MICHELLE LUJAN GRISHAM Governor

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

Date:	December 13, 2022
То:	Claudine Valerio-Salazar, Executive Director
Provider: Address: State/Zip:	EnSuenos Y Los Angelitos Development Center 1030 Salazar Rd. Taos, New Mexico 87571
E-mail Address:	cvs@eladc.org
CC: E-Mail Address:	Analisa Rugelio, Supported Living Coordinator / QI Coordinator / Trainer avigil@eladc.org
Region: Routine Survey: Verification Survey:	Northeast May 16 - 26, 2022 November 28 – December 8, 2022
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	Supported Living, Customized Community Supports, and Community Integrated Employment Services
Survey Type:	Verification
Team Leader:	Lei Lani Nava, MPH, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau
Team Members:	Verna Newman-Sikes, AA, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Dear Ms. Claudine Valerio-Salazar,

NEW MEXICO

Department of Health

Division of Health Improvement

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 May 16 – 26, 2022.*

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

<u>Partial Compliance with Standard Level Tags and Conditions of Participation Level Tags</u>: 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 # 1A08.2 Administrative Case File: Healthcare Requirements & Follow-up (Repeat Finding)
- Tag # 1A09 Medication Delivery Routine Medication Administration (New / Repeat Finding)

DIVISION OF HEALTH IMPROVEMENT

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However, due to the new/repeat deficiencies your agency will be required to contact your DDSD Regional Office for technical assistance and complete the Plan of Correction document attached at the end of this report. Please note that verification survey POC does not have the same timelines as the routine POC process. You are required to complete the Plan of Correction within 10 business days from the receipt of this letter. Once your POC is <u>approved</u>, you will have 3 business days to submit your documents.

Plan of Correction:

The POC must include the following:

- 1. Evidence your agency has contacted your DDSD Regional Office for technical assistance;
- 2. A Plan of Correction detailing Corrective Action for any new/repeat deficiencies and Quality Assurance/Quality Improvement processes to prevent your agency from receiving deficiencies in the future;

Submission of your Plan of Correction:

Please submit your agency's Plan of Correction using the format at the end of this report within 10 business days of receipt of this letter to the parties below:

1. Quality Management Bureau, Monica Valdez, Plan of Correction Coordinator MonicaE.Valdez@doh.nm.gov

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

As a reminder, you are to submit your documents electronically within 3 business days of the POC being approved.

- Documents not containing Protected Health Information (PHI) may be submitted to the POC Coordinator at Monicae.valdez@doh.nm.gov
- Documents <u>containing Protected Health Information (PHI)</u> may be submitted via Therap® S-Comm or you may contact the POC Coordinator at <u>Monicae.valdez@doh.nm.gov</u> to initiate a secure State email where you will reply and attach your documents. When possible, please submit requested documentation using a "zipped/compressed" file to reduce file size.
- You may also submit documents via another electronic format, i.e., flash drive. Please contact the POC Coordinator Monica Valdez at 505-273-1930 or <u>Monicae.valdez@doh.nm.gov</u> to make arrangements for delivery of your documents.

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

Please contact the Plan of Correction Coordinator, Monica Valdez at 505-273-1930 or <u>Monicae.valdez@doh.nm.gov</u> if you have questions about the survey or the report. 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:

Survey Process Employed:	
Administrative Review Start Date:	November 28, 2022
Contact:	EnSuenos Y Los Angelitos Development Center Claudine Valerio-Salazar, Executive Director
	DOH/DHI/QMB Lei Lani Nava, Team Lead/Healthcare Surveyor
On-site Entrance Conference Date:	Entrance conference was waived by provider.
Exit Conference Date:	December 8, 2022
Present:	EnSuenos Y Los Angelitos Development Center Claudine Valerio-Salazar, Executive Director Kimberly Tofoya, Human Resource Manager Joseph Rivera, Day Service Manager / Service Coordinator Beverly Rodriguez, Day Service / Residential Assistant Manager Analisa Vigil-Rugelio, Supported Living 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
	<u>DDSD - NE Regional Office</u> Angela Pacheco, Regional Director Kim Hamstra, Social Community Service Coordinator
Total Sample Size:	5
	1 - <i>Former Jackson</i> Class Members 4 - Non- <i>Jackson</i> Class Members
	4 - Supported Living 5 - Customized Community Supports 1 - Community Integrated Employment
Direct Support Personnel Records Reviewed	15 (Note: One DSP performs multiple roles as a Day Service Manager and Service Coordinator)
Direct Support Personnel Interviewed during Routine Survey	7 (Note: Interviews conducted by video / phone due to COVID- 19 Public Health Emergency)
Service Coordinator Records Reviewed	2 (Note: One Service Coordinator performs dual roles as a DSP and Day Service Manager)
Nurse Interview completed during Routine Survey	1
Administrative Processes and Records Review	red:

- 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: ^oIndividual Service Plans

°Progress on Identified Outcomes

- °Healthcare Plans
- °Medication Administration Records
- ^oMedical Emergency Response Plans
- °Therapy Evaluations and Plans
- ^oHealthcare Documentation Regarding Appointments and Required Follow-Up ^oOther 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

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 Living Care Arrangements and Community Inclusion 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

<u>Service Domain: Health, Welfare and Safety -</u> 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:

- The Informal Reconsideration of the Finding (IRF) request must be received in writing to the QMB Bureau Chief <u>within 10 business days</u> 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: <u>https://nmhealth.org/about/dhi/cbp/irf/</u>
- 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 <u>valerie.valdez@state.nm.us</u> 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.

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				Weighting			
Determination	LC	W		MEDIUM		н	IIGH
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 <u>and</u> 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:EnSuenos Y Los Angelitos Development Center - Northeast RegionProgram:Developmental Disabilities WaiverService:Supported Living, Customized Community Supports, and Community Integrated Employment ServicesSurvey Type:VerificationRoutine Survey:May 16 - 26, 2022Verification Survey:November 28 - December 8, 2022

Standard of Care	Routine Survey Deficiencies May 16 – 26, 2022	Verification Survey New and Repeat Deficiencies November 28 – December 8, 2022
	an ongoing basis, identifies, addresses and seeks to p	
	The provider supports individuals to access needed hea	
Tag # 1A08.2 Administrative Case File:	Condition of Participation Level Deficiency	Condition of Participation Level Deficiency
Healthcare Requirements & Follow-up Developmental Disabilities (DD) Waiver Service	After an analysis of the evidence, it has been	Repeat Finding:
Standards 2/26/2018; Re-Issue: 12/28/2018; Eff	determined there is a significant potential for a	Repeat I mullig.
1/1/2019	negative outcome to occur.	After an analysis of the evidence, it has been
Chapter 3 Safeguards: 3.1.1 Decision		determined there is a significant potential for a
Consultation Process (DCP): Health decisions are	Based on record review, the Agency did not provide	negative outcome to occur.
the sole domain of waiver participants, their	documentation of annual physical examinations	
guardians or healthcare decision makers.	and/or other examinations as specified by a licensed	Based on record review, the Agency did not provide
Participants and their healthcare decision makers	physician for 2 of 5 individuals receiving Living Care	documentation of annual physical examinations
can confidently make decisions that are compatible	Arrangements and Community Inclusion.	and/or other examinations as specified by a licensed
with their personal and cultural values. Provider		physician for 1 of 5 individuals receiving Living Care
Agencies are required to support the informed	Review of the administrative individual case files	Arrangements and Community Inclusion.
decision making of waiver participants by supporting	revealed the following items were not found,	
access to medical consultation, information, and	incomplete, and/or not current:	Review of the administrative individual case files
other available resources according to the following:		revealed the following items were not found,
1. The DCP is used when a person or his/her	Living Care Arrangements / Community	incomplete, and/or not current:
guardian/healthcare decision maker has concerns,	Inclusion (Individuals Receiving Multiple	Living Care Arrangements / Community
needs more information about health-related issues, or has decided not to follow all or part of an order,	Services):	Living Care Arrangements / Community Inclusion (Individuals Receiving Multiple
recommendation, or suggestion. This includes, but	Annual Physical:	Services):
is not limited to:	• Not Found (#3, 5)	Services).
a. medical orders or recommendations from the		Annual Physical:
Primary Care Practitioner, Specialists or other	Dental Exam:	• Not Found (#5)
licensed medical or healthcare practitioners	 Individual #5 - As indicated by collateral 	
such as a Nurse Practitioner (NP or CNP),	documentation reviewed, Exam was completed on	
Physician Assistant (PA) or Dentist;	3/14/2022. Exam was not linked / attached in	
b. clinical recommendations made by	Therap. (Note: Linked / attached in Therap during	
registered/licensed clinicians who are either	the on-site survey. Provider please complete POC	
members of the IDT or clinicians who have	for ongoing QA/QI.)	
performed an evaluation such as a video-		

fluoroscopy;

- health related recommendations or suggestions from oversight activities such as the Individual Quality Review (IQR) or other DOH review or oversight activities; and
- d. recommendations made through a Healthcare Plan (HCP), including a Comprehensive Aspiration Risk Management Plan (CARMP), or another plan.

2. When the person/guardian disagrees with a recommendation or does not agree with the implementation of that recommendation, Provider Agencies follow the DCP and attend the meeting coordinated by the CM. During this meeting:

- a. Providers inform the person/guardian of the rationale for that recommendation, so that the benefit is made clear. This will be done in layman's terms and will include basic sharing of information designed to assist the person/guardian with understanding the risks and benefits of the recommendation.
- b. The information will be focused on the specific area of concern by the person/guardian.
 Alternatives should be presented, when available, if the guardian is interested in considering other options for implementation.
- c. Providers support the person/guardian to make an informed decision.
- d. The decision made by the person/guardian during the meeting is accepted; plans are modified; and the IDT honors this health decision in every setting.

Chapter 20: Provider Documentation and Client Records: 20.2 Client Records Requirements: All DD Waiver Provider Agencies are required to create and maintain individual client records. The contents of client records vary depending on the unique needs of the person receiving services and the resultant information produced. The extent of documentation required for individual client records

Urgent Care:

 Individual #5 - As indicated by collateral documentation reviewed, Exam was completed on 2/13/2022. Exam was not linked / attached in Therap. (Note: Linked / attached in Therap during the on-site survey. Provider please complete POC for ongoing QA/QI.)

Primary Care:

• Individual #5 - As indicated by collateral documentation reviewed, Exam was completed on 1/4/2022. Exam was not linked / attached in Therap. (Note: Linked / attached in Therap during the on-site survey. Provider please complete POC for ongoing QA/QI.)

Wound Care:

- Individual #5 As indicated by collateral documentation reviewed, Appointment was completed on 3/3/2022. Appointment was not linked / attached in Therap. (Note: Linked / attached in Therap during the on-site survey. Provider please complete POC for ongoing QA/QI.)
- Individual #5 As indicated by collateral documentation reviewed, Appointment was completed on 4/5/2022. Appointment was not linked / attached in Therap. (Note: Linked / attached in Therap during the on-site survey. Provider please complete POC for ongoing QA/QI.)
- Individual #5 As indicated by collateral documentation reviewed, Appointment was completed on 4/26/2022. Appointment was not linked / attached in Therap. (Note: Linked / attached in Therap during the on-site survey. Provider please complete POC for ongoing QA/QI.)

per service type depends on the location of the file,	
the type of service being provided, and the	
information necessary.	
DD Waiver Provider Agencies are required to	
adhere to the following:	
1. Client records must contain all documents	
essential to the service being provided and essential	
to ensuring the health and safety of the person	
during the provision of the service.	
2. Provider Agencies must have readily accessible	
records in home and community settings in paper or	
electronic form. Secure access to electronic records	
through the Therap web-based system using	
computers or mobile devices is acceptable.	
3. Provider Agencies are responsible for ensuring	
that all plans created by nurses, RDs, therapists or	
BSCs are present in all needed settings.	
4. Provider Agencies must maintain records of all	
documents produced by agency personnel or	
contractors on behalf of each person, including any	
routine notes or data, annual assessments, semi-	
annual reports, evidence of training	
provided/received, progress notes, and any other	
interactions for which billing is generated.	
5. Each Provider Agency is responsible for	
maintaining the daily or other contact notes	
documenting the nature and frequency of service	
delivery, as well as data tracking only for the	
services provided by their agency.	
6. The current Client File Matrix found in Appendix	
A Client File Matrix details the minimum	
requirements for records to be stored in agency	
office files, the delivery site, or with DSP while	
providing services in the community.	
7. All records pertaining to JCMs must be retained permanently and must be made available to DDSD	
upon request, upon the termination or expiration of a	
provider agreement, or upon provider withdrawal	
from services.	
20.5.3 Health Passport and Physician	
Consultation Form: All Primary and Secondary	
Provider Agencies must use the <i>Health Passport</i> and	
The real of Agencies must use the meaning assport and	

Physician Consultation form from the Therap	
system. This standardized document contains	
individual, physician and emergency contact	
information, a complete list of current medical	
diagnoses, health and safety risk factors, allergies,	
and information regarding insurance, guardianship,	
and advance directives. The Health Passport also	
includes a standardized form to use at medical	
appointments called the <i>Physician Consultation</i>	
form. The <i>Physician Consultation</i> form contains a list	
of all current medications.	
Chapter 10: Living Care Arrangements (LCA)	
Living Supports-Supported Living: 10.3.9.6.1	
Monitoring and Supervision	
4. Ensure and document the following:	
a. The person has a Primary Care Practitioner.	
b. The person receives an annual physical	
examination and other examinations as	
recommended by a Primary Care	
Practitioner or specialist.	
c. The person receives annual	
dental check-ups and other	
check-ups as recommended by	
a licensed dentist.	
 d. The person receives a hearing test as recommended by a licensed audiologist. 	
e. The person receives eye	
examinations as recommended	
by a licensed optometrist or	
ophthalmologist.	
5. Agency activities occur as required for follow-up	
activities to medical appointments (e.g. treatment,	
visits to specialists, and changes in medication or	
daily routine).	
10.3.10.1 Living Care Arrangements (LCA)	
Living Supports-IMLS: 10.3.10.2 General	
Requirements: 9 . Medical services must be	
ensured (i.e., ensure each person has a licensed	
Primary Care Practitioner and receives an annual	
physical examination, specialty medical care as	
needed, and annual dental checkup by a licensed	
needed, and annual dental checkup by a licensed	

dentist). Chapter 13 Nursing Services: 13.2.3 General Requirements: 1. Each person has a licensed primary care practitioner and receives an annual physical examination and specialty medical/dental care as needed. Nurses communicate with these providers to share current health information.

Tag # 1A09 Medication Delivery Routine	Condition of Participation Level Deficiency	Condition of Participation Level Deficiency
Medication Administration Developmental Disabilities (DD) Waiver Service	After an analysis of the evidence, it has been	New / Repeat Finding:
Standards 2/26/2018; Re-Issue: 12/28/2018; Eff	determined there is a significant potential for a	
1/1/2019	negative outcome to occur.	After an analysis of the evidence, it has been
Chapter 20: Provider Documentation and Client		determined there is a significant potential for a
Records 20.6 Medication Administration Record	Medication Administration Records (MAR) were	negative outcome to occur.
(MAR): A current Medication Administration Record	reviewed for the months of April and May 2022.	
(MAR) must be maintained in all settings where		Medication Administration Records (MAR) were
medications or treatments are delivered. Family	Based on record review, 2 of 4 individuals had	reviewed for the months of October 2022.
Living Providers may opt not to use MARs if they are	Medication Administration Records (MAR), which	
the sole provider who supports the person with	contained missing medications entries and/or other	Based on record review, 1 of 4 individuals had
medications or treatments. However, if there are	errors:	Medication Administration Records (MAR), which
services provided by unrelated DSP, ANS for		contained missing medications entries and/or other
Medication Oversight must be budgeted, and a MAR	Individual #3	errors:
must be created and used by the DSP.	April 2022	
Primary and Secondary Provider Agencies are	Physician's Orders indicated the following	Individual #5
responsible for:	medication were to be given. The following	October 2022
1. Creating and maintaining either an	Medications were not documented on the	As indicated by the Medication Administration
electronic or paper MAR in their service	Medication Administration Records:	Records the individual is to take Docusate Sodium
setting. Provider Agencies may use the MAR	 Melatonin 5mg (1 time daily) 	100mg (1 time daily). According to the Physician's
in Therap, but are not mandated to do so.		Orders, Docusate Sodium 100mg (as needed).
2. Continually communicating any changes	Individual #5	Medication Administration Record and Physician's
about medications and treatments between	April 2022	Orders do not match.
Provider Agencies to assure health and safety.	Medication Administration Records contained	As indicated by the Madiastian Administration
7. Including the following on the MAR:	missing entries. No documentation found	As indicated by the Medication Administration Records the individual is to take Miralax Powder
a. The name of the person, a transcription of the physician's or licensed health care	indicating reason for missing entries:	17gram (1 time daily). According to the Physician's
provider's orders including the brand and	Mupirocin 2% Ointment (2 times daily) – Blank (46 (8 AN)) and 4(40, 26 (8 DN))	Orders, Polyethylene Glycol 17gm (as needed).
generic names for all ordered routine and	4/16 (8 AM), and 4/19, 26 (8PM)	Medication Administration Record and Physician's
PRN medications or treatments, and the	Medication Administration Records contain the	Orders do not match.
diagnoses for which the medications or	following medications. No Physician's Orders were	
treatments are prescribed;	found for the following medications:	Medication Administration Records contain the
b. The prescribed dosage, frequency and	Ergocalciferol 50,000 IU (1 time monthly)	following medications. No Physician's Orders were
method or route of administration; times	• Ergocalcheror 50,000 TO (T time monthly)	found for the following medications:
and dates of administration for all ordered	Saline Nasal Gel / Spray (2 times daily)	• Creams (1 time daily)
routine or PRN prescriptions or treatments;	• Same Nasar Ger/ Spray (2 times dairy)	
over the counter (OTC) or "comfort"	Mupirocin 2% Ointment (2 times 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	
<i>D. Administration of Drugs</i> 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.	
ponodi	

Standard of Care	Routine Survey Deficiencies May 16 – 26, 2022	Verification Survey New and Repeat Deficiencies November 28 – December 8, 2022
Service Domain: Service Plans: ISP Implementation		
frequency specified in the service plan.		
Tag # 1A08 Administrative Case File (Other	Standard Level Deficiency	COMPLETE
Required Documents)	•	
Tag # 1A32.1 Administrative Case File: Individual	Standard Level Deficiency	COMPLETE
Service Plan Implementation (Not Completed at	-	
Frequency)		
Tag # 1A32.2 Individual Service Plan	Standard Level Deficiency	COMPLETE
Implementation (Residential Implementation)		
Tag # LS14 Residential Service Delivery Site	Condition of Participation Level Deficiency	COMPLETE
Case File (ISP and Healthcare Requirements)		
Tag # LS14.1 Residential Service Delivery Site	Standard Level Deficiency	COMPLETE
Case File (Other Req. Documentation)		
Service Domain: Qualified Providers - The State me		
implements its policies and procedures for verifying the		
Tag # 1A22 Agency Personnel Competency	Standard Level Deficiency	COMPLETE
Tag # 1A26 Consolidated On-line Registry	Standard Level Deficiency	COMPLETE
Employee Abuse Registry	•	
Tag # 1A37 Individual Specific Training	Standard Level Deficiency	COMPLETE
Tag # 1A43.1 General Events Reporting: Individual Reporting	Condition of Participation Level Deficiency	COMPLETE
	an ongoing basis identifies addresses and seeks to	prevent occurrences of abuse, neglect and exploitation.
Individuals shall be afforded their basic human rights.		
Tag # 1A09.1 Medication Delivery PRN	Condition of Participation Level Deficiency	COMPLETE
Medication Administration		
Tag # 1A09.1.0 Medication Delivery	Standard Level Deficiency	COMPLETE
PRN Medication Administration		
Tag # 1A09.2 Medication Delivery Nurse	Condition of Participation Level Deficiency	COMPLETE
Approval for PRN Medication	······································	
Tag # 1A15.2 Administrative Case File:	Condition of Participation Level Deficiency	COMPLETE
Healthcare Documentation (Therap and Required		
Plans)		
Tag # 1A27.2 Duty to Report IRs Filed During On-	Standard Level Deficiency	COMPLETE
Site and/or IRs Not Reported by Provider	•	
Service Domain: Medicaid Billing/Reimbursement	 State financial oversight exists to assure that claims 	s are coded and paid for in accordance with the
reimbursement methodology specified in the approved	waiver.	
Tag # IS30 Customized Community Supports	Standard Level Deficiency	COMPLETE
Reimbursement	•	

	Verification Survey Plan of Correction, On-going QA/QI and Responsible Party	Completion Due
Tag # 1A08.2 Administrative Case File: Healthcare Requirements & Follow-up	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?): \rightarrow	
	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?): \rightarrow	
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?): \rightarrow	
	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?): \rightarrow	



PATRICK M. ALLEN Cabinet Secretary Designate

Date:	January 27, 2023
То:	Claudine Valerio-Salazar, Executive Director
Provider: Address: State/Zip:	EnSuenos Y Los Angelitos Development Center 1030 Salazar Rd. Taos, New Mexico 87571
E-mail Address:	cvs@eladc.org
CC: E-Mail Address:	Analisa Rugelio, Supported Living Coordinator / QI Coordinator / Trainer avigil@eladc.org
Region: Routine Survey: Verification Survey:	Northeast May 16 - 26, 2022 November 28 – December 8, 2022
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	Supported Living, Customized Community Supports, and Community Integrated Employment Services
Survey Type:	Verification

Dear Ms. Valerio-Salazar,

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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