NEW MEXICO
Department of Health
Division of Health Improvement

MICHELLE LUJAN GRISHAM
Governor

DAVID R. SCRASE, M.D.
Acting Cabinet Secretary

Date:	May 11, 2022
To:	Christine Chapman, DSP Supervisor / SC / Executive Director / Owner
Provider: Address: State/Zip:	Safe Harbor, Inc. 825 Quesenberry St. Las Cruces, New Mexico 88005
E-mail Address:	garychpm@aol.com
Region: Routine Survey: Verification Survey:	Southwest October 25 – November 5, 2021 April 18 – 28, 2022
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	Supported Living and Customized Community Supports
Survey Type:	Verification
Team Leader:	Lei Lani Nava, MPH, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau
Team Members:	Verna Newman-Sikes, AA, Credentials, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Dear Ms. Christine Chapman,

The Division of Health Improvement/Quality Management Bureau has completed a Verification survey of the services identified above. The purpose of the survey was to determine compliance with your Plan of Correction submitted to DHI regarding the *Routine Survey on October 25 – November 5, 2021*.

Determination of Compliance:

The Division of Health Improvement, Quality Management Bureau has determined your agency is now in:

Partial Compliance with Standard Level Tags and Conditions of Participation Level Tags: This determination is based on noncompliance with one to five (1 – 5) Condition of Participation Level Tags *(refer to Attachment D for details)*. The attached QMB Report of Findings indicates Standard Level and Condition of Participation Level deficiencies identified and requires completion and implementation of a Plan of Correction.

The following tags are identified as Condition of Participation Level:

- Tag # 1A09 Medication Delivery Routine Medication Administration ***(New / Repeat Findings)***
- Tag # 1A09.1 Medication Delivery PRN Medication Administration ***(New / Repeat Findings)***
- Tag # 1A09.2 Medication Delivery Nurse Approval for PRN Medication ***(New / Repeat Findings)***

The following tags are identified as Standard Level:

- Tag # 1A26 Consolidated On-line Registry Employee Abuse Registry ***(New / Repeat Findings)***

However, due to the new/repeat deficiencies your agency will be required to contact your DDSD Regional Office for technical assistance and follow up and complete the Plan of Correction document attached at the end of this report. Please respond to the Plan of Correction Coordinator within 10 business days of receipt of this letter.

Plan of Correction:
The attached Report of Findings identifies the new/repeat Standard Level deficiencies found during your agency's verification compliance review. You are required to complete and implement a Plan of Correction. Your agency has a total of 10 business days from the receipt of this letter. The Plan of Correction must include the following:

1. Evidence your agency has contacted your DDSD Regional Office for technical assistance;
2. A Plan of Correction detailing Quality Assurance/Quality Improvement processes to prevent your agency from receiving deficiencies in the future. Please use the format provided at the end of this report;
3. Documentation verifying that newly cited deficiencies have been corrected.

Submission of your Plan of Correction:
Please submit your agency's Plan of Correction and documentation verifying correction of survey deficiencies within 10 business days of receipt of this letter to the parties below:

1. **Quality Management Bureau, Attention: Plan of Correction Coordinator**
   5301 Central Ave. NE Suite 400, New Mexico 87108
   MonicaE.Valdez@state.nm.us

2. **Developmental Disabilities Supports Division Regional Office for region of service surveyed**

Failure to submit your POC within the allotted 10 business days may result in the imposition of a $200 per day Civil Monetary Penalty until it is received, completed and/or implemented.

Please call the Plan of Correction Coordinator Monica Valdez at 505-273-1930 if you have questions about the Report of Findings or Plan of Correction. Thank you for your cooperation and for the work you perform.

Sincerely,

Lei Lani Nava, MPH

Lei Lani Nava, MPH
Team Lead/Healthcare Surveyor
Division of Health Improvement
Quality Management Bureau

Survey Process Employed:

Administrative Review Start Date:	April 18, 2022
Contact:	**Safe Harbor, Inc.** Christine Chapman, DSP Supervisor / SC / Executive Director / Owner **DOH/DHI/QMB** Lei Lani Nava, MPH, Team Lead/Healthcare Surveyor
Exit Conference Date:	April 18, 2022
Present:	**Safe Harbor, Inc.** Christine Chapman, DSP Supervisor / SC / Executive Director / Owner Jannie Medley, Office Assistant Rebecca Ruiz, Office Manager **DOH/DHI/QMB** Lei Lani Nava, MPH, Team Lead/Healthcare Surveyor Amanda Castaneda-Holguin, MPA, Healthcare Surveyor Supervisor Verna Newman-Sikes, AA, Healthcare Surveyor **DDSD - SW Regional Office** David Chavez, Community Inclusion Specialist
Total Sample Size:	5 0 - *Jackson* Class Members 5 - Non-*Jackson* Class Members 5 - Supported Living 5 - Customized Community Supports
Persons Served Records Reviewed	5
Direct Support Personnel Records Reviewed	17 *(Note: 1 DSP perform dual role~~s~~ as Service Coordinator)*
Direct Support Personnel Interviewed during Routine Survey	3 *(Note: Interviews conducted by video / phone due to COVID-19 Public Health Emergency)*
Service Coordinator Records Reviewed	1 *(Note: 1 Service Coordinator perform dual role~~s~~ as DSP)*
Nurse Interview completed during Routine Survey	1

Administrative Processes and Records Reviewed:

- Medicaid Billing/Reimbursement Records for all Services Provided
- Accreditation Records
- Oversight of Individual Funds
- Individual Medical and Program Case Files, including, but not limited to:
    - °Individual Service Plans
    - °Progress on Identified Outcomes
    - °Healthcare Plans
    - °Medication Administration Records
    - °Medical Emergency Response Plans
    - °Therapy Evaluations and Plans
    - °Healthcare Documentation Regarding Appointments and Required Follow-Up

    - Other Required Health Information
- Internal Incident Management Reports and System Process / General Events Reports
- Personnel Files, including nursing and subcontracted staff
- Staff Training Records, Including Competency Interviews with Staff
- Agency Policy and Procedure Manual
- Caregiver Criminal History Screening Records
- Consolidated Online Registry/Employee Abuse Registry
- Human Rights Committee Notes and Meeting Minutes
- Evacuation Drills of Residences and Service Locations
- Quality Assurance / Improvement Plan

CC: Distribution List: DOH - Division of Health Improvement
DOH - Developmental Disabilities Supports Division
DOH - Office of Internal Audit
HSD - Medical Assistance Division
NM Attorney General's Office

Attachment B

Department of Health, Division of Health Improvement
QMB Determination of Compliance Process

The Division of Health Improvement, Quality Management Bureau (QMB) surveys compliance of the Developmental Disabilities Waiver (DDW) standards and other state and federal regulations. For the purpose of the LCA / CI survey the CMS waiver assurances have been grouped into four (4) Service Domains: Plan of Care (ISP Implementation); Qualified Providers; Health, Welfare and Safety; and Administrative Oversight (note that Administrative Oversight listed in this document is not the same as the CMS assurance of Administrative Authority. Used in this context it is related to the agency's operational policies and procedures, Quality Assurance system and Medicaid billing and reimbursement processes.)

The QMB Determination of Compliance process is based on provider compliance or non-compliance with standards and regulations identified during the on-site survey process and as reported in the QMB Report of Findings. All areas reviewed by QMB have been agreed to by DDSD and DHI/QMB and are reflective of CMS requirements. All deficiencies (non-compliance with standards and regulations) are identified and cited as either a Standard level deficiency or a Condition of Participation level deficiency in the QMB Reports of Findings. All deficiencies require corrective action when non-compliance is identified.

Each deficiency in your Report of Findings has been predetermined to be a Standard Level Deficiency, a Condition of Participation Level Deficiency, if below 85% compliance or a non-negotiable Condition of Participation Level Deficiency. Your Agency's overall Compliance Determination is based on a Scope and Severity Scale which takes into account the number of Standard and Condition Level Tags cited as well as the percentage of Individuals affected in the sample.

Conditions of Participation (CoPs)

CoPs are based on the Centers for Medicare and Medicaid Services, Home and Community-Based Waiver required assurances, in addition to the New Mexico Developmental Disability Waiver (DDW) Service Standards. The Division of Health Improvement (DHI), in conjunction with the Developmental Disability Support Division (DDSD), has identified certain deficiencies that have the potential to be a Condition of Participation Level, if the tag falls below 85% compliance based on the number of people affected. Additionally, there are what are called non-negotiable Conditions of Participation, regardless if one person or multiple people are affected. In this context, a CoP is defined as an essential / fundamental regulation or standard, which when out of compliance directly affects the health and welfare of the Individuals served. If no deficiencies within a Tag are at the level of a CoP, it is cited as a Standard Level Deficiency.

Service Domains and CoPs for <u>Living Care Arrangements and Community Inclusion</u> are as follows:

<u>Service Domain: Service Plan: ISP Implementation -</u> *Services are delivered in accordance with the service plan, including type, scope, amount, duration and frequency specified in the service plan.*

Potential Condition of Participation Level Tags, if compliance is below 85%:

- **1A08.3 –** Administrative Case File: Individual Service Plan / ISP Components
- **1A32 –** Administrative Case File: Individual Service Plan Implementation
- **LS14 –** Residential Service Delivery Site Case File (ISP and Healthcare Requirements)
- **IS14 –** CCS / CIES Service Delivery Site Case File (ISP and Healthcare Requirements)

<u>Service Domain: Qualified Providers -</u> *The State monitors non-licensed/non-certified providers to assure adherence to waiver requirements. The State implements its policies and procedures for verifying that provider training is conducted in accordance with State requirements and the approved waiver.*

Potential Condition of Participation Level Tags, if compliance is below 85%:

- **1A20 -** Direct Support Personnel Training
- **1A22 -** Agency Personnel Competency
- **1A37 –** Individual Specific Training

Non-Negotiable Condition of Participation Level Tags (one or more Individuals are cited):

- **1A25.1 –** Caregiver Criminal History Screening
- **1A26.1 –** Consolidated On-line Registry Employee Abuse Registry

Service Domain: Health, Welfare and Safety - *The State, on an ongoing basis, identifies, addresses and seeks to prevent occurrences of abuse, neglect and exploitation. Individuals shall be afforded their basic human rights. The provider supports individuals to access needed healthcare services in a timely manner.*

Potential Condition of Participation Level Tags, if compliance is below 85%:

- **1A08.2 –** Administrative Case File: Healthcare Requirements & Follow-up
- **1A09 –** Medication Delivery Routine Medication Administration
- **1A09.1 –** Medication Delivery PRN Medication Administration
- **1A15.2 –** Administrative Case File: Healthcare Documentation (Therap and Required Plans)

Non-Negotiable Condition of Participation Level Tags (one or more Individuals are cited):

- **1A05 –** General Requirements / Agency Policy and Procedure Requirements
- **1A07 –** Social Security Income (SSI) Payments
- **1A09.2 –** Medication Delivery Nurse Approval for PRN Medication
- **1A15 –** Healthcare Coordination - Nurse Availability / Knowledge
- **1A31 –** Client Rights/Human Rights
- **LS25.1 –** Residential Reqts. (Physical Environment - Supported Living / Family Living / Intensive Medical Living)

Attachment C

Guidelines for the Provider
Informal Reconsideration of Finding (IRF) Process

Introduction:
Throughout the QMB Survey process, surveyors are openly communicating with providers. Open communication means surveyors have clarified issues and/or requested missing information before completing the review through the use of the signed/dated "Document Request," or "Administrative Needs," etc. forms. Regardless, there may still be instances where the provider disagrees with a specific finding. Providers may use the following process to informally dispute a finding.

Instructions:
1. The Informal Reconsideration of the Finding (IRF) request must be received in writing to the QMB Bureau Chief **within 10 business days** of receipt of the final Report of Findings **(*Note: No extensions are granted for the IRF)*.**
2. The written request for an IRF *must* be completed on the QMB Request for Informal Reconsideration of Finding form available on the QMB website: https://nmhealth.org/about/dhi/cbp/irf/
3. The written request for an IRF must specify in detail the request for reconsideration and why the finding is inaccurate.
4. The IRF request must include all supporting documentation or evidence.
5. If you have questions about the IRF process, email the IRF Chairperson, Valerie V. Valdez at valerie.valdez@state.nm.us for assistance.

The following limitations apply to the IRF process:
- The written request for an IRF and all supporting evidence must be received within 10 business days.
- Findings based on evidence requested during the survey and not provided may not be subject to reconsideration.
- The supporting documentation must be new evidence not previously reviewed or requested by the survey team.
- Providers must continue to complete their Plan of Correction during the IRF process
- Providers may not request an IRF to challenge the sampling methodology.
- Providers may not request an IRF based on disagreement with the nature of the standard or regulation.
- Providers may not request an IRF to challenge the team composition.
- Providers may not request an IRF to challenge the DHI/QMB determination of compliance or the length of their DDSD provider contract.

A Provider forfeits the right to an IRF if the request is not received within 10 business days of receiving the report and/or does not include all supporting documentation or evidence to show compliance with the standards and regulations.

The IRF Committee will review the request; the Provider will be notified in writing of the ruling; no face-to-face meeting will be conducted.

When a Provider requests that a finding be reconsidered, it does not stop or delay the Plan of Correction process. **Providers must continue to complete the Plan of Correction, including the finding in dispute regardless of the IRF status.** If a finding is removed or modified, it will be noted and removed or modified from the Report of Findings. It should be noted that in some cases a Plan of Correction may be completed prior to the IRF process being completed. The provider will be notified in writing on the decisions of the IRF committee.

Attachment D

QMB Determinations of Compliance

Compliance:

The QMB determination of *Compliance* indicates that a provider has either no deficiencies found during a survey or that no deficiencies at the Condition of Participation Level were found. The agency has obtained a level of compliance such that there is a minimal potential for harm to individuals' health and safety. To qualify for a determination of *Compliance*, the provider must have received no Conditions of Participation Level Deficiencies and have a minimal number of Individuals on the sample affected by the findings indicated in the Standards Level Tags.

Partial-Compliance with Standard Level Tags:

The QMB determination of *Partial-Compliance with Standard Level Tags* indicates that a provider is in compliance with all Condition of Participation Level deficiencies but is out of compliance with a certain percentage of Standard Level deficiencies. This partial-compliance, if not corrected, may result in a negative outcome or the potential for more than minimal harm to individuals' health and safety. There are two ways to receive a determination of Partial Compliance with Standard Level Tags:

1. Your Report of Findings includes 16 or fewer Standards Level Tags with between 75% and 100% of the survey sample affected in any tag.

2. Your Report of Findings includes 17 or more Standard Level Tags with between 50% to 74% of the survey sample affected in any tag.

Partial-Compliance with Standard Level Tags and Condition of Participation Level Tags:

The QMB determination of *Partial-Compliance with Standard Level Tags and Condition of Participation Level Tags* indicates that a provider is out of compliance with one to five (1 – 5) Condition of Participation Level Tags. This partial-compliance, if not corrected, may result in a serious negative outcome or the potential for more than minimal harm to individuals' health and safety.

Non-Compliance:

The QMB determination of *Non-Compliance* indicates a provider is significantly out of compliance with both Standard Level deficiencies and Conditions of Participation level deficiencies. This non-compliance, if not corrected, may result in a serious negative outcome or the potential for more than minimal harm to individuals' health and safety. There are three ways an agency can receive a determination of Non-Compliance:

1. Your Report of Findings includes 17 or more total Tags with 0 to 5 Condition of Participation Level Tags with 75% to 100% of the survey sample affected in any Condition of Participation Level tag.

2. Your Report of Findings includes any amount of Standard Level Tags with 6 or more Condition of Participation Level Tags.

Compliance Determination	Weighting						
	LOW		MEDIUM			HIGH	
Total Tags:	up to 16	17 or more	up to 16	17 or more	Any Amount	17 or more	Any Amount
	and	and	and	and	And/or	and	And/or
COP Level Tags:	0 COP	0 COP	0 COP	0 COP	1 to 5 COP	0 to 5 CoPs	6 or more COP
	and	and	and	and		and	
Sample Affected:	0 to 74%	0 to 49%	75 to 100%	50 to 74%		75 to 100%	
"Non-Compliance"						**17 or more** Total Tags with **75 to 100%** of the Individuals in the sample cited in any CoP Level tag.	**Any Amount** of Standard Level Tags and **6 or more** Conditions of Participation Level Tags.
"Partial Compliance with Standard Level tags and Condition of Participation Level Tags"					**Any Amount** Standard Level Tags, plus **1 to 5** Conditions of Participation Level tags.		
"Partial Compliance with Standard Level tags"			**up to 16** Standard Level Tags with **75 to 100%** of the individuals in the sample cited in any tag.	**17 or more** Standard Level Tags with **50 to 74%** of the individuals in the sample cited any tag.			
"Compliance"	**Up to 16** Standard Level Tags with **0 to 74%** of the individuals in the sample cited in any tag.	**17 or more** Standard Level Tags with **0 to 49%** of the individuals in the sample cited in any tag.					

Agency: **Safe Harbor, Inc. – Southwest Region**
Program: Developmental Disabilities Waiver
Service: Supported Living and Customized Community Supports
Survey Type: Verification
Routine Survey: **October 25 – November 5, 2021**
Verification Survey: **April 18 – 28, 2022**

Standard of Care	Routine Survey Deficiencies October 25 – November 5, 2021	Verification Survey New and Repeat Deficiencies April 18 – 28, 2022
Service Domain: Qualified Providers – *The State monitors non-licensed/non-certified providers to assure adherence to waiver requirements. The State implements its policies and procedures for verifying that provider training is conducted in accordance with State requirements and the approved waiver.*		
Tag # 1A26 Consolidated On-line Registry Employee Abuse Registry	**Standard Level Deficiency**	**Standard Level Deficiency**
NMAC 7.1.12.8 - REGISTRY ESTABLISHED; PROVIDER INQUIRY REQUIRED: Upon the effective date of this rule, the department has established and maintains an accurate and complete electronic registry that contains the name, date of birth, address, social security number, and other appropriate identifying information of all persons who, while employed by a provider, have been determined by the department, as a result of an investigation of a complaint, to have engaged in a substantiated registry-referred incident of abuse, neglect or exploitation of a person receiving care or services from a provider. Additions and updates to the registry shall be posted no later than two (2) business days following receipt. Only department staff designated by the custodian may access, maintain and update the data in the registry. A. **Provider requirement to inquire of registry**. A provider, prior to employing or contracting with an employee, shall inquire of the registry whether the individual under consideration for employment or contracting is listed on the registry. B. **Prohibited employment.** A provider may not employ or contract with an individual to be an employee if the individual is listed on the registry as having a substantiated registry-referred incident of abuse, neglect or exploitation of a person receiving care or services from a provider. C. **Applicant's identifying information required**. In making the inquiry to the registry prior to employing or contracting with an employee, the provider shall use	Based on record review, the Agency did not maintain documentation in the employee's personnel records that evidenced inquiry into the Employee Abuse Registry prior to employment for 2 of 18 Agency Personnel. **The following Agency Personnel records contained evidence that indicated the Employee Abuse Registry check was completed after hire:** **Direct Support Personnel (DSP):** • #506 – Date of hire 9/15/2020, completed 9/16/2020. • #508 – Date of hire 7/28/2021, completed 7/29/2021.	**New / Repeat Findings:** Based on record review, the Agency did not maintain documentation in the employee's personnel records that evidenced inquiry into the Employee Abuse Registry prior to employment for 1 of 17 Agency Personnel. **The following Agency Personnel records contained evidence that indicated the Employee Abuse Registry check was completed after hire:** **Direct Support Personnel (DSP):** • #526 – Date of hire 11/29/2021, completed 12/08/2021.

identifying information concerning the individual under consideration for employment or contracting sufficient to reasonably and completely search the registry, including the name, address, date of birth, social security number, and other appropriate identifying information required by the registry. D. **Documentation of inquiry to registry**. The provider shall maintain documentation in the employee's personnel or employment records that evidences the fact that the provider made an inquiry to the registry concerning that employee prior to employment. Such documentation must include evidence, based on the response to such inquiry received from the custodian by the provider, that the employee was not listed on the registry as having a substantiated registry-referred incident of abuse, neglect or exploitation. E. **Documentation for other staff**. With respect to all employed or contracted individuals providing direct care who are licensed health care professionals or certified nurse aides, the provider shall maintain documentation reflecting the individual's current licensure as a health care professional or current certification as a nurse aide. F. **Consequences of noncompliance**. The department or other governmental agency having regulatory enforcement authority over a provider may sanction a provider in accordance with applicable law if the provider fails to make an appropriate and timely inquiry of the registry, or fails to maintain evidence of such inquiry, in connection with the hiring or contracting of an employee; or for employing or contracting any person to work as an employee who is listed on the registry. Such sanctions may include a directed plan of correction, civil monetary penalty not to exceed five thousand dollars ($5000) per instance, or termination or non-renewal of any contract with the department or other governmental agency.		

Standard of Care	Routine Survey Deficiencies October 25 – November 5, 2021	Verification Survey New and Repeat Deficiencies April 18 – 28, 2022
Service Domain: Health and Welfare – *The state, on an ongoing basis, identifies, addresses and seeks to prevent occurrences of abuse, neglect and exploitation. Individuals shall be afforded their basic human rights. The provider supports individuals to access needed healthcare services in a timely manner.*		
Tag # 1A09 Medication Delivery Routine Medication Administration	**Condition of Participation Level Deficiency**	**Condition of Participation Level Deficiency**
Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019 **Chapter 20: Provider Documentation and Client Records 20.6 Medication Administration Record (MAR):** A current Medication Administration Record (MAR) must be maintained in all settings where medications or treatments are delivered. Family Living Providers may opt not to use MARs if they are the sole provider who supports the person with medications or treatments. However, if there are services provided by unrelated DSP, ANS for Medication Oversight must be budgeted, and a MAR must be created and used by the DSP. Primary and Secondary Provider Agencies are responsible for: 1. Creating and maintaining either an electronic or paper MAR in their service setting. Provider Agencies may use the MAR in Therap, but are not mandated to do so. 2. Continually communicating any changes about medications and treatments between Provider Agencies to assure health and safety. 7. Including the following on the MAR: a. The name of the person, a transcription of the physician's or licensed health care provider's orders including the brand and generic names for all ordered routine and PRN medications or treatments, and the diagnoses for which the medications or treatments are prescribed; b. The prescribed dosage, frequency and method or route of administration; times and dates of administration for all ordered routine or PRN prescriptions or treatments; over the counter (OTC) or "comfort"	After an analysis of the evidence, it has been determined there is a significant potential for a negative outcome to occur. Medication Administration Records (MAR) were reviewed for the months of September and October 2021. Based on record review, 4 of 5 individuals had Medication Administration Records (MAR), which contained missing medications entries and/or other errors: Individual #1 October 2021 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Dronabinol 10 mg (2 times daily) – Blank 10/27 (7 AM) • Ibuprofen 200mg – Blank 10/19, 20, 26 and 27, 2021 Individual #2 October 2021 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Oxybutynin Chloride 5mg (2 times daily) – Blank 10/28 (8 AM) Individual #3 September 2021	**New / Repeat Findings:** After an analysis of the evidence, it has been determined there is a significant potential for a negative outcome to occur. Medication Administration Records (MAR) were reviewed for the month of March 2022. Based on record review, 2 of 5 individuals had Medication Administration Records (MAR), which contained missing medications entries and/or other errors: Individual #1 March 2022 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Clozapine 100mg (1 time daily) – Blank 3/31 (1:30 PM) Medication Administration Records contain the following medications. No Physician's Orders were found for the following medications: • CBD Oil (1 time daily) Individual #3 March 2022 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Chlorhexidine 0.12% (2 times daily) – Blank 3/2 (7 PM)

medications or treatments and all self-selected herbal or vitamin therapy; c. Documentation of all time limited or discontinued medications or treatments; d. The initials of the individual administering or assisting with the medication delivery and a signature page or electronic record that designates the full name corresponding to the initials; e. Documentation of refused, missed, or held medications or treatments; f. Documentation of any allergic reaction that occurred due to medication or treatments; and g. For PRN medications or treatments: i. instructions for the use of the PRN medication or treatment which must include observable signs/symptoms or circumstances in which the medication or treatment is to be used and the number of doses that may be used in a 24-hour period; ii. clear documentation that the DSP contacted the agency nurse prior to assisting with the medication or treatment, unless the DSP is a Family Living Provider related by affinity of consanguinity; and iii. documentation of the effectiveness of the PRN medication or treatment. **Chapter 10 Living Care Arrangements** ***10.3.4 Medication Assessment and Delivery:*** Living Supports Provider Agencies must support and comply with: 1. the processes identified in the DDSD AWMD training; 2. the nursing and DSP functions identified in the Chapter 13.3 Part 2- Adult Nursing Services; 3. all Board of Pharmacy regulations as noted in Chapter 16.5 Board of Pharmacy; and	Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Artificial Tears 1.4% (2 times daily) – Blank 9/30 (7 AM and 7 PM) Individual #4 October 2021 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Carbidopa-Levodopa 25-100mg (5 times daily) – Blank 10/27 (6 PM) • Oxcarbazepine 150mg (2 times daily) – Blank 10/27 (8 PM) • Polyethylene Glycol 3350 17 Gram/Dose (1 time daily) Blank 10/27 (8 PM)	

4. documentation requirements in a Medication Administration Record (MAR) as described in Chapter 20.6 Medication Administration Record (MAR). **NMAC 16.19.11.8 MINIMUM STANDARDS:** A. MINIMUM STANDARDS FOR THE DISTRIBUTION, STORAGE, HANDLING AND RECORD KEEPING OF DRUGS: (d) The facility shall have a Medication Administration Record (MAR) documenting medication administered to residents, **including over-the-counter medications.** This documentation shall include: (i) Name of resident; (ii) Date given; (iii) Drug product name; (iv) Dosage and form; (v) Strength of drug; (vi) Route of administration; (vii) How often medication is to be taken; (viii) Time taken and staff initials; (ix) Dates when the medication is discontinued or changed; (x) The name and initials of all staff administering medications. **Model Custodial Procedure Manual** ***D. Administration of Drugs*** Unless otherwise stated by practitioner, patients will not be allowed to administer their own medications. Document the practitioner's order authorizing the self-administration of medications. All PRN (As needed) medications shall have complete detail instructions regarding the administering of the medication. This shall include: ➢ symptoms that indicate the use of the medication, ➢ exact dosage to be used, and ➢ the exact amount to be used in a 24-hour period.		

Tag # 1A09.1 Medication Delivery PRN Medication Administration	Condition of Participation Level Deficiency	Condition of Participation Level Deficiency
Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019 **Chapter 20: Provider Documentation and Client Records 20.6 Medication Administration Record (MAR):** A current Medication Administration Record (MAR) must be maintained in all settings where medications or treatments are delivered. Family Living Providers may opt not to use MARs if they are the sole provider who supports the person with medications or treatments. However, if there are services provided by unrelated DSP, ANS for Medication Oversight must be budgeted, and a MAR must be created and used by the DSP. Primary and Secondary Provider Agencies are responsible for: 1. Creating and maintaining either an electronic or paper MAR in their service setting. Provider Agencies may use the MAR in Therap, but are not mandated to do so. 2. Continually communicating any changes about medications and treatments between Provider Agencies to assure health and safety. 7. Including the following on the MAR: a. The name of the person, a transcription of the physician's or licensed health care provider's orders including the brand and generic names for all ordered routine and PRN medications or treatments, and the diagnoses for which the medications or treatments are prescribed; b. The prescribed dosage, frequency and method or route of administration; times and dates of administration for all ordered routine or PRN prescriptions or treatments; over the counter (OTC) or "comfort" medications or treatments and all self-selected herbal or vitamin therapy; c. Documentation of all time limited or discontinued medications or treatments;	After an analysis of the evidence, it has been determined there is a significant potential for a negative outcome to occur. Medication Administration Records (MAR) were reviewed for the months of 10/2021. Based on record review, 3 of 5 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #1 October 2021 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Tylenol 500mg – PRN – 10/1 -13 (given 1 time) • Alprazolam 2mg – PRN – 10/4, 11 (given 2 times), 10/5, 10 (given 1 time), 10/13 (given 3 times) No Time of Administration was noted on the Medication Administration Record for the following PRN medication: • Tylenol 500mg – PRN – 10/1 -13 Individual #4 October 2021 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Milk of Magnesia 1200mg – PRN – 10/19 (given 1 time) No Time of Administration was noted on the Medication Administration Record for the following PRN medication:	After an analysis of the evidence, it has been determined there is a significant potential for a negative outcome to occur. Medication Administration Records (MAR) were reviewed for the month of March 2022. Based on record review, 3 of 5 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #1 March 2022 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Propranolol HCL 20mg – PRN – 3/20 - 21, 24, 25 (given 1 time), 3/31 (given 2 times) • Clorazepate Dipotassium 15mg – PRN – 3/12, 21, 24 - 25 (given 1 time) Medication Administration Records contain the following medications. No Physician's Orders were found for the following medications: • Oxycodone 10mg (PRN) Individual #3 March 2022 Medication Administration Records contain the following medications. No Physician's Orders were found for the following medications: • Procto-med HC 2.5% CRM/PE App: (Hydrocortisone) (PRN) Individual #5 March 2022

d. The initials of the individual administering or assisting with the medication delivery and a signature page or electronic record that designates the full name corresponding to the initials; e. Documentation of refused, missed, or held medications or treatments; f. Documentation of any allergic reaction that occurred due to medication or treatments; and g. For PRN medications or treatments: i. instructions for the use of the PRN medication or treatment which must include observable signs/symptoms or circumstances in which the medication or treatment is to be used and the number of doses that may be used in a 24-hour period; ii. clear documentation that the DSP contacted the agency nurse prior to assisting with the medication or treatment, unless the DSP is a Family Living Provider related by affinity of consanguinity; and iii. documentation of the effectiveness of the PRN medication or treatment. **Chapter 10 Living Care Arrangements** ***10.3.4 Medication Assessment and Delivery:*** Living Supports Provider Agencies must support and comply with: 1. the processes identified in the DDSD AWMD training; 2. the nursing and DSP functions identified in the Chapter 13.3 Part 2- Adult Nursing Services; 3. all Board of Pharmacy regulations as noted in Chapter 16.5 Board of Pharmacy; and 4. documentation requirements in a Medication Administration Record (MAR) as described in Chapter 20.6 Medication Administration Record (MAR).	• Milk of Magnesia 1200mg – PRN – 10/19 (given 1 time) Individual #5 October 2021 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Tylenol 500mg – PRN – 10/20 – 27 (given 1 time) No Time of Administration was noted on the Medication Administration Record for the following PRN medication: • Tylenol 500mg – PRN – 10/20 – 27 (given 1 time)	Medication Administration Records contain the following medications. No Physician's Orders were found for the following medications: • Milk of Magnesia Saline Laxative (PRN)

Tag # 1A09.2 Medication Delivery Nurse Approval for PRN Medication	Condition of Participation Level Deficiency	Condition of Participation Level Deficiency
Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019 **Chapter 13 Nursing Services: *13.2.12 Medication Delivery:*** Nurses are required to: 1. Be aware of the New Mexico Nurse Practice Act, and Board of Pharmacy standards and regulations. 2. Communicate with the Primary Care Practitioner and relevant specialists regarding medications and any concerns with medications or side effects. 3. Educate the person, guardian, family, and IDT regarding the use and implications of medications as needed. 4. Administer medications when required, such as intravenous medications; other specific injections; via NG tube; non-premixed nebulizer treatments or new prescriptions that have an ordered assessment. 5. Monitor the MAR or treatment records at least monthly for accuracy, PRN use and errors. 6. Respond to calls requesting delivery of PRNs from AWMD trained DSP and non-related (surrogate or host) Family Living Provider Agencies. 7. Assure that orders for PRN medications or treatments have: a. clear instructions for use; b. observable signs/symptoms or circumstances in which the medication is to be used or withheld; and c. documentation of the response to and effectiveness of the PRN medication administered. 8. Monitor the person's response to the use of routine or PRN pain medication and contact the prescriber as needed regarding its effectiveness. 9. Assure clear documentation when PRN medications are used, to include: a. DSP contact with nurse prior to assisting with medication. i. The only exception to prior consultation	After an analysis of the evidence, it has been determined there is a significant potential for a negative outcome to occur. Based on record review, the Agency did not maintain documentation of PRN authorization as required by standard for 2 of 5 Individuals. Individual #1 October 2021 No documentation of the verbal authorization from the Agency nurse prior to each administration/assistance of PRN medication was found for the following PRN medication: • Alprazolam 2mg – PRN – 10/4/2021 (given 2 times), 10/5, 10 (given 1 time), 10/13 (given 3 times) Individual #5 October 2021 No documentation of the verbal authorization from the Agency nurse prior to each administration/assistance of PRN medication was found for the following PRN medication: • Tylenol 500mg – PRN – 10/23 – 25 (given 1 time)	After an analysis of the evidence, it has been determined there is a significant potential for a negative outcome to occur. Based on record review, the Agency did not maintain documentation of PRN authorization as required by standard for 1 of 5 Individuals. Individual #1 March 2022 No documentation of the verbal authorization from the Agency nurse prior to each administration/assistance of PRN medication was found for the following PRN medication: • Propranolol HCL 20mg – PRN – 3/8 (given 1 time) • Clorazepate Dipotassium 15mg – PRN – 3/12 (given 1 time)

with the agency nurse is to administer selected emergency medications as listed on the Publications section of the DOH-DDSD - Clinical Services Website https://nmhealth.org/about/ddsd/pgsv/clinical/ . b. Nursing instructions for use of the medication. c. Nursing follow-up on the results of the PRN use. d. When the nurse administers the PRN medication, the reasons why the medications were given and the person's response to the medication.		

Standard of Care	Routine Survey Deficiencies October 25 – November 5, 2021	Verification Survey New and Repeat Deficiencies April 18 – 28, 2022
Service Domain: Service Plans: ISP Implementation – *Services are delivered in accordance with the service plan, including type, scope, amount, duration and frequency specified in the service plan.*		
Tag # 1A08 Administrative Case File (Other Required Documents)	**Standard Level Deficiency**	**COMPLETE**
Tag # 1A08.3 Administrative Case File: Individual Service Plan / ISP Components	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # 1A08.1 Administrative and Residential Case File: Progress Notes	**Standard Level Deficiency**	**COMPLETE**
Tag # 1A32 Administrative Case File: Individual Service Plan Implementation	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # 1A32.1 Administrative Case File: Individual Service Plan Implementation *(Not Completed at Frequency)*	**Standard Level Deficiency**	**COMPLETE**
Tag # 1A32.2 Individual Service Plan Implementation (Residential Implementation)	**Standard Level Deficiency**	**COMPLETE**
Tag # LS14 Residential Service Delivery Site Case File (ISP and Healthcare Requirements)	**Condition of Participation Level Deficiency**	**COMPLETE**
Service Domain: Qualified Providers – *The State monitors non-licensed/non-certified providers to assure adherence to waiver requirements. The State implements its policies and procedures for verifying that provider training is conducted in accordance with State requirements and the approved waiver.*		
Tag # 1A43.1 General Events Reporting: Individual Reporting	**Standard Level Deficiency**	**COMPLETE**
Service Domain: Health and Welfare – *The state, on an ongoing basis, identifies, addresses and seeks to prevent occurrences of abuse, neglect and exploitation. Individuals shall be afforded their basic human rights. The provider supports individuals to access needed healthcare services in a timely manner.*		
Tag # 1A08.2 Administrative Case File: Healthcare Requirements & Follow-up	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # 1A03 Continuous Quality Improvement System & Key Performance Indicators (KPIs)	**Standard Level Deficiency**	**COMPLETE**
Tag # 1A09.0 Medication Delivery Routine Medication Administration	**Standard Level Deficiency**	**COMPLETE**
Tag # 1A15.2 Administrative Case File: Healthcare Documentation (Therap and Required Plans)	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # LS25 Residential Health & Safety (Supported Living / Family Living / Intensive Medical Living)	**Standard Level Deficiency**	**COMPLETE**
Service Domain: Medicaid Billing/Reimbursement – *State financial oversight exists to assure that claims are coded and paid for in accordance with the reimbursement methodology specified in the approved waiver.*		
Tag #1A12 All Services Reimbursement	**No Deficient Practices Found**	**COMPLETE**

	Verification Survey Plan of Correction, On-going QA/QI and Responsible Party	Completion Date
Tag # 1A26 Consolidated On-line Registry Employee Abuse Registry	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)*: → **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to affect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)*: →	
Tag # 1A09 Medication Delivery Routine Medication Administration	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)*: → **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to affect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)*: →	

Tag # 1A09.1 Medication Delivery PRN Medication Administration	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)*: → **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to affect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)*: →	
Tag # 1A09.2 Medication Delivery Nurse Approval for PRN Medication	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)*: → **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to affect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)*: →	

NEW MEXICO
Department of Health
Division of Health Improvement

MICHELLE LUJAN GRISHAM
Governor

DAVID R. SCRASE, M.D.
Acting Cabinet Secretary

Date: June 16, 2022

To: Christine Chapman, DSP Supervisor / SC / Executive Director / Owner

Provider: Safe Harbor, Inc.
Address: 825 Quesenberry St.
State/Zip: Las Cruces, New Mexico 88005

E-mail Address: garychpm@aol.com

Region: Southwest
Routine Survey: October 25 – November 5, 2021
Verification Survey: April 18 – 28, 2022

Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: Supported Living and Customized Community Supports

Survey Type: Verification

Dear Ms. Chapman:

The Division of Health Improvement/Quality Management Bureau has received, reviewed and approved the supporting documents you submitted for your Plan of Correction. The documents you provided verified that all previously cited survey Deficiencies have been corrected.

The Plan of Correction process is now complete.

Furthermore, your agency is now determined to be in Compliance with all Conditions of Participation.

To maintain ongoing compliance with standards and regulations, continue to use the Quality Assurance (self-auditing) processes you described in your Plan of Correction.

Consistent use of these Quality Assurance processes will enable you to identify and promptly respond to problems, enhance your service delivery, and result in fewer deficiencies cited in future QMB surveys.

Thank you for your cooperation with the Plan of Correction process, for striving to come into compliance with standards and regulations, and for helping to provide the health, safety and personal growth of the people you serve.

Sincerely,

Monica Valdez, BS

Monica Valdez, BS
Healthcare Surveyor Advanced/Plan of Correction Coordinator
Quality Management Bureau/DHI

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