Date:	September 9, 2025
To:	Julian Sandoval, Administrative Assistant
Provider: Address: State/Zip:	Trinity Asset Management, LLC 108 Roehl Road NW Albuquerque, New Mexico 87107
E-mail Address:	admin@tamsupport.net
CC: E-Mail Address:	Melissa Morenus, Director melissa@tamsupport.net
CC: E-Mail Address:	McKenzie Chiarelli, Chief Operating Officer mckenzie@tamsupport.net
Region: Survey Date:	Metro August 11 - 21, 2025
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	Supported Living, Customized Community Supports
Survey Type:	Initial
Team Leader:	Monica Valdez, BS, Advanced Healthcare Surveyor / Plan of Correction Coordinator, Division of Health Improvement/Quality Management Bureau
Team Members:	Armida Medina, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Dear Mr. Sandoval:

The Division of Health Improvement/Quality Management Bureau has completed a compliance survey of the services identified above. The purpose of the survey was to determine compliance with federal and state standards; to assure the health, safety, and welfare of individuals receiving services through the Developmental Disabilities Waiver; and to identify opportunities for improvement. This Report of Findings will be shared with the Developmental Disabilities Supports Division for their use in determining your current and future provider agreements. Upon receipt of this letter and Report of Findings your agency must immediately correct all deficiencies which place Individuals served at risk of harm.

Determination of Compliance:
The Division of Health Improvement, Quality Management Bureau has determined your agency is 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 # 1A22 Agency Personnel Competency
- Tag# 1A08.2 Administrative Case File: Healthcare Requirements & Follow-up
- Tag # 1A09.1 Medication Delivery PRN Medication Administration

The following tags are identified as Standard Level:

- Tag # 1A20 Direct Support Professional Training
- Tag # 1A26 Employee Abuse Registry
- Tag # 1A33 Board of Pharmacy: Med. Storage

Plan of Correction:

The attached Report of Findings identifies the deficiencies found during your agency's on-site compliance review. You are required to complete and implement a Plan of Correction. Your agency has a total of 45 business days (10 business days to submit your POC for approval and 35 days to implement your *approved* Plan of Correction) from the receipt of this letter.

You were provided information during the exit meeting portion of your on-site survey. Please refer to this information (Attachment A) for specific instruction on completing your Plan of Correction. At a minimum your Plan of Correction should address the following for each Tag cited:

Corrective Action for Current Citation:

- How is the deficiency going to be corrected? (i.e. obtained documents, retrain staff, individuals and/or staff no longer in service, void/adjusts completed, etc.) This can be specific to each deficiency cited or if possible an overall correction, i.e. all documents will be requested and filed as appropriate.

On-going Quality Assurance/Quality Improvement Processes:

- What is going to be done on an ongoing basis? (i.e. file reviews, etc.)
- How many individuals is this going to effect? (i.e. percentage of individuals reviewed, number of files reviewed, etc.)
- How often will this be completed? (i.e. weekly, monthly, quarterly, etc.)
- Who is responsible? (responsible position within your agency)
- What steps will be taken if issues are found? (i.e. retraining, requesting documents, filing RORA, etc.)
- How is this integrated in your agency's QIS, QI Committee reviews and annual report?

Submission of your Plan of Correction:

Please submit your agency's Plan of Correction in the available space on the two right-hand columns of the Report of Findings. *(See attachment "A" for additional guidance in completing the Plan of Correction)*.

Within 10 business days of receipt of this letter your agency Plan of Correction must be submitted to the parties below:

1. **Quality Management Bureau, Verna Newman-Sikes, Healthcare Surveyor Advanced / Plan of Correction Coordinator** at verna.newman-sikes@hca.nm.gov

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

Upon notification from QMB that your *Plan of Correction has been approved*, you must implement all remedies and corrective actions to come into compliance. If your Plan of Correction is denied, you must resubmit a revised plan as soon as possible for approval, as your POC approval and all remedies must be completed within 45 business days of the receipt of this letter.

Failure to submit your POC within the allotted 10 business days or complete and implement your Plan of Correction within the total 45 business days allowed may result in the imposition of a $200 per day Civil Monetary Penalty until it is received, completed and/or implemented.

Billing Deficiencies:
If you have deficiencies noted in this report of findings under the *Service Domain: Medicaid Billing/Reimbursement*, you must complete a "Void/Adjust" claim or remit the identified overpayment via a check within 30 calendar days of the date of this letter to HCA/OIG/PIU, *though this is not the preferred method of payment*. If you choose to pay via check, please include a copy of this letter with the payment. Make the check payable to the New Mexico Human Services Department and mail to:

Attention: *Lisa Medina-Lujan*
HCA/OIG/Program Integrity Unit
PO Box 2348
1474 Rodeo Road
Santa Fe, New Mexico 87505

If you have questions and would like to speak with someone at HCA/OIG/PIU, please contact:

Lisa Medina-Lujan (Lisa.Medina-Lujan@hca.nm.gov)

Please be advised that there is a one-week lag period for applying payments received by check to Void/Adjust claims. During this lag period, your other claim payments may be applied to the amount you owe even though you have sent a refund, reducing your payment amount. For this reason, we recommend that you allow the system to recover the overpayment instead of sending in a check.

Request for Informal Reconsideration of Findings (IRF):
If you disagree with a finding of deficient practice, you have 10 business days upon receipt of this notice to request an IRF. Submit your request for an IRF in writing to:

ATTN: QMB Bureau Chief
Request for Informal Reconsideration of Findings
5300 Homestead Rd NE, Suite 200-211
Albuquerque, NM 87110
Attention: IRF request/QMB

See Attachment "C" for additional guidance in completing the request for Informal Reconsideration of Findings. The request for an IRF will not delay the implementation of your Plan of Correction which must be completed within 45 total business days (10 business days to submit your POC for approval and 35 days to implement your *approved* Plan of Correction). Providers may not appeal the nature or interpretation of the standard or regulation, the team composition or sampling methodology. If the IRF approves the modification or removal of a finding, you will be advised of any changes.

Please contact the Plan of Correction Coordinator, Verna Newman-Sikes at 575-449-5843 or email at: verna.newman-sikes@hca.nm.gov 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,

Monica Valdez, BS

Monica Valdez, BS
Team Lead/Healthcare Surveyor
Division of Health Improvement
Quality Management Bureau

Survey Process Employed:

Administrative Review Start Date:	August 11, 2025
Contact:	**Trinity Asset Management, LLC** Julian Sandoval, Administrative Assistant **HCA/DHI/QMB** Monica Valez, BS, Team Lead/ Advanced Healthcare Surveyor/Plan of Correction Coordinator
Entrance Conference Date:	August 11, 2025
Present:	**Trinity Asset Management, LLC** Julian Sandoval, Administrative Assistant McKenzie Chiarelli, Chief Operating Officer **HCA/DHI/QMB** Monica Valez, BS, Team Lead/ Advanced Healthcare Surveyor/Plan of Correction Coordinator Armida Medina, Healthcare Surveyor Jamie Pond, BS, Staff Manager
Exit Conference Date:	August 21, 2025
Present:	**Trinity Asset Management, LLC** Melissa Morenus, Director Julian Sandoval, Administrative Assistant McKenzie Chiarelli, Chief Operating Officer **HCA/DHI/QMB** Monica Valez, BS, Team Lead/ Advanced Healthcare Surveyor/Plan of Correction Coordinator Armida Medina, Healthcare Surveyor Jamie Pond, BS, Staff Manager **DDSD – Metro Regional Office** Katie Hellebust, Provider Support Coordinator
Administrative Locations Visited:	0 (*Administrative portion of survey completed remotely*)
Total Wellness Visits Completed (Individuals Seen):	4
Total Compliance Survey Sample Size:	4 4 - Supported Living 3 - Customized Community Supports
Total Compliance Survey Homes Visits	1
❖ Supported Living Homes Visited	1 *Note: The following Individuals share a SL residence:* • #1, 2, 3, 4

Persons Served Records Reviewed	4
Persons Served Interviewed	1
Persons Served Observed	3 *(Note: 3 Individuals were observed, as one individual was engaged in another different activity and two were not responsive to surveyors' questions)*
Direct Support Professional Records Reviewed	13
Direct Support Professional Interviewed	1
Service Coordinator Records Reviewed	1
Nurse Interview	2

Administrative Processes and Records Reviewed:

- Medicaid Billing/Reimbursement Records for all Services Provided
- Oversight of Individual Funds
- Individual Agency / Residential / Site Case Files, including, but not limited to:
    - Individual Service Plans
    - Progress on Identified Outcomes
    - Healthcare Plans
    - Medication Administration Records
    - Physician Orders
    - Therapy Plans
    - Healthcare Documentation Regarding Appointments and Required Follow-Up
    - Other Required Health Information / Therap Required Documents
- Internal Incident Management Reports and System Process / General Events Reports
- Personnel Files:
    - Training Records
    - Caregiver Criminal History Screening Records
    - Employee Abuse Registry
- Interviews with the Individuals and Agency Personnel
- Human Rights Committee Notes and Meeting Minutes
- Quality Assurance / Improvement Plan
- Agency Policy and Procedure Manual

CC: Distribution List: HCA - Division of Health Improvement
HCA - Developmental Disabilities Supports Division
HCA - Medical Assistance Division

Attachment A

Provider Instructions for Completing the QMB Plan of Correction (POC) Process

Introduction:
After a QMB Compliance Survey, your QMB Report of Findings will be sent to you via e-mail.

Each provider must develop and implement a Plan of Correction (POC) that identifies specific quality assurance and quality improvement activities the agency will implement to correct deficiencies and prevent continued deficiencies and non-compliance.

Agencies must submit their Plan of Correction within ten (10) business days from the date you receive the QMB Report of Findings. (Providers who do not submit a POC within 10 business days may be referred to the DDSD Regional Office for purposes of contract management or the Internal Review Committee [IRC] for possible actions or sanctions).

Agencies must fully implement their approved Plan of Correction within 45 business days (10 business days to submit your POC for approval and 35 days to implement your approved Plan of Correction) from the date they receive the QMB Report of Findings. Providers who fail to complete a POC within the 45-business days allowed will be referred to the IRC for possible actions or sanctions.

Requests for technical assistance must be requested through your Regional DDSD Office.

The POC process cannot resolve disputes regarding findings. If you wish to dispute a finding on the official Report of Findings, you must file an Informal Reconsideration of Findings (IRF) request within ten (10) business days of receiving your report. Please note that you must still submit a POC for findings that are in question (see Attachment C).

Instructions for Completing Agency POC:

Required Content
Your Plan of Correction should provide a step-by-step description of the methods to correct each deficient practice cited to prevent recurrence and information that ensures the regulation cited comes into and remains in compliance. The remedies noted in your POC are expected to be added to your Agency's required, annual Quality Assurance (QA) Plan.

If a deficiency has already been corrected since the on-site survey, the plan should state how it was corrected, the completion date (date the correction was accomplished), and how possible recurrence of the deficiency will be prevented.

The following details should be considered when developing your Plan of Correction:

The Plan of Correction must address each deficiency cited in the Report of Findings unless otherwise noted with a "No Plan of Correction Required statement." The Plan of Correction must address the five (5) areas listed below:

1. How the specific and realistic corrective action will be accomplished for individuals found to have been affected by the deficient practice.
2. How the agency will identify other individuals who have the potential to be affected by the same deficient practice, and how the agency will act to protect those individuals in similar situations.
3. What Quality Assurance measures will be put into place and what systemic changes made to ensure the deficient practice will not recur.
4. Indicate how the agency plans to monitor its performance to make certain solutions are sustained. The agency must develop a QA plan for ensuring correction is achieved and sustained. This QA plan must be implemented, and the corrective action is evaluated for its effectiveness. The plan of correction is integrated into the agency quality assurance system; and

5. Include dates when corrective actions will be completed. The corrective action completion dates must be acceptable to the State.

The following details should be considered when developing your Plan of Correction:

- Details about how and when Individual Served, agency personnel and administrative and service delivery site files are audited by agency personnel to ensure they contain required documents;
- Information about how medication administration records are reviewed to verify they contain all required information before they are distributed to service sites, as they are being used, and after they are completed;
- Your processes for ensuring that all required agency personnel are trained on required DDSD required trainings;
- How accuracy in billing/reimbursement documentation is assured;
- How health, safety is assured;
- For Case Management providers, how Individual Service Plans are reviewed to verify they meet requirements, how the timeliness of level of care (LOC) packet submissions and consumer visits are tracked;
- Your process for gathering, analyzing and responding to quality data indicators; and,
- Details about Quality Targets in various areas, current status, analyses about why targets were not met, and remedies implemented.

Note: **Instruction or in-service of staff alone may not be a sufficient plan of correction.** This is a good first step toward correction, but additional steps must be taken to ensure the deficiency is corrected and will not recur.

Completion Dates

- The plan of correction must include a **completion date** (entered in the far right-hand column) for each finding. Be sure the date is **realistic** in the amount of time your Agency will need to correct the deficiency; not to exceed 45 total business days.
- Direct care issues should be corrected immediately and monitored appropriately.
- Some deficiencies may require a staged plan to accomplish total correction.
- Deficiencies requiring replacement of equipment, etc., may require more time to accomplish correction but should show reasonable time frames.

Initial Submission of the Plan of Correction Requirements

1. The Plan of Correction must be completed on the official QMB Survey Report of Findings/Plan of Correction Form and received by QMB within ten (10) business days from the date you received the report of findings.
2. For questions about the POC process, call the POC Coordinator, Verna Newman-Sikes at 575-449-5843 or email at: verna.newman-sikes@hca.nm.gov for assistance.
3. For Technical Assistance (TA) in developing or implementing your POC, contact your respective Regional DDSD Office.
4. Submit your POC to via email to verna.newman-sikes@hca.nm.gov. Please also submit your POC to your Developmental Disabilities Supports Division Regional Office for region of service surveyed.
5. ***Do not submit supporting documentation*** (evidence of compliance) to QMB ***until after*** **your POC has been approved** by the QMB.
6. QMB will notify you when your POC has been "approved" or "denied."
    a. During this time, whether your POC is "approved," or "denied," you will have a maximum of 45-business days from the date of receipt of your Report of Findings to correct all survey deficiencies.
    b. If your POC is denied, it must be revised and resubmitted as soon as possible, as the 45-business day limit is in effect.
    c. If your POC is denied a second time your agency may be referred to the Internal Review Committee.
    d. You will receive written confirmation when your POC has been approved by QMB and a final deadline for completion of your POC.
    e. Please note that all POC correspondence will be sent electronically unless otherwise requested.
7. Failure to submit your POC within 10 business days without prior approval of an extension by QMB will result in a referral to the Internal Review Committee and the possible implementation of monetary penalties and/or sanctions.

POC Document Submission Requirements

Once your POC has been approved by the QMB Plan of Correction Coordinator, you must submit copies of documents as evidence that all deficiencies have been corrected. You must also submit evidence of the ongoing Quality Assurance/Quality Improvement processes.

1. Your internal documents are due within a *maximum* of 45-business days of receipt of your Report of Findings.
2. Please submit your documents electronically according to the following: If documents do not contain protected Health information (PHI) then you may submit your documents electronically scanned and attached to the State email account. If documents contain PHI **do not** submit PHI directly to the State email account. *You may submit PHI **only** when **replying** to a **secure** email received from the State email account*. When possible, please submit requested documentation using a "zipped/compressed" file to reduce file size. You may also submit documents via S-Comm (Therap), or another electronic format, i.e., flash drive.
3. All submitted documents *must be annotated*; please be sure the tag numbers and Identification numbers are indicated on each document submitted. Documents which are not annotated with the Tag number and Identification number may not be accepted.
4. Do not submit original documents; Please provide copies or scanned electronic files for evidence. Originals must be maintained in the agency file(s) per DDSD Standards.
5. In lieu of some documents, you may submit copies of file or home audit forms that clearly indicate cited deficiencies have been corrected, other attestations of correction must be approved by the Plan of Correction Coordinator prior to their submission.
6. When billing deficiencies are cited, you must provide documentation to justify billing and/or void and adjust forms submitted to Conduent for the deficiencies cited in the Report of Findings.

Revisions, Modifications or Extensions to your Plan of Correction (post QMB approval) must be made in writing and submitted to the Plan of Correction Coordinator, prior to the due date and are approved on a case-by-case basis. No changes may be made to your POC or the timeframes for implementation without written approval of the POC Coordinator.

Attachment B

Division of Health Improvement / Quality Management Bureau
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:

Service Domain: Service Plan: ISP Implementation - *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)

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.*

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
- **1A09.2 –** Medication Delivery Nurse Approval for PRN Medication
- **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
- **1A15 –** Healthcare Documentation - Nurse Availability
- **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: Microsoft Word - IRF-QMB-Form.doc The written request for an IRF must specify in detail the request for reconsideration and why the finding is inaccurate.
3. 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@hca.nm.gov 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 standard level tags, as well as, 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 two 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: **Trinity Asset Management, LLC – Metro Region**
Program: Developmental Disabilities Waiver
Service: Supported Living, Customized Community Supports
Survey Type: Initial
Survey Date: **August 11 – 21, 2025**

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Completion Date
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 # 1A20 Direct Support Professional Training	**Standard Level Deficiency**		
Developmental Disabilities Waiver Service Standards Orig. Issue:11/1/2021 Reissue eff. 7.1.2024 **Chapter 17 Training Requirements: 17.1 Training Requirements for Direct Support Professional and Direct Support Supervisors:** Direct Support Professional (DSP) and Direct Support Supervisors (DSS) include staff and contractors from agencies providing the following services: Supported Living, Family Living, CIHS, IMLS, CCS, CIE and Crisis Supports… The training shall address at least the following: • Individual Specific Training • First Aid • CPR • Assisting With Medication Delivery (AWMD) Part 1 Session 1 & 2 **…** **17.1.13 Training Requirements for Service Coordinators (SC):** Service Coordinators (SCs) refer to staff at agencies providing the following services: Supported Living, Family Living, Customized In-home Supports, Intensive Medical Living, Customized Community Supports, Community Integrated Employment, and Crisis Supports…The training shall address at least the following: • Individual Specific Training	Based on record review, the Agency did not ensure Orientation and Training requirements were met for 1 of 14 Direct Support Professional, Direct Support Supervisory Personnel and / or Service Coordinators. Review of Agency training records found no evidence of the following required HCA/DDSD trainings being completed: **First Aid:** • Not Found (#507) **CPR:** • Not Found (#507)	**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?)*: →	

• First Aid • CPR • Assisting With Medication Delivery (AWMD) Part 1 Session 1 & 2 … ***(see DDW Standards Chapter 17 Training Requirements for all training specifics)***			

Tag # 1A22 Agency Personnel Competency	Condition of Participation Level Deficiency		
Developmental Disabilities Waiver Service Standards Orig. Issue:11/1/2021 Reissue eff. 7.1.2024 **Chapter 17 Training Requirements:** **17.9 Individual-Specific Training Requirements:** 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 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. The trainer must 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	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 1 of 1 Direct Support Professional. **When DSP were asked, if the Individual's had Health Care Plans, where could they be located and if they had been trained, the following was reported:** • DSP #504 stated, "Body mass, skin and wound, BM's, make sure he's hydrated, constipation, paralysis." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual also requires Health Care Plans for Aspiration and Pain. (Individual #1) • DSP #504 stated, "CARMP. I do not see any in here for him." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual also requires a Health Care Plan for Gastroesophageal Reflux Disease. (Individual #2)	**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?)*: →	

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…			

Tag # 1A26 Employee Abuse Registry	Standard Level Deficiency		
NMAC 8.370.8 - EMPLOYEE ABUSE REGISTRY: This rule applies to a broad range of New Mexico providers of health care and services and employees of these providers who are not licensed health care professionals or certified nurse aides. This rule requires that providers check with the registry and avoid employing an individual on the registry. This rule provides for the investigation and determination of complaints alleging abuse, neglect or exploitation of recipients of care or services by employees. This rule further requires listing employees with substantiated registry-referred abuse, neglect or exploitation on the registry, following an opportunity for a hearing. This rule supplements other pre-employment screening requirements currently applicable to health care providers, such as the requirement for criminal history screening of caregivers employed by care providers subject to the Caregiver Criminal History Screening Act, Sections 29-17-1 et seq. NMSA 1978, and that Act's implementing rule, 8.370.5 NMAC. It also supplements requirements for pre-employment screening of certified nurse aides applicable to nursing facilities pursuant to 42 CFR Sections 483.75(e) and 488.335; and 8.370.25 NMAC. This rule does not address the consequences of abuse, neglect, or exploitation for which a provider, as distinguished from an employee, is responsible. **8.370.8.8 REGISTRY ESTABLISHED; PROVIDER INQUIRY REQUIRED:** Upon the effective date of this rule, the authority 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	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 4 of 14 Agency Personnel. **The following Agency Personnel records contained evidence that indicated the Employee Abuse Registry check was completed after hire:** **Direct Support Professional (DSP):** • #500 – Date of hire 3/21/2025, completed 4/8/2025. • #501 – Date of hire 3/26/2025, completed 4/8/2025. • #507 – Date of hire 6/7/2025, completed 6/10/2025. **Service Coordination Personnel (SC):** • #513 – Date of hire 3/18/2025, completed 4/8/2025.	**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?)*: →	

information of all persons who, while employed by a provider, have been determined by the authority, 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 business days following receipt. Only authority 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. **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.			

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Completion Date
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**		
Developmental Disabilities Waiver Service Standards Orig. Issue:11/1/2021 Reissue eff. 7.1.2024 **Chapter 3 Safeguards: 3.1 Decisions about Health Care or Other Treatment: Decision Consultation Process:** There are a variety of approaches and available resources to support decision making when desired by the person. The decision consultation process assists participants and their health care decision makers to document their decisions. It is important for provider agencies to communicate with guardians to share with the Interdisciplinary Team (IDT) Members any medical, behavioral, or psychiatric information as part of an individual's routine medical or psychiatric care. **3.1.1 Decision about Health Care or Other Treatment Decision Consultation:** Decisions are the sole domain of waiver participants; their guardians or healthcare decision makers and decisions can be made that are compatible with their personal and cultural values. Provider Agencies and Interdisciplinary Teams (IDTs) are required to support the informed decisions made by supporting access to medical consultation, information, and other available resources according to the following: The Decision Consultation Process (DCP) is documented on the Decision Consultation Form (DCF) and is used for recommendations when a person or his/her guardian/healthcare decision maker has concerns, needs more information, or has decided not to follow all or part of a	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 provide documentation of annual physical examinations and/or other examinations as specified by a licensed physician for 1 of 4 individuals receiving Living Care Arrangements and Community Inclusion. Review of the Agency administrative individual case files revealed the following items were not found, not current and/or did not meet the requirement: **Annual Dental Exam:** • Individual #2 - As indicated by collateral documentation reviewed, the exam was not found. Per the Appendix A Client File matrix, Dental Exams are to be conducted annually.	**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?)*: →	

recommendation from a professional or clinician… **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 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. Records must contain information of concerns related to abuse, neglect or exploitation. 3. 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 are acceptable. 4. Provider Agencies are responsible for ensuring that all plans created by nurses, RDs, therapists or BSCs are present in all settings. 5. 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.			

6. 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. 7. 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. 8. All records must be retained for six (6) years 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 Communication and Documentation in Therap:** Therap is a secure online documentation system required to be used by specific New Mexico DD Waiver Provider Agencies. Use of the required elements of Therap are intended to improve agency monitoring, health care coordination for individuals, and overall quality of services. **20.5.3 *Health Passport and Consultation Form:*** The Health Passport and Consultation form are generated within Therap. The standardized combination of documents includes all information that are required for medical consultation during an appointment and other health coordination activities: 1. The Primary Provider must keep the Health Passport and Consultation form updated in concert with critical information and changes from the IDT, including secondary provider agencies, medical providers for the individual. The Health Passport pulls from Individual Demographics, Health Tracking and eCHAT. a. The primary provider must notify secondary providers when a new eCHAT is completed or contact information is updated.			

2. The Primary and Secondary Provider Agencies must ensure that a current copy of the *Health Passport* and *Consultation* forms are printed and available at all service delivery sites. a. Updated forms must be sent to each site after eCHAT and/or Contact Updates. b. Outdated version of both unused forms must be removed from all sites. 3. Primary and Secondary Provider Agencies must assure that the current *Health Passport* and *Consultation* form accompany each person when taken by the provider to a medical appointment, urgent care/emergency room visits, emergency service encounter, or are admitted to a hospital or nursing home for details see Health Tracking: Appointments **20.5.4 *Health Tracking*** **20.5.5 *Nursing Assessment Tracking*** **Chapter 13 Nursing Services: 13.2.3 General Requirements Related to Orders, Implementation, and Oversight:** 1. Each person has a licensed primary care practitioner and receives an annual physical examination, dental care and specialized medical/behavioral care as needed. PPN communicate with providers regarding the person as needed. 2. Orders from licensed healthcare providers are implemented promptly and carried out until discontinued. a. The nurse will contact the ordering or on call practitioner as soon as possible if the order cannot be implemented due to the person's or guardian's refusal or due to other issues delaying implementation of the order. The nurse must clearly document the issues and all attempts to resolve the problems with all involved parties.			

b. Not implementing orders by a licensed healthcare provider is considered neglect, unless a Decision Consultation Form is filled out by participant or guardian, or a healthcare decision maker making this decision. c. Based on prudent nursing practice, if a nurse determines to hold a practitioner's order, they are required to immediately document the circumstances and rationale for this decision and to notify the ordering or on call practitioner as soon as possible, but no later than the next business day. d. If the person resides with their biological family, and there are no nursing services budgeted, the family is responsible for implementation or follow up on all orders from all providers.			

Tag # 1A09.1 Medication Delivery PRN Medication Administration	Condition of Participation Level Deficiency		
Developmental Disabilities Waiver Service Standards Orig. Issue:11/1/2021 Reissue eff. 7.1.2024 **Chapter 10 Living Care Arrangements (LCA): 10.3.5 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 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 5.7 Medication Administration Record (MAR) **Chapter 20 Provider Documentation and Client Records: 20.5.7 Medication Administration Record (MAR):** Administration of medications apply to all provider agencies of the following services: living supports, customized community supports, community integrated employment, intensive medical living supports. 1. Primary and secondary provider agencies are to utilize the Medication Administration Record (MAR) online in Therap. 2. Medication/Treatment must be recorded online per assisting with medication delivery per the DDSD Assisting with Medication Delivery (AWMD) program. 3. Family Living Providers may opt not to use MARs if they are the **sole** provider who supports the person and are related by affinity or consanguinity. However, if there are services provided by unrelated DSP, ANS for Medication Oversight must be budgeted, a	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 July 2025. Based on record review, 2 of 4 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #1 July 2025 Physician's Orders indicated the following medication were to be given. The following Medications were not documented on the Medication Administration Records: • Anbesol Max Liquid 20% (PRN) • Antacid/Anti-Gas - Susp (PRN) • Artificial Tears (PRN) Individual #4 July 2025 Medication Administration Records contain the following medications. No Physician's Orders were found for the following medications: • Acetaminophen 325 mg (PRN)	**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?)*: →	

MAR online in Therap must be created and used by the DSP. 4. Provider Agencies must configure and use the MAR when assisting with medication. 5. Provider Agencies Continually communicate any changes about medications and treatments between Provider Agencies to assure health and safety. 6. Provider agencies must include 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 and PRN medications and other treatments; all over the counter (OTC) or "comfort" medications or treatments; all self-selected herbal preparation approved by the prescriber, and/or vitamin therapy approved by prescriber. c. Documentation of all time limited or discontinued medications or treatments. d. The initials of the person administering or assisting with medication delivery. e. Documentation of refused, missed, or held medications or treatments. f. Documentation of any allergic reaction that occurred due to medication or treatments. g. For PRN medications or treatments including all physician approved over the counter medications and herbal or other supplements: 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 follow-up detailed documentation that the DSP contacted the agency nurse or physician service prior to assisting with the medication or treatment; and iii. documentation of the effectiveness of the PRN medication or treatment. **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 # 1A33 Board of Pharmacy: Med. Storage	Standard Level Deficiency		
New Mexico Board of Pharmacy Model Custodial Drug Procedures Manual **E. Medication Storage:** 1. Prescription drugs will be stored in a locked cabinet and the key will be in the care of the administrator or designee. 2. Drugs to be taken by mouth will be separate from all other dosage forms. 3. A locked compartment will be available in the refrigerator for those items labeled "Keep in Refrigerator." The temperature will be kept in the 36°F - 46°F range. An accurate thermometer will be kept in the refrigerator to verify temperature. 4. Separate compartments are required for each resident's medication. 5. All medication will be stored according to their individual requirement or in the absence of temperature and humidity requirements, controlled room temperature (68-77°F) and protected from light. Storage requirements are in effect 24 hours a day. 6. Medication no longer in use, unwanted, outdated, or adulterated will be placed in a quarantine area in the locked medication cabinet and held for destruction by the consultant pharmacist. **8. References** A. Adequate drug references shall be available for facility staff **H. Controlled Substances (Perpetual Count Requirement)** 1. Separate accountability or proof-of-use sheets shall be maintained, for each controlled substance, indicating the following information: a. date b. time administered	Based on observation, the Agency did not ensure proper storage of medication for 1 of 4 individuals. Observation included: Individual #2 • Hydroxyzine HCL 25 mg - Is no longer in use according to documentation found and not kept in a separate place, as required by regulation.	**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?)*: →	

c. name of patient d. dose e. practitioner's name f. signature of person administering or assisting with the administration the dose g. balance of controlled substance remaining. **NMAC 16.19.11 DRUG CONTROL** **(a)** All state and federal laws relating to storage, administration and disposal of controlled substances and dangerous drugs shall be complied with. **(b)** Separate sheets shall be maintained for controlled substances records indicating the following information for each type and strength of controlled substances: date, time administered, name of patient, dose, physician's name, signature of person administering dose, and balance of controlled substance in the container. **(c)** All drugs shall be stored in locked cabinets, locked drug rooms, or state of the art locked medication carts. **(d)** Medication requiring refrigeration shall be kept in a secure locked area of the refrigerator or in the locked drug room. **(e)** All refrigerated medications will be kept in separate refrigerator or compartment from food items. **(f)** Medications for each patient shall be kept and stored in their originally received containers, and stored in separate compartments. Transfer between containers is forbidden, waiver shall be allowed for oversize containers and controlled substances at the discretion of the drug inspector. **(g)** Prescription medications for external use shall be kept in a locked cabinet separate from other medications. **(h)** No drug samples shall be stocked in the licensed facility. **(i)** All drugs shall be properly labeled with the following information:			

(i) Patient's full name; **(ii)** Physician's name; **(iii)** Name, address and phone number of pharmacy; **(iv)** Prescription number; **(v)** Name of the drug and quantity; **(vi)** Strength of drug and quantity; **(vii)** Directions for use, route of administration; **(viii)** Date of prescription (date of refill in case of a prescription renewal); **(ix)** Expiration date where applicable: The dispenser shall place on the label a suitable beyond-use date to limit the patient's use of the medication. Such beyond-use date shall be not later than (a) the expiration date on the manufacturer's container, or (b) one year from the date the drug is dispensed, whichever is earlier; **(x)** Auxiliary labels where applicable; **(xi)** The Manufacturer's name; **(xii)** State of the art drug delivery systems using unit of use packaging require items i and ii above, provided that any additional information is readily available at the nursing station. Developmental Disabilities Waiver Service Standards Orig. Issue:11/1/2021 Reissue eff. 7.1.2024 **Chapter 10 Living Care Arrangement (LCA):** **10.3.7 Requirements for Each Residence:** Provider Agencies must assure that each residence is clean, safe, and comfortable, and each residence accommodates individual daily living, social and leisure activities. In addition, the Provider Agency must ensure the residence: … 8. has safe storage of all medications with dispensing instructions for each person that are consistent with the Assistance with Medication (AWMD) training or each person's ISP; …			

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Completion Date
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**		
NMAC 8.302.2 BILLING FOR MEDICAID SERVICES Developmental Disabilities Waiver Service Standards Orig. Issue:11/1/2021 Reissue eff. 7.1.2024 **Chapter 21: Billing Requirements; 23.1 Recording Keeping and Documentation Requirements:** DD Waiver Provider Agencies must maintain all records necessary to demonstrate proper provision of services for Medicaid billing. At a minimum, Provider Agencies must adhere to the following: 1. The level and type of service provided must be supported in the ISP and have an approved budget prior to service delivery and billing. 2. Comprehensive documentation of direct service delivery must include, at a minimum: a. the agency name; b. the name of the recipient of the service; c. the location of the service; d. the date of the service; e. the type of service; f. the start and end times of the service; g. the signature and title of each staff member who documents their time; and 3. Details of the services provided. A Provider Agency that receives payment for treatment, services, or goods must retain all medical and business records for a period of at least six years from the last payment date, until ongoing audits are settled, or until involvement of the state Attorney General is completed regarding settlement of any claim, whichever is longer…	Based on record review, the Agency maintained all the records necessary to fully disclose the nature, quality, amount and medical necessity of services furnished to an eligible recipient who is currently receiving DDW services for 4 of 4 individuals. *Progress notes and billing records supported billing activities for the months of April, May and June 2025 for the following services:* • Supported Living • Customized Community Supports		

21.4 Electronic Visit Verification: Section 12006(a) of the 21st Century Cures Act (the Cures Act) requires that states implement Electronic Visit Verification (EVV) for all Medicaid services under the umbrella of personal care and home health care that require an in-home visit by a provider. The EVV system verifies the: a. **Type** of service performed. b. **Individual receiving** the service. c. **Date** of service. d. **Location** of service delivery. e. **Individual providing** the service. f. **Time** the service begins and ends. **21.7 Billable Activities**: Specific billable activities are defined in the scope of work and service requirements for each DD Waiver service. In addition, any billable activity must also be consistent with the person's approved ISP. **21.9 Billable Units**: The unit of billing depends on the service type. The unit may be a 15-minute interval, a daily unit, a monthly unit, or a dollar amount. The unit of billing is identified in the current DD Waiver Rate Table. Provider Agencies must correctly report service units. **21.9.1 Requirements for Daily Units:** For services billed in daily units, Provider Agencies must adhere to the following: 1. A day is considered 24 hours from midnight to midnight. 2. If 12 or fewer hours of service are provided, then one-half unit shall be billed. A whole unit can be billed if more than 12 hours of service is provided during a 24-hour period. 3. The maximum allowable billable units cannot exceed 340 calendar days per ISP year or 170 calendar days per six months.			

21.9.2 Requirements for Monthly Units: For services billed in monthly units, a Provider Agency must adhere to the following: 1. A month is considered a period of 30 calendar days. 2. Face-to-face billable services shall be provided during a month where any portion of a monthly unit is billed. 3. Monthly units can be prorated by a half unit. **21.9.4 Requirements for 15-minute and hourly units:** For services billed in 15-minute or hourly intervals, Provider Agencies must adhere to the following: 1. When time spent providing the service is not exactly 15 minutes or one hour, Provider Agencies are responsible for reporting time correctly following NMAC 8.302.2. 2. Services that last in their entirety less than eight minutes cannot be billed.			

HEALTH CARE
AUTHORITY

Michelle Lujan Grisham, Governor
Kari Armijo, Secretary
Alex Castillo Smith, Deputy Secretary
Kathy Slater Huff, Deputy Secretary
Niki Kozlowski, Acting Deputy Secretary
Dana Flannery, Medicaid Director

Date:	October 14, 2025
To:	Jefferson Kee, Executive Director
Provider:	Tohatchi Area of Opportunity & Services, Inc.
Address:	1658 S. Second Street
State/Zip:	Gallup, New Mexico 87301
E-mail Address:	Jeff.Kee@taos-inc.org
CC:	Marcella Franklin, Quality Assurance Manager
E-Mail Address:	Marcella.Franklin@taos-inc.org
Region:	Northwest
Survey Date:	July 21 – August 1, 2025
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	Supported Living and Customized Community Supports
Survey Type:	Routine

Dear Mr. Kee and Ms. Franklin:

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,

Kayla R. Benally, BSW

Kayla R. Benally, BSW
Healthcare Surveyor Advanced/Plan of Correction Coordinator
DHI - Quality Management Bureau