 mg of Lorazepam three times daily as needed for anxiety. Physician's orders for Resident 5, dated May 27, 2025, June 8, 2025, and June 24, 2025, included orders for the resident to receive 0.5 mg of Lorazepam every eight hours as needed for anxiety. Physician's orders for Resident 5, dated July 10, 2025, July 24, 2025, July 30, 2025, revealed that the resident was to receive 0.5 mg of Lorazepam every eight hours, may use three times daily, as needed for anxiety. Physician's orders for Resident 5, dated August 8, 2025, and August 25, 2025, revealed that the resident was to receive 0.5 mg of Lorazepam every eight hours as needed for anxiety. Review of Resident 5's Medication Administration Record (MAR) for May, 2025 through August, 2025, revealed that the resident received 0.5 mg of Lorazepam on May 10 at 1:20 p.m.; May 14 at 11:20 p.m.; May 18 at 10:01 a.m.; May 23 at 5:58 p.m.; May 27 at 7:15 p.m.; May 28 at 8:15 a.m.; May 31 at 3:38 p.m.; June 2 at 8:33 a.m.; June 3 at 8:11 a.m.; June 6 at 3:56 p.m.; June 10 at 3:43 p.m.; June 14 at 4:10 p.m.; July 10 at 5:48 p.m.; July 29 at 12:05 p.m.; August 14 at 2:15 p.m.; August 21 at 12:14 p.m.; and August 25 at 2:22 p.m. Review of Resident 5's clinical record, revealed no documented evidence that non-pharmacological interventions were attempted prior to administering the as needed Lorazepam on the above-mentioned dated and times. Interview with the Director of Nursing on September 4, 2025, at 3:25 p.m. confirmed that there was no documented evidence that non-pharmacological interventions were attempted prior to administering as needed Lorazepam to Resident 5 on the above-mentioned dates/times. 28 Pa. Code 211.2(d)(3) Medical director 28 Pa. Code: 211.9(a)(1) Pharmacy services. 28 Pa. Code: 211.12 (d)(5) Nursing services.</p>		