Date:	March 16, 2022
To:	Michael Gemme, Administrator
Provider: Address: State/Zip:	Los Lunas Community Program 1000 Main Street, NW Los Lunas, New Mexico 87031
E-mail Address:	Michael.Gemme@state.nm.us
CC: E-mail Address:	Joseph Chavez, QA Director Joseph.Chavez12@state.nm.us
Region: Routine Survey: Verification Survey:	Metro September 13 - 24, 2021 February 14 – 23, 2022
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	Supported Living, Intensive Medical Living, Customized Community Supports, and Community Integrated Employment Services
Survey Type:	Verification
Team Leader:	Verna Newman-Sikes, AA, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau
Team Members:	Lei Lani Nava, MPH, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Dear Mr. Gemme;

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 September 13 – 24, 2021*.

Determination of Compliance

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

Compliance: This determination is based on your agency's compliance with Condition of Participation level and Standard level requirements. Deficiencies found only affect a small percentage of the Individuals on the survey sample *(refer to Attachment D for details)*. The attached QMB Report of Findings indicates Standard Level deficiencies identified and requires implementation of a Plan of Correction.

The following tags are identified as Standard Level:

- Tag # 1A22 Agency Personnel Competency ***(New / Repeat Findings)***
- Tag # 1A09.0 Medication Delivery Routine Medication Administration ***(New / Repeat Findings)***

- Tag # 1A09.1 Medication Delivery PRN Medication Administration ***(New / Repeat Findings)***

However, due to the new/repeat deficiencies your agency will be referred to the Internal Review Committee (IRC). Your agency will also 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 or email at: MonicaE.Valdez@state.nm.us 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,

Verna Newman-Sikes, AA

Verna Newman-Sikes, AA
Team Lead/Healthcare Surveyor
Division of Health Improvement
Quality Management Bureau

Survey Process Employed:

Administrative Review Start Date:	February 14, 2022
Contact:	**Los Lunas Community Program (NMDOH)** Joseph Chavez, QA Director **DOH/DHI/QMB** Verna Newman-Sikes, AA, Team Lead/Healthcare Surveyor
Exit Conference Date:	February 23, 2022
Present:	**Los Lunas Community Program (NMDOH)** Michael Gemme, Administrator Joseph Chavez, QA Director Onecimo Mirabal, Program Director Kent Montoya, Director of Nursing Anthony Fragua, Community Inclusion Manager **DOH/DHI/QMB** Verna Newman-Sikes, AA, Team Lead/Healthcare Surveyor Lei Lani Nava, MPH, Healthcare Surveyor Amanda Castañeda-Holguin, MPA, Healthcare Surveyor Supervisor **DDSD - Metro Regional Office** Alicia Otolo, Social and Community Service Coordinator
Administrative Locations Visited:	0 *(Note: No administrative locations visited due to COVID- 19 Public Health Emergency.)*
Total Sample Size:	18 7 - *Jackson* Class Members 11 - Non-*Jackson* Class Members 13 - Supported Living 5 - Intensive Medical Living Supports 16 - Customized Community Supports 6 - Community Integrated Employment
Persons Served Records Reviewed	18
Direct Support Personnel Records Reviewed	141
Direct Support Personnel Interviewed during Routine Survey	12 *(Note: Interviews conducted by video / phone due to COVID- 19 Public Health Emergency)*
Service Coordinator Records Reviewed	3
Nurse Interview completed during Routine Survey	1

Administrative Processes and Records Reviewed:

- Medicaid Billing/Reimbursement Records for all Services Provided
- Accreditation Records
- 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: **Los Lunas Community Program (NMDOH) - Metro Region**
Program: Developmental Disabilities Waiver
Service: Supported Living, Intensive Medical Living, Customized Community Supports, and Community Integrated Employment Services
Survey Type: Verification
Routine Survey: **September 13 - 24, 2021**
Verification Survey: **February 14 – 23, 2022**

Standard of Care	Routine Survey Deficiencies September 13 - 24, 2021	Verification Survey New and Repeat Deficiencies February 14 – 23, 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 # 1A22 Agency Personnel Competency	**Condition of Participation Level Deficiency**	**Standard Level Deficiency**
Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019 **Chapter 13: Nursing Services *13.2.11 Training and Implementation of Plans:*** 1. RNs and LPNs are required to provide Individual Specific Training (IST) regarding HCPs and MERPs. 2. The agency nurse is required to deliver and document training for DSP/DSS regarding the healthcare interventions/strategies and MERPs that the DSP are responsible to implement, clearly indicating level of competency achieved by each trainee as described in Chapter 17.10 Individual-Specific Training. **Chapter 17: Training Requirement** **17.10 Individual-Specific Training:** The following are elements of IST: defined standards of performance, curriculum tailored to teach skills and knowledge necessary to meet those standards of performance, and formal examination or demonstration to verify standards of performance, using the established DDSD training levels of awareness, knowledge, and skill. Reaching an **awareness level** may be accomplished by reading plans or other information. The trainee is cognizant of information related to a person's specific condition. Verbal or written recall of basic information or knowing where to access the	After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur. Based on interview, the Agency did not ensure training competencies were met for 3 of 12 Direct Support Personnel. **When DSP were asked, if the Individual had a Positive Behavioral Supports Plan (PBSP), have you been trained on the PBSP and what does the plan cover, the following was reported:** • DSP #517 stated, "Yes, give her space, redirect her, we have to stand out of her way. Time to breathe, come and address her in a different way, do incentives like go out and maybe give her a treat." According to the Individual Specific Training Section of the ISP, the Individual does not require a Positive Behavioral Supports Plan. (Individual #19) **When DSP were asked, if they received training on the Individual's Behavioral Crisis Intervention Plan (BCIP) and if so, what the plan covered, the following was reported:** • DSP #558 stated, "I don't think he has one." According to the Individual Specific Training	**New/Repeat Finding:** Per the Plan of Correction approved on 11/22/2021, "#587 will be retrained. Service Coordinators will retrain DSP by November 30, 2021". No evidence training was completed by 11/30/2021. *(Note: Training was completed during the on-site survey. Provider please complete POC for ongoing QA/QI.)*

information can verify awareness. Reaching a **knowledge level** may take the form of observing a plan in action, reading a plan more thoroughly, or having a plan described by the author or their designee. Verbal or written recall or demonstration may verify this level of competence. Reaching a **skill level** involves being trained by a therapist, nurse, designated or experienced designated trainer. The trainer shall demonstrate the techniques according to the plan. Then they observe and provide feedback to the trainee as they implement the techniques. This should be repeated until competence is demonstrated. Demonstration of skill or observed implementation of the techniques or strategies verifies skill level competence. Trainees should be observed on more than one occasion to ensure appropriate techniques are maintained and to provide additional coaching/feedback. Individuals shall receive services from competent and qualified Provider Agency personnel who must successfully complete IST requirements in accordance with the specifications described in the ISP of each person supported. 1. IST must be arranged and conducted at least annually. IST includes training on the ISP Desired Outcomes, Action Plans, strategies, and information about the person's preferences regarding privacy, communication style, and routines. More frequent training may be necessary if the annual ISP changes before the year ends. 2. IST for therapy-related WDSI, HCPs, MERPs, CARMPs, PBSA, PBSP, and BCIP, must occur at least annually and more often if plans change, or if monitoring by the plan author or agency finds incorrect implementation, when new DSP or CM are assigned to work with a person, or when an existing DSP or CM requires a refresher. 3. The competency level of the training is based on the IST section of the ISP. 4. The person should be present for and involved in IST whenever possible. 5. Provider Agencies are responsible for tracking of IST requirements.	Section of the ISP, the individual has a Behavioral Crisis Intervention Plan. (Individual #2) • DSP #587 stated, "No, I don't think so I have never heard of that for him ever." According to the Individual Specific Training Section of the ISP, the individual has a Behavioral Crisis Intervention Plan. (Individual #17) **When DSP were asked, if the Individual had any food and / or medication allergies that could be potentially life threatening, the following was reported:** • DSP #517 stated, "All Antibiotics." As indicated by the Health Passport the individual is also allergic to Codeine and Darvocet-N. (Individual #11)	

6. Provider Agencies must arrange and ensure that DSP's are trained on the contents of the plans in accordance with timelines indicated in the Individual-Specific Training Requirements: Support Plans section of the ISP and notify the plan authors when new DSP are hired to arrange for trainings. 7. If a therapist, BSC, nurse, or other author of a plan, healthcare or otherwise, chooses to designate a trainer, that person is still responsible for providing the curriculum to the designated trainer. The author of the plan is also responsible for ensuring the designated trainer is verifying competency in alignment with their curriculum, doing periodic quality assurance checks with their designated trainer, and re-certifying the designated trainer at least annually and/or when there is a change to a person's plan.		

Standard of Care	Routine Survey Deficiencies September 13 - 24, 2021	Verification Survey New and Repeat Deficiencies February 14 – 23, 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 # 1A09.0 Medication Delivery Routine Medication Administration	**Standard Level Deficiency**	**Standard 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"	Medication Administration Records (MAR) were reviewed for the months of August and September 2021. Based on record review, 2 of 19 individuals had Medication Administration Records (MAR), which contained missing medications entries and/or other errors: Individual #4 August 2021 Medication Administration Records did not contain the strength of the medication which is to be given: • Acidophilus Probiotic (1 time daily) Individual #8 September 2021 Medication Administration Records did not contain the strength of the medication which is to be given: • Digestive Enzyme (3 times daily) Medication Administration Records did not contain the route of administration for the following medications: • Melatonin 3mg (1 time daily)	**New/Repeat Finding:** Medication Administration Records (MAR) were reviewed for the month of January 2022. Based on record review, 1 of 18 individuals had Medication Administration Records (MAR), which contained missing medications entries and/or other errors: Individual #4 January 2022 Medication Administration Records did not contain the strength of the medication which is to be given: • Daily-Vite (1 time daily)

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		

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	Standard 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; d. The initials of the individual administering	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 August and September 2021. Based on record review, 6 of 19 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #6 August 2021 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Benadryl 25 mg – PRN – 8/31 (given 1 time) • Pepto-Bismol Suspension 262 mg/15 ml PRN – 8/20 (given 1 time) Individual #9 August 2021 Medication Administration Records contain the following medications. No Physician's Orders were found for the following medications: • Pepto-Bismol Suspension (PRN) Individual #11 August 2021 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Acetaminophen 325 mg. – PRN – 8/17, 25, 27, 28 (given 1 time) • Cyclobenzaprine 10mg. – PRN – 8/5, 25, 27 (given 1 time)	**New/Repeat Finding:** Medication Administration Records (MAR) were reviewed for the month of January 2022. Based on record review, 1 of 18 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #11 January 2022 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Cyclobenzaprine 10 mg – PRN – 1/8 & 12 (given 1 time)

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).	September 2021 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Acetaminophen 325 mg— PRN – 9/10, 14, 15 (given 1 time) 9/13 (given 2 times) • Cyclobenzaprine 10 mg – PRN – 9/10 (given 1 time) Individual #12 August 2021 As indicated by the Medication Administration Records the individual is to apply Boudreaux paste 40% (PRN) as needed for skin irritation. According to the Physician's Orders, Boudreaux 16% ointment (PRN) apply to affected area topically at every brief change. Medication Administration Record and Physician's Orders do not match. Medication Administration Records contain the following medications. No Physician's Orders were found for the following medications: • Prune Juice (PRN) Individual #16 August 2021 No evidence of documented Signs/Symptoms were found for the following PRN medication: • Excedrin Migraine 250 mg-65 mg – PRN – 8/31 (given 1 time). No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Excedrin Migraine 250 mg-65 mg – PRN – 8/31 (given 1 time). Medication Administration Records Indicated Excedrin Migraine 250 mg-65 mg was given. MAR	

	did not indicate the exact dosage each time the med was assisted or administered for the following dates: • 8/31 Individual #19 August 2021 Medication Administration Records contain the following medications. No Physician's Orders were found for the following medications: • Acetaminophen 325 mg (PRN) • Ibuprofen 200 mg (PRN) • MI Acid Suspension 200-200-20 mg/5ml (PRN) September 2021 No evidence of documented Signs/Symptoms were found for the following PRN medication: • Acetaminophen 325 mg – PRN – 9/7 (given 1 time) No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Acetaminophen 325 mg – PRN – 9/7 (given 1 time)	

Standard of Care	Routine Survey Deficiencies September 13 - 24, 2021	Verification Survey New and Repeat Deficiencies February 14 – 23, 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 # 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 Medication Delivery Routine Medication Administration	**Standard Level Deficiency**	**COMPLETE**
Tag # 1A09.1.0 Medication Delivery PRN Medication Administration	**Standard Level Deficiency**	**COMPLETE**
Tag # 1A09.2 Medication Delivery Nurse Approval for PRN Medication	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # 1A15.2 Administrative Case File: Healthcare Documentation (Therap and Required Plans)	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # 1A31 Client Rights / Human Rights	**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 # LS26 Supported Living Reimbursement	**Standard Level Deficiency**	**COMPLETE**

	Verification Survey Plan of Correction, On-going QA/QI and Responsible Party	Completion Date
Tag # 1A22 **Agency Personnel Competency**	**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 # 1A09.0 **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?)*: →	

NEW MEXICO
Department of Health
Division of Health Improvement

MICHELLE LUJAN GRISHAM
Governor

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

Date: April 8, 2022

To: Michael Gemme, Administrator

Provider: Los Lunas Community Program
Address: 1000 Main Street, NW
State/Zip: Los Lunas, New Mexico 87031

E-mail Address: Michael.Gemme@state.nm.us

CC: Joseph Chavez, QA Director
E-mail Address: Joseph.Chavez12@state.nm.us

Region: Metro
Routine Survey: September 13 - 24, 2021
Verification Survey: February 14 – 23, 2022

Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: Supported Living, Intensive Medical Living, Customized Community Supports, and Community Integrated Employment Services

Survey Type: Verification

Dear Mr. Gemme:

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

Q.22.3.DDW.D1977.5.VER.09.22.098