NEW MEXICO
Department of Health
Division of Health Improvement

MICHELLE LUJAN GRISHAM
Governor

PATRICK M. ALLEN
Cabinet Secretary

Date:	October 2, 2023
To:	Kami Silva, Executive Director
Provider: Address: State/Zip:	Lessons of Life LLC 1720 S. Telshor Blvd Las Cruces, New Mexico 88011
E-mail Address:	ksilva@lessonsoflifellc.com
Region: Survey Date:	Southeast and Southwest August 21 – September 1, 2023
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	Supported Living, Family Living, Customized In-Home Supports; Customized Community Supports, and Community Integrated Employment Services
Survey Type:	Routine
Team Leader:	Verna Newman-Sikes, AA, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau
Team Members:	Kaydee Conticelli, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; William Easom, MPA, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; Kayla Hartsfield, BS, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; Sally Rel, MS, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; Amanda Castaneda-Holguin, MPA, Healthcare Surveyor Supervisor, Division of Health Improvement/Quality Management Bureau

Dear Ms. Silva:

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 # LS14 Residential Service Deliver Site Case File (ISP and Healthcare Requirements)
- Tag # 1A09 Medication Delivery Routine Medication Administration
- Tag # 1A09.1 Medication Delivery PRN Medication Administration
- Tag # 1A09.2 Medication Delivery Nurse Approval for PRN Medication
- Tag # 1A15.2 Administrative Case File: Healthcare Documentation (Therap and Required Plans)

The following tags are identified as Standard Level:

- Tag # 1A08.1 Administrative and Residential Case File: Progress notes
- Tag # 1A32 Administrative Case File: Individual Service Plan Implementation
- Tag # 1A32.1 Administrative Case File: Individual Service plan Implementation (Not Completed at Frequency)
- Tag # 1A32.2 Individual Service Plan Implementation (Residential Implementation)
- Tag # LS14.1 Residential Service Delivery Site Case File (Other Required Documentation)
- Tag # 1A20 Direct Support Professional Training
- Tag # 1A22 Agency Personnel Competency
- Tag # 1A26 Employee Abuse Registry
- Tag # 1A43.1 General Events Reporting: Individual Reporting
- Tag # 1A09.1.0 Medication Delivery PRN Medication Administration
- Tag # 1A27.2 Duty to Report IRs Filed During On-Site and/or IRs Not Reported by Provider
- Tag # 1A29 Complaints / Grievances Acknowledgement
- Tag # 1A39 Assistive Technology and Adaptive Equipment
- Tag # 1A50.1 Individual: Scope of Services *(Individual Interviews)*
- Tag # LS06 Family Living Requirements
- Tag # LS25 Residential Health & Safety (Supported Living / Family Living)
- Tag # IS30 Customized Community Supports Reimbursement
- Tag # LS27 Family Living Reimbursement

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, Monica Valdez, Plan of Correction Coordinator** at MonicaE.Valdez@doh.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 HSD/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*
HSD/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 HSD/OIG/PIU, please contact:

Lisa Medina-Lujan (Lisa.Medina-Lujan@hsd.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 300-331
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, Monica Valdez at 505-273-1930 or email at: MonicaE.Valdez@doh.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,

Verna Newman-Sikes, AA

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

Survey Process Employed:

Administrative Review Start Date:	August 21, 2023
Contact:	**Lessons of Life LLC** Kami Silva, Executive Director **DOH/DHI/QMB** Verna Newman-Sikes, AA, Team Lead/Healthcare Surveyor
On-site Entrance Conference Date:	*Entrance conference was waived by provider.*
Exit Conference Date:	September 1, 2023
Present:	**Lessons of Life LLC** Kami Silva, Executive Director **DOH/DHI/QMB** Verna Newman-Sikes, AA, Team Lead/Healthcare Surveyor William Easom, MPA, Healthcare Surveyor Kaydee Conticelli, Healthcare Surveyor Kayla Hartsfield, BS, Healthcare Surveyor Sally Rel, MS, Healthcare Surveyor Amanda Castaneda-Holguin MPA, Healthcare Surveyor Supervisor **DDSD – Metro Regional Office** Marie Velasco, DD Waiver Program Manager **DDSD – SW Regional Office** Jaime Lopez, Generalist
Total Sample Size:	26 2 - *Former Jackson Class Members* 24 - Non-*Jackson* Class Members 11 - Supported Living 10 - Family Living 3 - Customized In-Home Supports 15 - Customized Community Supports 4 - Community Integrated Employment
Total Homes Visits	20
❖ Supported Living Homes Visited	10 *Note: The following Individuals share a SL residence:* • #6, 15
❖ Family Living Homes Visited	10
Persons Served Records Reviewed	26
Persons Served Interviewed	22

Persons Served Observed	2 *(Note: 2 Individuals were observed, as one Individual chose not to participate in the interview process and one Individual was sleeping)*
Persons Served Not Seen and/or Not Available	2 *(Note: 2 Individuals were not available during the on-site survey)*
Direct Support Professional Records Reviewed	170
Direct Support Professional Interviewed	28
Substitute Care/Respite Personnel Records Reviewed	29
Service Coordinator Records Reviewed	6
Nurse Interview	1

Administrative Processes and Records Reviewed:

- Medicaid Billing/Reimbursement Records for all Services Provided
- Accreditation Records
- Oversight of Individual Funds
- Individual Medical and Program Case Files, including, but not limited to:
    - °Individual Service Plans
    - °Progress on Identified Outcomes
    - °Healthcare Plans
    - °Medical Emergency Response Plans
    - °Medication Administration Records
    - °Physician Orders
    - °Therapy Evaluations and Plans
    - °Healthcare Documentation Regarding Appointments and Required Follow-Up
    - °Other Required Health Information
- Internal Incident Management Reports and System Process / General Events Reports
- Personnel Files, including nursing and subcontracted staff
- Staff Training Records, Including Competency Interviews with Staff
- Agency Policy and Procedure Manual
- Caregiver Criminal History Screening Records
- Consolidated Online Registry/Employee Abuse Registry
- Human Rights Committee Notes and Meeting Minutes
- 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

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.

If you have questions about the Plan of Correction process, call the Plan of Correction Coordinator at 505-273-1930 or email at MonicaE.Valdez@doh.nm.gov. 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, Monica Valdez at 505-273-1930 or email at MonicaE.Valdez@doh.nm.gov for assistance.
3. For Technical Assistance (TA) in developing or implementing your POC, contact your Regional DDSD Office.
4. Submit your POC to Monica Valdez, POC Coordinator via email at MonicaE.valdez@doh.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 Xerox State Healthcare, LLC 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

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:

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 Professional Training
- **1A22 -** Agency Personnel Competency

- **1A37 –** Individual Specific Training

Non-Negotiable Condition of Participation Level Tags (one or more Individuals are cited):
- **1A25.1 –** Caregiver Criminal History Screening
- **1A26.1 –** Consolidated On-line Registry Employee Abuse Registry

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

Potential Condition of Participation Level Tags, if compliance is below 85%:
- **1A08.2 –** Administrative Case File: Healthcare Requirements & Follow-up
- **1A09 –** Medication Delivery Routine Medication Administration
- **1A09.1 –** Medication Delivery PRN Medication Administration
- **1A15.2 –** Administrative Case File: Healthcare Documentation (Therap and Required Plans)

Non-Negotiable Condition of Participation Level Tags (one or more Individuals are cited):
- **1A05 –** General Requirements / Agency Policy and Procedure Requirements
- **1A07 –** Social Security Income (SSI) Payments
- **1A09.2 –** Medication Delivery Nurse Approval for PRN Medication
- **1A15 –** Healthcare Coordination - Nurse Availability / Knowledge
- **1A31 –** Client Rights/Human Rights
- **LS25.1 –** Residential Reqts. (Physical Environment - Supported Living / Family Living / Intensive Medical Living)

Attachment C

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

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

Instructions:
1. The Informal Reconsideration of the Finding (IRF) request must be received in writing to the QMB Bureau Chief **within 10 business days** of receipt of the final Report of Findings **(*Note: No extensions are granted for the IRF)*.**
2. The written request for an IRF *must* be completed on the QMB Request for Informal Reconsideration of Finding form available on the QMB website: https://nmhealth.org/about/dhi/cbp/irf/
3. The written request for an IRF must specify in detail the request for reconsideration and why the finding is inaccurate.
4. The IRF request must include all supporting documentation or evidence.
5. If you have questions about the IRF process, email the IRF Chairperson, Valerie V. Valdez at valerie.valdez@doh.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 one to five (1 – 5) Condition of Participation Level Tags. This partial-compliance, if not corrected, may result in a serious negative outcome or the potential for more than minimal harm to individuals' health and safety.

Non-Compliance:

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

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

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

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

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Completion Date
Service Domain: Service Plans: ISP Implementation – *Services are delivered in accordance with the service plan, including type, scope, amount, duration and frequency specified in the service plan.*			
Tag # 1A08.1 Administrative and Residential Case File: Progress Notes	**Standard Level Deficiency**		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **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. 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. 3. Provider Agencies are responsible for ensuring that all plans created by nurses,	Based on record review, the Agency did not maintain progress notes and other service delivery documentation for 5 of 26 Individuals. Review of the Agency individual case files revealed the following items were not found: **Residential Case File:** **Supported Living Progress Notes/Daily Contact Logs:** • Individual #1 - None found for 8/1 – 10, 13 – 17, 20, 2023. *(Date of home visit: 8/22/2023)* **Family Living Progress Notes/Daily Contact Logs:** • Individual #9 - None found for 8/1 – 15, 2023. *(Date of home visit: 8/23/2023)* • Individual #13 - None found for 8/1 – 21, 2023. *(Date of home visit: 8/22/2023)* • Individual #25 - None found for 8/1 – 15, 2023. *(Date of home visit: 8/24/2023)* • Individual #28 - None found for 8/1 – 22, 2023. *(Date of home visit: 8/23/2023)*	**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?)***:** →	

RDs, therapists or BSCs are present in all 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.			

Tag # 1A32 Administrative Case File: Individual Service Plan Implementation	Standard Level Deficiency		
NMAC 7.26.5.16.C and D Development of the ISP. Implementation of the ISP. The ISP shall be implemented according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan. C. The IDT shall review and discuss information and recommendations with the individual, with the goal of supporting the individual in attaining desired outcomes. The IDT develops an ISP based upon the individual's personal vision statement, strengths, needs, interests and preferences. The ISP is a dynamic document, revised periodically, as needed, and amended to reflect progress towards personal goals and achievements consistent with the individual's future vision. This regulation is consistent with standards established for individual plan development as set forth by the commission on the accreditation of rehabilitation facilities (CARF) and/or other program accreditation approved and adopted by the developmental disabilities division and the department of health. It is the policy of the developmental disabilities division (DDD), that to the extent permitted by funding, each individual receive supports and services that will assist and encourage independence and productivity in the community and attempt to prevent regression or loss of current capabilities. Services and supports include specialized and/or generic services, training, education and/or treatment as determined by the IDT and documented in the ISP. D. The intent is to provide choice and obtain opportunities for individuals to live, work and play with full participation in their communities. The following principles provide direction and purpose in planning for individuals with	Based on administrative record review, the Agency did not implement the ISP according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan for 1 of 26 individuals. As indicated by Individuals ISP the following was found with regards to the implementation of ISP Outcomes: **Family Living Data Collection/Data Tracking/Progress with regards to ISP Outcomes:** Individual #9 • None found regarding: Live Outcome/Action Step: "… will work on activity" for 7/2023. Action step is to be completed 1 time per week.	**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?)*: →	

developmental disabilities. [05/03/94; 01/15/97; Recompiled 10/31/01] Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 6 Individual Service Plan (ISP): 6.9** ISP Implementation and Monitoring All DD Waiver Provider Agencies with a signed SFOC are required to provide services as detailed in the ISP. The ISP must be readily accessible to Provider Agencies on the approved budget. (See Section II Chapter 20: Provider Documentation and Client Records) CMs facilitate and maintain communication with the person, their guardian, other IDT members, Provider Agencies, and relevant parties to ensure that the person receives the maximum benefit of their services and that revisions to the ISP are made as needed. All DD Waiver Provider Agencies are required to cooperate with monitoring activities conducted by the CM and the DOH. Provider Agencies are required to respond to issues at the individual level and agency level as described in Section II Chapter 16: Qualified Provider Agencies. **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. 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.			

Tag # 1A32.1 Administrative Case File: Individual Service Plan Implementation *(Not Completed at Frequency)*	Standard Level Deficiency		
NMAC 7.26.5.16.C and D Development of the ISP. Implementation of the ISP. The ISP shall be implemented according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan. C. The IDT shall review and discuss information and recommendations with the individual, with the goal of supporting the individual in attaining desired outcomes. The IDT develops an ISP based upon the individual's personal vision statement, strengths, needs, interests and preferences. The ISP is a dynamic document, revised periodically, as needed, and amended to reflect progress towards personal goals and achievements consistent with the individual's future vision. This regulation is consistent with standards established for individual plan development as set forth by the commission on the accreditation of rehabilitation facilities (CARF) and/or other program accreditation approved and adopted by the developmental disabilities division and the department of health. It is the policy of the developmental disabilities division (DDD), that to the extent permitted by funding, each individual receive supports and services that will assist and encourage independence and productivity in the community and attempt to prevent regression or loss of current capabilities. Services and supports include specialized and/or generic services, training, education and/or treatment as determined by the IDT and documented in the ISP. D. The intent is to provide choice and obtain opportunities for individuals to live, work and play with full participation in their communities. The following principles provide direction and purpose in planning for individuals with developmental disabilities. [05/03/94; 01/15/97; Recompiled 10/31/01]	Based on administrative record review, the Agency did not implement the ISP according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan for 2 of 26 individuals. As indicated by Individuals ISP the following was found with regards to the implementation of ISP Outcomes: **Supported Living Data Collection / Data Tracking/Progress with regards to ISP Outcomes**: Individual #10 • According to the Live Outcome; Action Step for "… will search for a simple recipe to prepare" is to be completed 1 time per week. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 6/2023. **Customized Community Supports Data Collection/Data Tracking/Progress with regards to ISP Outcomes**: Individual #22 • According to the Fun Outcome; Action Step for "… will practice songs" is to be completed 1 time per week. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 5/2023.	**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?)*: →	

Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 6 Individual Service Plan (ISP): 6.9** ISP Implementation and Monitoring All DD Waiver Provider Agencies with a signed SFOC are required to provide services as detailed in the ISP. The ISP must be readily accessible to Provider Agencies on the approved budget. (See Section II Chapter 20: Provider Documentation and Client Records) CMs facilitate and maintain communication with the person, their guardian, other IDT members, Provider Agencies, and relevant parties to ensure that the person receives the maximum benefit of their services and that revisions to the ISP are made as needed. All DD Waiver Provider Agencies are required to cooperate with monitoring activities conducted by the CM and the DOH. Provider Agencies are required to respond to issues at the individual level and agency level as described in Section II Chapter 16: Qualified Provider Agencies. **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. 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.			

Tag # 1A32.2 Individual Service Plan Implementation (Residential Implementation)	Standard Level Deficiency		
NMAC 7.26.5.16.C and D Development of the ISP. Implementation of the ISP. The ISP shall be implemented according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan. C. The IDT shall review and discuss information and recommendations with the individual, with the goal of supporting the individual in attaining desired outcomes. The IDT develops an ISP based upon the individual's personal vision statement, strengths, needs, interests and preferences. The ISP is a dynamic document, revised periodically, as needed, and amended to reflect progress towards personal goals and achievements consistent with the individual's future vision. This regulation is consistent with standards established for individual plan development as set forth by the commission on the accreditation of rehabilitation facilities (CARF) and/or other program accreditation approved and adopted by the developmental disabilities division and the department of health. It is the policy of the developmental disabilities division (DDD), that to the extent permitted by funding, each individual receive supports and services that will assist and encourage independence and productivity in the community and attempt to prevent regression or loss of current capabilities. Services and supports include specialized and/or generic services, training, education and/or treatment as determined by the IDT and documented in the ISP. D. The intent is to provide choice and obtain opportunities for individuals to live, work and play with full participation in their communities. The following principles provide direction and	Based on residential record review, the Agency did not implement the ISP according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan for 3 of 21 individuals. As indicated by Individuals ISP the following was found with regards to the implementation of ISP Outcomes: **Supported Living Data Collection/Data Tracking / Progress with regards to ISP Outcomes**: Individual #1 • According to the Live Outcome; Action Step for "… will care for plants" is to be completed 3 times per week. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 8/1 – 18, 2023. (Date of home visit: 8/22/2023) **Family Living Data Collection/Data Tracking / Progress with regards to ISP Outcomes:** Individual #13 • According to the Live Outcome; Action Step for "… will complete one load of laundry (wash, dry, and put away), for the duration of the ISP year" is to be completed 1 time per week. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 8/1 – 18, 2023. (Date of home visit: 8/22/2023) • According to the Fun Outcome; Action Step for "… will participate in 30 minutes of physical activity" is to be completed 3 times per week. Evidence found indicated it was	**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?)*: →	

purpose in planning for individuals with developmental disabilities. [05/03/94; 01/15/97; Recompiled 10/31/01] Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 6 Individual Service Plan (ISP): 6.9** ISP Implementation and Monitoring All DD Waiver Provider Agencies with a signed SFOC are required to provide services as detailed in the ISP. The ISP must be readily accessible to Provider Agencies on the approved budget. (See Section II Chapter 20: Provider Documentation and Client Records) CMs facilitate and maintain communication with the person, their guardian, other IDT members, Provider Agencies, and relevant parties to ensure that the person receives the maximum benefit of their services and that revisions to the ISP are made as needed. All DD Waiver Provider Agencies are required to cooperate with monitoring activities conducted by the CM and the DOH. Provider Agencies are required to respond to issues at the individual level and agency level as described in Section II Chapter 16: Qualified Provider Agencies. **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	not being completed at the required frequency as indicated in the ISP for 8/1 – 18, 2023. (Date of home visit: 8/22/2023) Individual #28 • None found regarding: Live Outcome/Action Step: "… will listen to music" for 8/2023. Action step is to be completed 2 times per week. (Date of home visit: 8/23/2023)		

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 are acceptable. 3. Provider Agencies are responsible for ensuring that all plans created by nurses, RDs, therapists or BSCs are present in all 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.			

Tag # LS14 Residential Service Delivery Site Case File (ISP and Healthcare Requirements)	Condition of Participation Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 6 Individual Service Plan (ISP)** The CMS requires a person-centered service plan for every person receiving HCBS. The DD Waiver's person-centered service plan is the ISP. **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. 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. 3. Provider Agencies are responsible for ensuring that all plans created by nurses, RDs, therapists or BSCs are present in all settings. 4. Provider Agencies must maintain records of all documents produced by agency personnel or contractors on behalf of each	After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur. Based on record review, the Agency did not maintain a complete and confidential case file in the residence for 9 of 21 Individuals receiving Living Care Arrangements. Review of the residential individual case files revealed the following items were not found, incomplete, and/or not current: **Healthcare Passport:** • Not Found (#1) • Not Current (#3, 6, 16, 18, 28) **Health Care Plans:** • Body Mass Index (#3) • Bowel and Bladder (#15) • Gastrointestinal (#15) • States of Care/Hygiene (#3) **Medical Emergency Response Plans:** • Allergies (#29) • Body Mass Index (#3,15) • Cardiac Circulatory Condition (#15) • Falls (#2) • Gastrointestinal (#6,15) • History of Hepatitis C (#3) • Oral Hygiene (#3)	**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?)*: →	

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. **20.5.4 Health Passport and Physician Consultation Form:** All Primary and Secondary Provider Agencies must use the *Health Passport* and *Physician Consultation* form generated from an e-CHAT in 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 *Physician Consultation* form. The *Physician Consultation* form contains a list of all current medications.			

Chapter 13 Nursing Services: 13.2.9.1 Health Care Plans (HCP): Health Care Plans are created to provide guidance for the Direct Support Professionals (DSP) to support health related issues. Approaches that are specific to nurses may also be incorporated into the HCP. Healthcare Plans are based upon the eCHAT and the nursing assessment of the individual's needs. **13.2.9.2 Medical Emergency Response Plan (MERP): 1)** The agency nurse is required to develop a Medical Emergency Response Plan (MERP) for all conditions automatically triggered and marked with an "R" in the e-CHAT summary report. The agency nurse should use their clinical judgment and input from. 2) MERPs are required for persons who have one or more conditions or illnesses that present a likely potential to become a life-threatening situation.			

Tag # LS14.1 Residential Service Delivery Site Case File (Other Req. Documentation)	Standard Level Deficiency		
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. 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. 3. Provider Agencies are responsible for ensuring that all plans created by nurses, RDs, therapists or BSCs are present in all 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	Based on record review, the Agency did not maintain a complete and confidential case file in the residence for 1 of 21 Individuals receiving Living Care Arrangements. Review of the residential individual case files revealed the following items were not found, incomplete, and/or not current: **Behavior Crisis Intervention Plan:** • Not Found (#10)	**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?)*: →	

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.			

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 Eff 11/1/2021 **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. 1. DSP/DSS must successfully complete within 30 calendar days of hire and prior to working alone with a person in service: a. Complete IST requirements in accordance with the specifications described in the ISP of each person supported and as outlined in Chapter 17.9 Individual Specific Training below. b. Complete DDSD training in standards precautions located in the New Mexico Waiver Training Hub. c. Complete and maintain certification in First Aid and CPR. The training materials shall meet OSHA requirements/guidelines. d. Complete relevant training in accordance with OSHA requirements (if job involves exposure to hazardous chemicals). e. Become certified in a DDSD-approved system of crisis prevention and intervention (e.g., MANDT, Handle with Care, Crisis Prevention and Intervention (CPI)) before using Emergency Physical Restraint (EPR). Agency DSP and DSS shall maintain certification in a DDSD-approved system if any person they	Based on record review, the Agency did not ensure Orientation and Training requirements were met for 2 of 176 Direct Support Professional, Direct Support Supervisory Personnel and / or Service Coordinators. Review of Agency training records found no evidence of the following required DOH/DDSD trainings being completed: **First Aid:** • Not Found (#579, 697)	**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?)*: →	

support has a BCIP that includes the use of EPR. f. Complete and maintain certification in a DDSD-approved Assistance with Medication Delivery (AWMD) course if required to assist with medication delivery. g. Complete DDSD training regarding the HIPAA located in the New Mexico Waiver Training Hub. **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. 1. A SC must successfully complete within 30 calendar days of hire and prior to working alone with a person in service: a. Complete IST requirements in accordance with the specifications described in the ISP of each person supported, and as outlined in the Chapter 17.10 Individual-Specific Training below. b. Complete DDSD training in standard precautions located in the New Mexico Waiver Training Hub. c. Complete and maintain certification in First Aid and CPR. The training materials shall meet OSHA requirements/guidelines. d. Complete relevant training in accordance with OSHA requirements (if job involves exposure to hazardous chemicals). e. Become certified in a DDSD-approved system of crisis prevention and intervention (e.g., MANDT, Handle with Care, CPI) before using emergency physical restraint. Agency SC shall maintain certification in a DDSD-			

approved system if a person they support has a Behavioral Crisis Intervention Plan that includes the use of emergency physical restraint. f. Complete and maintain certification in AWMD if required to assist with medications. g. Complete DDSD training regarding HIPAA located in the New Mexico Waiver Training Hub.			

Tag # 1A22 Agency Personnel Competency	Standard Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **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 additional coaching/feedback. Individuals shall receive services from competent and qualified Provider Agency personnel who must successfully complete IST	Based on interview, the Agency did not ensure training competencies were met for 3 of 28 Direct Support Professional. **When DSP were asked, what State Agency do you report suspected Abuse, Neglect or Exploitation to, the following was reported:** • DSP #582 stated, "I mean I don't know what State agency. If I seem funny answering these questions it is cause I am getting over a stomach sickness so I might not be answering them right." Staff was not able to identify the State Agency as Division of Health Improvement. **When DSP were asked, if the Individual had Positive Behavioral Supports Plan (PBSP), If have they had been trained on the PBSP and what does the plan cover, the following was reported:** • DSP #520 stated, "I don't know." According to the Individual Specific Training Section of the ISP the Individual requires a Positive Behavioral Supports Plan. (Individual #10) **When DSP were asked, if the Individual had Behavioral Crisis Intervention Plan (BCIP), If have they had been trained on the BCIP and what does the plan cover, the following was reported:** • DSP 684 stated, "He says he doesn't have one, he is sitting here next to me. I don't think he does." According to the Individual Specific Training Section of the ISP the individual has Behavioral Crisis Intervention Plan. (Individual #3) **When DSP were asked, if they knew what the Individual's health condition / diagnosis**	**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?)*: →	

requirements in accordance with the specifications described in the ISP of each person supported. 1. IST must be arranged and conducted at least annually. IST includes training on the ISP Desired Outcomes, Action Plans, Teaching and Support Strategies, and information about the person's preferences regarding privacy, communication style, and routines. More frequent training may be necessary if the annual ISP changes before the year ends. 2. IST for therapy-related Written Direct Support Instructions (WDSI), Healthcare Plans (HCPs), Medical Emergency Response Plan (MERPs), Comprehensive Aspiration Risk Management Plans (CARMPs), Positive Behavior Supports Assessment (PBSA), Positive Behavior Supports Plans (PBSPs), and Behavior Crisis Intervention Plans (BCIPs), PRN Psychotropic Medication Plans (PPMPs), and Risk Management Plans (RMPs) must occur at least annually and more often if plans change, or if monitoring by the plan author or agency finds problems with implementation, when new DSP or CM are assigned to work with a person, or when an existing DSP or CM requires a refresher. 3. The competency level of the training is based on the IST section of the ISP. 4. The person should be present for and involved in IST whenever possible. 5. Provider Agencies are responsible for tracking of IST requirements. 6. Provider Agencies must arrange and ensure that DSP's and CIE's are trained on the contents of the plans in accordance with timelines indicated in the Individual-Specific Training Requirements: Support Plans section of the ISP and notify the plan authors when new DSP are hired to arrange for trainings.	**or when the information could be found, the following was reported:** • DSP #684 stated, "I went through his book, but I didn't go over it thoroughly it and we are not home right now. I am not sure what health conditions he has or his diagnoses." Per the Health Passport the Individual has a diagnosis of Opioid Dependence, Major Depressive disorder, Generalized Anxiety disorder, Autistic disorder, Asperger's Syndrome, Attention-Deficit disorder, Insomnia, and Hearing loss. (Individual #3) **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 #684 stated, "I assume yes because he is on the waiver, he just had surgery on his back. I do not have his book." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual requires Health Care Plans for Body Mass Index and Status of Care/Hygiene. (Individual #3) **When DSP were asked, if the Individual had any food and / or medication allergies that could be potentially life threatening, the following was reported:** • DSP #684 stated, " No." As indicated by the Health Passport the individual is allergic to Lamictal. (Individual #3)		

7. If a therapist, BSC, nurse, or other author of a plan, healthcare or otherwise, chooses to designate a trainer, that person is still responsible for providing the curriculum to the designated trainer. The author of the plan is also responsible for ensuring the designated trainer is verifying competency in alignment with their curriculum, doing periodic quality assurance checks with their designated trainer, and re-certifying the designated trainer at least annually and/or when there is a change to a person's plan.			

Tag # 1A26 Employee Abuse Registry	Standard Level Deficiency		
NMAC 7.1.12.8 - REGISTRY ESTABLISHED; PROVIDER INQUIRY REQUIRED: Upon the effective date of this rule, the department has established and maintains an accurate and complete electronic registry that contains the name, date of birth, address, social security number, and other appropriate identifying information of all persons who, while employed by a provider, have been determined by the department, as a result of an investigation of a complaint, to have engaged in a substantiated registry-referred incident of abuse, neglect or exploitation of a person receiving care or services from a provider. Additions and updates to the registry shall be posted no later than two (2) business days following receipt. Only department staff designated by the custodian may access, maintain and update the data in the registry. A. **Provider requirement to inquire of registry**. A provider, prior to employing or contracting with an employee, shall inquire of the registry whether the individual under consideration for employment or contracting is listed on the registry. B. **Prohibited employment.** A provider may not employ or contract with an individual to be an employee if the individual is listed on the registry as having a substantiated registry-referred incident of abuse, neglect or exploitation of a person receiving care or services from a provider. C. **Applicant's identifying information required**. In making the inquiry to the registry prior to employing or contracting with an employee, the provider shall use identifying information concerning the individual under consideration for employment or contracting sufficient to reasonably and completely search the registry, including the name, address, date of birth, social security number, and other	Based on record review, the Agency did not maintain documentation in the employee's personnel records that evidenced inquiry into the Employee Abuse Registry prior to employment for 1 of 205 Agency Personnel. **The following Agency Personnel records contained evidence that indicated the Employee Abuse Registry check was completed after hire:** **Service Coordination Personnel (SC):** • #669 – Date of hire 5/18/2022, completed 5/20/2022.	**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?)*: →	

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

Tag # 1A43.1 General Events Reporting: Individual Reporting	Standard Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 19 Provider Reporting Requirements:** DOH-DDSD collects and analyzes system wide information for quality assurance, quality improvement, and risk management in the DD Waiver Program. Provider Agencies are responsible for tracking and reporting to DDSD in several areas on an individual and agency wide level. The purpose of this chapter is to identify what information Provider Agencies are required to report to DDSD and how to do so. **19.2 General Events Reporting (GER):** The purpose of General Events Reporting (GER) is to report, track and analyze events, which pose a risk to adults in the DD Waiver program, but do not meet criteria for ANE or other reportable incidents as defined by the IMB. Analysis of GER is intended to identify emerging patterns so that preventative action can be taken at the individual, Provider Agency, regional and statewide level. On a quarterly and annual basis, DDSD analyzes GER data at the provider, regional and statewide levels to identify any patterns that warrant intervention. Provider Agency use of GER in Therap is required as follows: 1. DD Waiver Provider Agencies approved to provide Customized In- Home Supports, Family Living, IMLS, Supported Living, Customized Community Supports, Community Integrated Employment, Adult Nursing and Case Management must use the GER 2. DD Waiver Provider Agencies referenced above are responsible for entering specified information into a Therap GER module entry per standards set through the Appendix B GER Requirements and as identified by DDSD.	Based on record review, the Agency did not follow the General Events Reporting requirements as indicated by the policy for 2 of 26 individuals. **The following General Events Reporting records contained evidence that indicated the General Events Report was not entered and / or approved within 2 business days and / or entered within 30 days for medication errors:** **Individual #27** • General Events Report (GER) indicates on 1/9/2023 the Individual was transported to the ER due to losing her balance. (Injury). GER was approved 4/26/2023. • General Events Report (GER) indicates on 8/22/2023 the Individual sprained her ankle and was transported to the ER. (Injury). GER was approved 8/25/2023. **The following events were not reported in the General Events Reporting System as required by policy:** **Individual #9** • Documentation reviewed indicates on 5/15/2023 the Individual was transported to the ER due to the Feeding tube falling out. (Hospital). No GER was found.	**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?)*: →	

3. At the Provider Agency's discretion additional events, which are not required by DDSD, may also be tracked within the GER section of Therap. Events that are tracked for internal agency purposes and do not meet reporting requirements per DD Waiver Service Standards must be marked with a notification level of "Low" to indicate that it is being used internal to the provider agency. 4. GER does not replace a Provider Agency's obligations to report ANE or other reportable incidents as described in Chapter 18: Incident Management System. 5. GER does not replace a Provider Agency's obligations related to healthcare coordination, modifications to the ISP, or any other risk management and QI activities. 6. Each agency that is required to participate in General Event Reporting via Therap should ensure information from the staff and/or individual with the most direct knowledge is part of the report. a. Each agency must have a system in place that assures all GERs are approved per Appendix B GER Requirements and as identified by DDSD. b. Each is required to enter and approve GERs within 2 business days of discovery or observation of the reportable event. **19.2.1 Events Required to be Reported in GER:** The following events need to be reported in the Therap GER: when they occur during delivery of Supported Living, Family Living, Intensive Medical Living, Customized In-Home Supports, Customized Community Supports, Community Integrated Employment or Adult Nursing Services for DD Waiver participants aged 18 and older: 1. Emergency Room/Urgent Care/Emergency Medical Services			

2. Falls Without Injury 3. Injury (including Falls, Choking, Skin Breakdown and Infection) 4. Law Enforcement Use 5. All Medication Errors 6. Medication Documentation Errors 7. Missing Person/Elopement 8. Out of Home Placement- Medical: Hospitalization, Long Term Care, Skilled Nursing or Rehabilitation Facility Admission 9. PRN Psychotropic Medication 10. Restraint Related to Behavior 11. Suicide Attempt or Threat 12. COVID-19 Events to include COVID-19 vaccinations.			

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 # 1A09 Medication Delivery Routine Medication Administration	**Condition of Participation Level Deficiency**		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **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 20.6 Medication Administration Record (MAR) **Chapter 20 Provider Documentation and Client Records: 20.6 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. Providers have until November 1, 2022, to have a current Electronic Medication Administration Record online in Therap in all settings where medications or treatments are delivered. 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,	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 and August 2023. Based on record review, 2 of 12 individuals had Medication Administration Records (MAR), which contained missing medications entries and/or other errors: Individual #6 August 2023 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Aripiprazole (Abilify) 5 mg (1 time daily) – Blank 8/20, 21 (8:00 PM) • Clonazepam (Klonopin) 0.5 mg (2 times daily) – Blank 8/20, 21 (8:00 PM) • Docusate Sodium (Colace) 100 mg (1 time daily) – Blank 8/20, 21 (8:00 PM) • Fluvoxamine Maleate (Luvox) 100 mg (1 time daily) – Blank 8/20, 21 (8:00 PM) • Magnesium Oxide 500 mg (1 time daily) – Blank 8/20, 8/21 (8:00 PM) • Metamucil Powder 3.4 g/12g (2 times daily) – Blank 8/20, 21 (8:00 PM)	**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?)***:** →	

ANS for Medication Oversight must be budgeted, a 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 communicating 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	Individual #15 August 2023 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Oxybutyin CL ER (Ditropan) 10 mg (1 time daily) – Blank 8/20, 21 (8:00 PM) • Risperidone (Risperdal) 0.5 mg (1 time daily) – Blank 8/20, 21 (8:00 PM) • Simvastatin (Zocor) 20 mg (1 time daily) – Blank 8/20, 21 (8:00 PM) • Trazadone (Desyrel) 50 mg (1 time daily) – Blank 8/20, 21 (8:00 PM) • ZENPEP (Pancrelipase) 25000 unit (3 times daily) – Blank 8/20 (8:00 PM), 8/21 (3:00 PM, 8:00 PM)		

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 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 # 1A09.1 Medication Delivery PRN Medication Administration	Condition of Participation Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **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 20.6 Medication Administration Record (MAR) **Chapter 20 Provider Documentation and Client Records: 20.6 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. Providers have until November 1, 2022, to have a current Electronic Medication Administration Record online in Therap in all settings where medications or treatments are delivered. 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 MAR online in Therap must be created and used by the DSP.	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 and August 2023. Based on record review, 11 of 12 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #1 August 2023 As indicated by the Medication Administration Record the individual is to take the following medication. The following medications were not in the Individual's home. • Cough Drops (PRN) • Diphenhydramine (Benadryl) 25mg (PRN) • Imodium (Loperamide) 1mg/7.5ml (PRN) • Milk of Magnesium (PRN) • Pink Bismuth (Pepto Bismol) 262 mg/15ml (PRN) • Polyethylene Glycol 3350/17gm (PRN) • Sore Throat spray (Chloraseptic) (PRN) • Tums (Calcium Carbonate Antacid) 500mg (PRN) • Tussin DM (Robitussin DM) 10-100mg/ml (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?)*: →	

4. Provider Agencies must configure and use the MAR when assisting with medication. 5. Provider Agencies Continually communicating 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	• Aloe Vera gel 100% (PRN) • Saline Nasal Spray (PRN) • Sunscreen SPF 30 (PRN) • Triamcinolone Acitonide 0.5 % (PRN) • Triple Antibiotic Ointment (PRN) Individual #2 August 2023 As indicated by the Medication Administration Record the individual is to take the following medication. The following medications were not in the Individual's home. • Loratadine (Claritin) 10mg (PRN) • Magnesium Hydroxide Milk of Magnesium 400mg/5ml (PRN) • Polyethylene Glycol 3350/17gm (PRN) • Robitussin DM-SF 200-20 mg/10ml (PRN) • Tums (Calcium Carbonate Antacid) 500mg (PRN) • Aloe Vera gel (PRN) • Hemorrhoid Cream 14.4%-0.25%-1%-15% (PRN) • Hydrocortisone (Anti-Itch) 1% (PRN) • Pataday 0.2 % Eye Drop (PRN) • Saline Nasal Spray 0.45% (PRN) • Triple Antibiotic Ointment (PRN)		

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 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:	• Vicks (Chest Vaporizing medicated rub) (PRN) • Zinc Oxide 40% (Boudreaux's Butt Paste) (PRN) Individual #3 August 2023 As indicated by the Medication Administration Record the individual is to take the following medication. The following medications were not in the Individual's home. • Acetaminophen ES (Tylenol ES) 500mg (PRN) • Pink Bismuth (Pepto Bismol) 525 mg/30ml (PRN) • Tums (Calcium Carbonate Antacid) 500mg (PRN) • Dextromethorphan (Tussin DM/Robitussin DM Sugar Free) 20-200mg/ml (PRN) • Diphenhydramine (Benadryl) 25mg (PRN) • Ibuprofen (Advil) 200mg (PRN) • Ibuprofen (Motrin) 800mg (PRN) • Loratadine (Claritin) Non-Drowsy 10mg (PRN) • Imodium (Loperamide) 2mg (PRN) • Magnesium Hydroxide Milk of Magnesium 400mg/5ml (PRN) • Pataday/Olopatadine HCL 0.2 % Eye Drop (PRN) • Polyethylene Glycol 3350/17gm (PRN)		

➢ 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.	Individual #6 August 2023 As indicated by the Medication Administration Record the individual is to take the following medication. The following medications were not in the Individual's home. • Acetaminophen ES (Tylenol ES) 500mg (PRN) • Benadryl (Diphenhydramine) 25mg (PRN) • Ibuprofen (Motrin/Advil) 200mg (PRN) • Imodium (Loperamide) 2mg (PRN) • Milk of Magnesium (Magnesium Hydroxide) RS 400mg/5ml (PRN) • MiraLAX Polyethylene Glycol 3350/17gm (PRN) • Mylanta (Magnesium Hydroxide/alum/sim) 200-200 mg-20mg/5ml (PRN) • Pink Bismuth (Pepto Bismol) 525mg/30ml (PRN) • Tums (Calcium Carbonate Antacid) 500mg (PRN) • Tussin DM (Robitussin DM) 200 mg/20mg /10ml (PRN) • Aloe Vera gel (PRN) • Boudreaux's Butt Paste (Zinc Oxide) 40% (PRN) • Hemorrhoid Cream 14.4%-0.25%-1%-15% (PRN) • Hydrocortisone 1% (PRN)		

	• Insect Repellent Spray (PRN) • Pataday/Olopatadine HCL 0.2 % Eye Drop (PRN) • Saline Nasal Spray 0.45% (PRN) • Sunscreen SPF 50 (PRN) • Triple Antibiotic Ointment (PRN) • Vicks (Chest Vaporizing medicated rub) (PRN) Individual #8 July 2023 Physician's Orders indicated the following medication were to be given. The following Medications were not documented on the Medication Administration Records: • Ibuprofen 200 mg (PRN) • Milk of Magnesia 400 mg/5 ml (PRN) • Mylanta 200-200-20 mg/5 ml (PRN) • Pepto Bismol/Bismuth Subsalicylate 525 mg/30 ml (PRN) • Imodium/Loperamide 2 mg (PRN) • Benadryl/Diphenhydramine 25 mg (PRN) August 2023 As indicated by the Medication Administration Record the individual is to take the following medication. The following medications were not in the Individual's home. • Acetaminophen ES (Tylenol ES) 500mg (PRN) • Loratadine (Claritin) 10mg (PRN)		

	• MiraLAX Polyethylene Glycol 3350 (PRN) • Robitussin DM 200mg/20mg /10ml (PRN) • Tums (Calcium Carbonate Antacid) 500mg (PRN) • Aloe Vera gel (PRN) • Hemorrhoid Cream 14.4%-0.25%-1%-15% (PRN) • Hydrocortisone 1% (PRN) • Insect Repellent Spray (PRN) • Pataday/Olopatadine HCL 0.2 % Allergy Relief Eye Drop (PRN) • Saline Nasal Spray 0.45% (PRN) • Zinc Oxide (Boudreaux's Butt Paste) 40% (PRN) Individual #10 August 2023 As indicated by the Medication Administration Record the individual is to take the following medication. The following medications were not in the Individual's home. • Imodium (Loperamide) 2mg (PRN) • Milk of Magnesium (Magnesium Hydroxide) 400mg/5ml (PRN) • Mylanta (Magnesium Hydroxide/alum/sim) 200-200 mg-20mg/5ml (PRN) • Pepto Bismol (Bismuth Subsalicylate) 525 mg/30ml (PRN)		

	• Polyethylene Glycol (MiraLAX) 3350/17 gm (PRN) • Aloe Vera gel (PRN) • Hemorrhoid Cream 14.4%-0.25%-1%-15% (PRN) • Insect Repellent Spray (PRN) • Pataday/Olopatadine HCL 0.2 % Allergy Relief Eye Drop (PRN) • Deep Sea (ayr saline) 0.65% (PRN) • Sunscreen SPF 50 (PRN) • Vicks (Chest Vaporizing medicated rub) (PRN) • Boudreaux's Butt Paste (Zinc Oxide) 40% (PRN) Individual #15 August 2023 As indicated by the Medication Administration Record the individual is to take the following medication. The following medications were not in the Individual's home. • Acetaminophen ES (Tylenol ES) 500mg (PRN) • Benadryl (Diphenhydramine) 25mg (PRN) • Ibuprofen (Motrin/Advil) 200mg (PRN) • Imodium (Loperamide) 2mg (PRN) • Milk of Magnesium (Magnesium Hydroxide) RS 400mg/5ml (PRN)		

	• MiraLAX Polyethylene Glycol 3350/17gm (PRN) • Mylanta (Magnesium Hydroxide/alum/sim) 200-200 mg-20mg/5ml (PRN) • Pepto Bismol (Bismuth Subsalicylate) 525 mg/30ml (PRN) • Tums (Calcium Carbonate Antacid) 500mg (PRN) • Tussin/Robitussin DM (Dextromethorphan) 200mg/20mg /10ml (PRN) • Aloe Vera gel (PRN) • Hemorrhoid Cream 14.4%-0.25%-1%-15% (PRN) • Hydrocortisone 1% (PRN) • Insect Repellent Spray (PRN) • Pataday/Olopatadine HCL 0.2 % Eye Drop (PRN) • Saline Nasal Spray 0.45% (PRN) • Sunscreen SPF 50 (PRN) • Triple Antibiotic Ointment (PRN) • Vicks (Chest Vaporizing medicated rub) (PRN) • Zinc Oxide (Boudreaux's Butt Paste) 40% (PRN) Individual #16 August 2023 As indicated by the Medication Administration Record the individual is to		

	take the following medication. The following medications were not in the Individual's home. • Imodium (Loperamide) 2mg (PRN) • Loratadine (Claritin) Non-Drowsy 10mg (PRN) • Milk of Magnesium Regular Strength 400mg/5ml (PRN) • Mylanta 200-200mg-20mg/5ml (PRN) • Robitussin/Tussin DM (Dextromethorphan) Sugar Free 20mg/200mg (PRN) • Pataday/Olopatadine HCL 0.2 % Allergy Relief Eye Drop (PRN) • Zinc Oxide (Boudreaux's Butt Paste) 40% (PRN) Individual #17 August 2023 As indicated by the Medication Administration Record the individual is to take the following medication. The following medications were not in the Individual's home. • Mylanta (Magnesium Hydroxide/alum/sim) 200-200mg-20mg/5ml (PRN) • Tussin/Robitussin DM (Dextromethorphan) 200mg/20mg /10ml (PRN) • Tums (Calcium Carbonate Antacid) 500mg (PRN) • Aloe Vera gel (PRN) • Hemorrhoid Cream 14.4%-0.25%-1%-15% (PRN)		

	• Hydrocortisone 1% (PRN) • Insect Repellent Spray (PRN) • Pataday/Olopatadine HCL 0.2 % Eye Drop (PRN) • Saline Nasal Spray 0.45% (PRN) • Sunscreen SPF 50 (PRN) • Triple Antibiotic Ointment (PRN) • Vicks (Chest Vaporizing medicated rub) (PRN) • Zinc Oxide (Boudreaux's Butt Paste) 40% (PRN) • Benadryl (Diphenhydramine) 25mg (PRN) • Benzonatate (Tessalon per) 100mg (PRN) • Loratadine (Claritin) Non-Drowsy 10mg (PRN) • Milk of Magnesium (Magnesium Hydroxide) RS 400mg/5ml (PRN) Individual #18 August 2023 As indicated by the Medication Administration Record the individual is to take the following medication. The following medications were not in the Individual's home. • Benadryl (Diphenhydramine) 25mg (PRN) • Imodium (Loperamide) 2mg (PRN) • Milk of Magnesium (Magnesium Hydroxide) 400mg/5ml (PRN)		

	• Pepto Bismol (Bismuth Subsalicylate) 525mg/30ml (PRN) • MiraLAX Polyethylene Glycol 3350/17gm (PRN) • Mylanta (Magnesium Hydroxide) 200-200 mg-20mg/5ml (PRN) • Tums (Calcium Carbonate Antacid) 500mg (PRN) • Aloe Vera gel (PRN) • Hemorrhoid Cream 14.4%-0.25%-1%-15% (PRN) • Saline Nasal Spray 0.45% (PRN) • Sunscreen SPF 50 (PRN) • Triple Antibiotic Ointment (PRN) Individual #29 August 2023 As indicated by the Medication Administration Record the individual is to take the following medication. The following medications were not in the Individual's home. • Benadryl (Diphenhydramine) 25mg (PRN) • Imodium (Loperamide) 2mg (PRN) • Loratadine (Claritin) Non-Drowsy 10mg (PRN) • Milk of Magnesium (Magnesium Hydroxide) 400mg/5ml (PRN) • Mylanta (Magnesium Hydroxide/alum/sim) 200-200 mg-20mg/5ml (PRN)		

	• Pepto Bismol (Bismuth Subsalicylate) 525 mg/30ml (PRN) • Tums (Calcium Carbonate Antacid) 500mg (PRN) • Tussin/Robitussin DM (DM-Sugar Free) 200mg/20mg /10ml (PRN) • Aloe Vera gel (PRN) • Hemorrhoid Cream 14.4%-0.25%-1%-15% (PRN) • Hydrocortisone 1% (PRN) • Insect Repellent Spray (PRN) • Pataday/Olopatadine HCL 0.2 % Allergy Relief Eye Drop (PRN) • Saline Nasal Spray 0.45% (PRN) • Sunscreen SPF 50 (PRN) • Vicks (Chest Vaporizing medicated rub) (PRN) • Zinc Oxide (Boudreaux's Butt Paste) 40% (PRN)		

Tag # 1A09.1.0 Medication Delivery PRN Medication Administration	Standard Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **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 20.6 Medication Administration Record (MAR) **Chapter 20 Provider Documentation and Client Records: 20.6 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. Providers have until November 1, 2022, to have a current Electronic Medication Administration Record online in Therap in all settings where medications or treatments are delivered. 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 MAR online in Therap must be created and used by the DSP.	Medication Administration Records (MAR) were reviewed for the months of July and August 2023. Based on record review, 2 of 12 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #10 July 2023 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Banophen 25mg – PRN – 7/12 (given 1 time) Individual #29 August 2023 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Clonazepam (Klonopin) 0.5 mg – PRN – 8/8 (given 1 time)	**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?)*: →	

4. Provider Agencies must configure and use the MAR when assisting with medication. 5. Provider Agencies Continually communicating 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 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 # 1A09.2 Medication Delivery Nurse Approval for PRN Medication	Condition of Participation Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **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 20.6 Medication Administration Record (MAR) **Chapter 13 Nursing Services: 13.2 General Nursing Services Requirements and Scope of Services:** The following general requirements are applicable for all RNs and LPNs in the DD Waiver. This section represents the scope of nursing services. Refer to Chapter 10 Living Care Arrangements (LCA) for residential provider agency responsibilities related to nursing. Refer to Chapter 11.6 Customized Community Supports (CCS) for agency responsibilities related to nursing. **13.3.2.3 Medication Oversight:** Medication Oversight by a DD Waiver nurse is required in Family Living when a person lives with a non-related Family Living provider; for all JCMs; and whenever non-related DSP provide AWMD medication supports. 1. The nurse must respond to calls requesting delivery of PRN medications from AWMD trained DSP, non-related Family Living providers. 2. Family Living providers related by affinity or consanguinity (blood, adoption, or marriage) are not required to contact the	After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur. Based on record review, the Agency did not maintain documentation of PRN authorization as required by standard for 1 of 12 Individuals. Individual #29 August 2023 No documentation of the verbal authorization from the Agency nurse prior to each administration / assistance of PRN medication was found for the following PRN medication: • Clonazepam (Klonopin) 0.5 mg – PRN – 8/14 (given 1 time)	**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?)*: →	

nurse prior to assisting with delivery of a PRN medication. **13.2.8.1.3 Assistance with Medication Delivery by Staff (AWMD):** For people who do not meet the criteria to self-administer medications independently or with physical assistance, trained staff may assist with medication delivery if: 1. Criteria in the MAAT are met. 2. Current written consent has been obtained from the person/guardian/surrogate healthcare decision maker. 3. There is a current Primary Care Practitioner order to receive AWMD by staff. 4. Only AWMD trained staff, in good standing, may support the person with this service. 5. All AWMD trained staff must contact the on-call nurse prior to assisting with a PRN medication of any type. a. Exceptions to this process must comply with the DDSD Emergency Medication list as part of a documented MERP with evidence of DSP training to skill level.			

Tag # 1A15.2 Administrative Case File: Healthcare Documentation (Therap and Required Plans)	Condition of Participation Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 3: Safeguards: Decisions about Health Care or Other Treatment: Decision Consultation and Team Justification Process:** There are a variety of approaches and available resources to support decision making when desired by the person. The decision consultation and team justification processes assist 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. For current forms and resources please refer to the DOH Website: https://nmhealth.org/about/ddsd/. 3.1.1 **Decision Consultation Process (DCP):** Health decisions are the sole domain of waiver participants, their guardians or healthcare decision makers. Participants and their healthcare decision makers can confidently make decisions that are compatible with their personal and cultural values. Provider Agencies and Interdisciplinary Teams (IDTs) are required to support the informed decision making of waiver participants by supporting access to medical consultation, information, and other available resources 1. The Decision Consultation Process (DCP) is documented on the Decision Consultation and Team Justification Form (DC/TJF) and is used for health related issues when a person or their guardian/healthcare decision maker has concerns, needs more information about these types of issues or has decided not to follow all or part of a healthcare-related order, recommendation, or suggestion. This includes, but is not limited to: a. medical orders or recommendations from the Primary Care Practitioner, Specialists or other licensed medical or healthcare	After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur. Based on record review, the Agency did not maintain the required documentation in the Individuals Agency Record as required by standard for 6 of 26 individual Review of the administrative individual case files revealed the following items were not found, incomplete, and/or not current: **Healthcare Passport:** • Did not contain Name of Physician (#1, 10, 22, 28) • Did not contain Insurance Information (#1, 10, 22, 29) • Did not contain Guardianship/Healthcare Decision Maker (#8, 22, 28, 29) • Did not contain Emergency Contact Information (#10, 22, 28)	**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?)*: →	

practitioners such as a Nurse Practitioner (NP or CNP), Physician Assistant (PA) or Dentist; b. clinical recommendations made by registered/licensed clinicians who are either members of the IDT (e.g., nurses, therapists, dieticians, BSCs or PRS Risk Evaluator) or clinicians who have performed evaluations such as a video-fluoroscopy; c. health related recommendations or suggestions from oversight activities such as the Individual Quality Review (IQR); and d. recommendations made by a licensed professional through a Healthcare Plan (HCP), including a Comprehensive Aspiration Risk Management Plan (CARMP), a Medical Emergency Response Plan (MERP) or another plan such as a Risk Management Plan (RMP) or a Behavior Crisis Intervention Plan (BCIP). **Chapter 10 Living Care Arrangements: Supported Living Requirements: 10.4.1.5.1 Monitoring and Supervision:** Supported Living Provider Agencies must: 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. 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).			

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. 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. 3. Provider Agencies are responsible for ensuring that all plans created by nurses, RDs, therapists or BSCs are present in all 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 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. **20.5.4 Health Passport and Physician Consultation Form:** All Primary and Secondary Provider Agencies must use the *Health Passport* and *Physician Consultation* form generated from an e-CHAT in 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 *Physician Consultation* form. The *Physician Consultation* form contains a list of all current medications. **Chapter 13 Nursing Services: 13.1 Overview of The Nurse's Role in The DD Waiver and Larger Health Care System:** Routine medical and healthcare services are accessed through the person's Medicaid State Plan benefits and through Medicare and/or private insurance for persons who have these additional types of insurance coverage. DD Waiver health related services are specifically designed to support the person in the community setting and complement but may not duplicate those medical or health related services provided by the Medicaid State Plan or other insurance systems. Nurses play a pivotal role in supporting persons and their guardians or legal Health Care Decision makers within the DD Waiver and are a key link with the larger healthcare system in New Mexico. DD Waiver Nurses identify and support the person's preferences regarding health decisions; support health awareness and self-management of medications and health conditions; assess, plan, monitor and manage health related issues; provide education; and share information among the IDT members including DSP in a variety of settings, and share information with natural supports when requested by individual or			

guardian. Nurses also respond proactively to chronic and acute health changes and concerns, facilitating access to appropriate healthcare services. This involves communication and coordination both within and beyond the DD Waiver. DD Waiver nurses must contact and consistently collaborate with the person, guardian, IDT members, Direct Support Professionals and all medical and behavioral providers including Medical Providers or Primary Care Practitioners (physicians, nurse practitioners or physician assistants), Specialists, Dentists, and the Medicaid Managed Care Organization (MCO) Care Coordinators. **13.2.7 Documentation Requirements for all DD Waiver Nurses** **13.2.8 Electronic Nursing Assessment and Planning Process** **13.2.8.1 Medication Administration Assessment Tool (MAAT)** **13.2.8.2 Aspiration Risk Management Screening Tool (ARST)** **13.2.8.3 The Electronic Comprehensive Health Assessment Tool (e-CHAT)** **13.2.9.1 Health Care Plans (HCP)** **13.2.9.2 Medical Emergency Response Plan (MERP)**			

Tag # 1A27.2 Duty to Report IRs Filed During On-Site and/or IRs Not Reported by Provider	Standard Level Deficiency		
NMAC 7.1.14.8 INCIDENT MANAGEMENT SYSTEM REPORTING REQUIREMENTS FOR COMMUNITY-BASED SERVICE PROVIDERS: **A. Duty to report:** **(1)** All community-based providers shall immediately report alleged crimes to law enforcement or call for emergency medical services as appropriate to ensure the safety of consumers. **(2)** All community-based service providers, their employees and volunteers shall immediately call the department of health improvement (DHI) hotline at 1-800-445-6242 to report abuse, neglect, exploitation, suspicious injuries or any death and also to report an environmentally hazardous condition which creates an immediate threat to health or safety. **B. Reporter requirement.** All community-based service providers shall ensure that the employee or volunteer with knowledge of the alleged abuse, neglect, exploitation, suspicious injury, or death calls the division's hotline to report the incident. **C. Initial reports, form of report, immediate action and safety planning, evidence preservation, required initial notifications:** **(1) Abuse, neglect, and exploitation, suspicious injury or death reporting:** Any person may report an allegation of abuse, neglect, or exploitation, suspicious injury or a death by calling the division's toll-free hotline number 1-800-445-6242. Any consumer, family member, or legal guardian may call the division's hotline to report an allegation of abuse, neglect, or exploitation, suspicious injury or death directly, or may report through the community-based service provider who, in addition to calling the hotline, must also utilize the division's abuse, neglect, and exploitation or report of death form.	Based on interview and observation, the Agency did not report suspected abuse, neglect, or exploitation**,** unexpected and natural/expected deaths; or other reportable incidents as required to the Division of Health Improvement. **During the on-site survey on August 21 – September 1, 2023 12:00 PM, surveyors observed the following:** During QMB's home visit on 8/23/2023 for Individual #21, Surveyors drove up to the residence. The outside had debris such as: couches, appliances, and trash that surrounded the front and back of the residence. The wooden rail and steps leading up to the front door were broken and detached from the home. Surveyors had to stretch their legs across the broken steps and grab on to the doorframe to pull themselves up in order to enter the home. When entering the home, Surveyors noticed several dogs in cages with dog feces on the Living Room floor and hallway. DSP #635 told Surveyors there was no running water in the home. Surveyors did observe human feces in the toilet. Surveyors could smell human body odor and animal feces throughout the residence. The ceiling had holes that were covered up with sheets and blankets in all of the rooms to the mobile home. In the kitchen, rodent droppings were observed on the countertops and on the pans that were hanging in between the cabinets on hooks. Surveyors observed knee high piles of trash and clothes in all rooms. Incident report was reported to APS and DHI. **As a result of what was observed the following incident(s) was reported:**	**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?)***:** →	

The abuse, neglect, and exploitation or report of death form and instructions for its completion and filing are available at the division's website, http://dhi.health.state.nm.us, or may be obtained from the department by calling the division's toll free hotline number, 1-800-445-6242. **(2) Use of abuse, neglect, and exploitation or report of death form and notification by community-based service providers:** In addition to calling the division's hotline as required in Paragraph (2) of Subsection A of 7.1.14.8 NMAC, the community-based service provider shall also report the incident of abuse, neglect, exploitation, suspicious injury, or death utilizing the division's abuse, neglect, and exploitation or report of death form consistent with the requirements of the division's abuse, neglect, and exploitation reporting guide. The community-based service provider shall ensure all abuse, neglect, exploitation or death reports describing the alleged incident are completed on the division's abuse, neglect, and exploitation or report of death form and received by the division within 24 hours of the verbal report. If the provider has internet access, the report form shall be submitted via the division's website at http://dhi.health.state.nm.us; otherwise it may be submitted via fax to 1-800-584-6057. The community-based service provider shall ensure that the reporter with the most direct knowledge of the incident participates in the preparation of the report form. **(3) Limited provider investigation:** No investigation beyond that necessary in order to be able to report the abuse, neglect, or exploitation and ensure the safety of consumers is permitted until the division has completed its investigation. **(4) Immediate action and safety planning:** Upon discovery of any alleged incident of abuse, neglect, or exploitation, the community-based service provider shall:	**Individual #21** • A State ANE Report was filed as a result of the following: On 8/23/2023 1:00 PM, an Incident report was reported to DHI.		

(a) develop and implement an immediate action and safety plan for any potentially endangered consumers, if applicable; **(b)** be immediately prepared to report that immediate action and safety plan verbally, and revise the plan according to the division's direction, if necessary; and **(c)** provide the accepted immediate action and safety plan in writing on the immediate action and safety plan form within 24 hours of the verbal report. If the provider has internet access, the report form shall be submitted via the division's website at http://dhi.health.state.nm.us; otherwise it may be submitted by faxing it to the division at 1-800-584-6057. **(5) Evidence preservation:** The community-based service provider shall preserve evidence related to an alleged incident of abuse, neglect, or exploitation, including records, and do nothing to disturb the evidence. If physical evidence must be removed or affected, the provider shall take photographs or do whatever is reasonable to document the location and type of evidence found which appears related to the incident. **(6) Legal guardian or parental notification:** The responsible community-based service provider shall ensure that the consumer's legal guardian or parent is notified of the alleged incident of abuse, neglect and exploitation within 24 hours of notice of the alleged incident unless the parent or legal guardian is suspected of committing the alleged abuse, neglect, or exploitation, in which case the community-based service provider shall leave notification to the division's investigative representative. **(7) Case manager or consultant notification by community-based service providers:** The responsible community-based service provider shall notify the consumer's case manager or consultant within 24 hours that an alleged incident involving abuse, neglect, or exploitation has been reported to the division. Names of other consumers and employees may			

be redacted before any documentation is forwarded to a case manager or consultant. **(8) Non-responsible reporter:** Providers who are reporting an incident in which they are not the responsible community-based service provider shall notify the responsible community-based service provider within 24 hours of an incident or allegation of an incident of abuse, neglect, and exploitation.			

Tag # 1A29 Complaints / Grievances Acknowledgement	Standard Level Deficiency		
NMAC 7.26.3.6: A. These regulations set out rights that the department expects all providers of services to individuals with developmental disabilities to respect. These regulations are intended to complement the department's Client Complaint Procedures (7 NMAC 26.4) [now 7.26.4 NMAC]. **NMAC 7.26.3.13 Client Complaint Procedure Available.** A complainant may initiate a complaint as provided in the client complaint procedure to resolve complaints alleging that a service provider has violated a client's rights as described in Section 10 [now 7.26.3.10 NMAC]. The department will enforce remedies for substantiated complaints of violation of a client's rights as provided in client complaint procedure. [09/12/94; 01/15/97; Recompiled 10/31/01] **NMAC 7.26.4.13 Complaint Process: A. (2).** The service provider's complaint or grievance procedure shall provide, at a minimum, that: **(a)** the client is notified of the service provider's complaint or grievance procedure Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Appendix A Client File Matrix**	Based on record review, the Agency did not provide documentation, the complaint procedure had been made available to individuals or their legal guardians for 1 of 26 individuals. Review of the Agency individual case files revealed the following items were not found and/or incomplete: **Grievance/Complaint Procedure Acknowledgement:** • Not Current (#14)	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)*: → **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to affect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)*: →	

Tag # 1A39 Assistive Technology and Adaptive Equipment	Standard Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 12 Professional Services: 12.4.1 Participatory Approach:** The "Participatory Approach" is person-centered and asserts that no one is too severely disabled to benefit from assistive technology and other therapy supports that promote participation in life activities. The Participatory Approach rejects the premise that an individual shall be "ready" or demonstrate certain skills before assistive technology can be provided to support function. **12.4.7.3 Assistive Technology (AT) Services, Remote Personal Support Technology (RPST) and Environmental Modifications:** Therapists support the person to access and utilize AT, RPST and Environmental Modifications through the following requirements: 1. Therapists are required to be or become familiar with AT and RPST related to that therapist's practice area and used or needed by individuals on that therapist's caseload. 2. Therapists are required to provide a current AT Inventory to each Living Supports and CCS site where AT is used, for each person using AT related to that therapist's scope of service. 3. Therapists are required to initiate or update the AT Inventory annually, by the 190th day following the person's ISP effective date, so that it accurately identifies the assistive technology currently in use by the individual and related to that therapist's scope of service. 4. Therapists are required to maintain professional documentation related to the delivery of services related to AT, RPST and Environmental Modifications. (Refer to	Based on interview, the Agency did not ensure the necessary support mechanisms and devices, including the rationale for the use of assistive technology or adaptive equipment is in place for 1 of 26 Individuals. **When DSP were asked, if the Individual require any type assistive device or adaptive equipment and if they had been trained on the equipment, the following was reported:** • DSP #554 stated, "None." Per the Health and Safety section on the ISP, the individual requires Eyeglasses, Hearing Aids, and Orthotic Shoes. (Individual #25)	**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?)*: →	

Chapter 14: Other Services for more information about these services.) 5. Therapists must respond to requests to perform in-home evaluations and make recommendations for environmental modifications, as appropriate. **Chapter 10 Living Care Arrangements (LCA): 10.3.8 Requirements for Each Residence: Scope of Living Supports (Supported Living, Family Living, and IMLS)** 7. ensuring readily available access to and assistance with use of a person's adaptive equipment, augmentative communication, remote personal support technology (RPST) and assistive technology (AT) devices, including monitoring and support related to maintenance of such equipment and devices to ensure they are in working order; **Chapter 11 Community Inclusion:** Exploring, facilitating, developing, requesting, and implementing job accommodations and the use of assistive technology to help an individual be successful in employment			

Tag # 1A50.1 Individual: Scope of Services *(Individual Interviews)*	Standard Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 4 Person-Centered Planning (PCP):** **4.1 Essential Elements of Person-Centered Planning (PCP):** Person-centered planning is a process that places a person at the center of planning their life and supports. The CMS requires use of PCP in the development of the ISP. It is an ongoing process that is the foundation for all aspects of the DD Waiver Program and DD Waiver Provider Agencies' work with people with I/DD. The process is designed to identify the strengths, capacities, preferences, and needs of the person. The process may include other people chosen by the person, who are able to serve as important contributors to the process. Overall, PCP involves person-centered thinking, person-centered service planning, and person-centered practice. PCP enables and assists the person to identify and access a personalized mix of paid and non-paid services and supports to assist him or her to achieve personally defined outcomes in the community. **4.1.1 Person-Centered Thinking:** Person-centered thinking involves a process of examining the individual's values, strengths, needs and skills to set the foundation for ISP development. Person-centered thinking respects and supports the person with I/DD to develop strategies to: 1. have informed choices; 2. exercise the same basic civil and human rights as other citizens; 3. have personal control over the life they prefer in the community of choice; 4. be valued for contributions to their community; and 5. be supported through a network of resources, both natural and paid.	Based on interview, the Agency did not provide the essential elements of person centered planning as indicated in Individuals interview for 1 of 26 individuals. **When the Individuals receiving services were asked, if there was an accessible vehicle available to transport them to work, appointments, shopping, activities of their choosing, the following was reported:** • Individual #21 stated, "No, we have to get rides from friends but no one wants to help us."	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)*: → **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to affect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)*: →	

Tag # LS06 Family Living Requirements	Standard Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 10 Living Care Arrangements (LCA) Living Supports Family Living:** **10.3.9.2.1** Monitoring and Supervision Family Living Provider Agencies must: 1. Provide and document monthly face-to-face consultation in the Family Living home conducted by agency supervisors or internal service coordinators with the DSP and the person receiving services to include: a. reviewing implementation of the person's ISP, Outcomes, Action Plans, and associated support plans, including HCPs, MERPs, Health Passport, PBSP, CARMP, WDSI; b. scheduling of activities and appointments and advising the DSP regarding expectations and next steps, including the need for IST or retraining from a nurse, nutritionist, therapists or BSC; and c. assisting with resolution of service or support issues raised by the DSP or observed by the supervisor, service coordinator, or other IDT members. 2. Monitor that the DSP implement and document progress of the AT inventory, Remote Personal Support Technology (RPST), physician and nurse practitioner orders, therapy, HCPs, PBSP, BCIP, PPMP, RMP, MERPs, and CARMPs. **10.3.9.2.1.1 Home Study:** An on-site Home Study is required to be conducted by the Family Living Provider agency initially, annually, and if there are any changes in the home location, household makeup, or other significant event. 1. The agency person conducting the Home Study must have a bachelor's degree in Human Services or related field or be at least 21 years of age, HS Diploma or GED	Based on record review, the Agency did not complete all DDSD requirements for approval of each direct support provider for 4 of 10 individuals. Review of the Agency files revealed the following items were not found, incomplete, and/or not current: **Family Living (Annual Update) Home Study:** • Individual #8 – Incomplete, did not include Health and Safety checklist. Last completed on 3/30/2023. • Individual #9 – Incomplete, did not include Health and Safety checklist. Last completed on 4/24/2023. • Individual #13 – Incomplete, did not include Health and Safety checklist. Last completed on 4/18/2023. • Individual #25 - Incomplete, did not include Health and Safety checklist. Last completed on 4/4/2023.	**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?)*: →	

and a minimum of 1-year experience with I/DD. 2. The Home Study must include a health and safety checklist assuring adequate and safe: a. Heating, ventilation, air conditioning cooling; b. Fire safety and Emergency exits within the home; c. Electricity and electrical outlets; and d. Telephone service and access to internet, when possible. 3. The Home Study must include a safety inspection of other possible hazards, including: a. Swimming pools or hot tubs; b. Traffic Issues; c. Water temperature that does not exceed a safe temperature (110^0 F). Anyone with a history of being unsafe in or around water while bathing, grooming, etc. or with a history of at least one scalding incident will have a regulated temperature control valve or device installed in the home. d. Any needed repairs or modifications 4. The home setting must comply with the CMS Final Settings Rule and ensure tenant protections, privacy, and autonomy.			

Tag # LS25 Residential Health & Safety (Supported Living / Family Living / Intensive Medical Living)	Standard Level Deficiency		

Developmental Disabilities Waiver Service Standards Eff 11/1/2021
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:
1. has basic utilities, i.e., gas, power, water, telephone, and internet access;
2. supports telehealth, and/ or family/friend contact on various platforms or using various devices;
3. has a battery operated or electric smoke detectors or a sprinkler system, carbon monoxide detectors, and fire extinguisher;
4. has a general-purpose first aid kit;
5. has accessible written documentation of evacuation drills occurring at least three times a year overall, one time a year for each shift;
6. has water temperature that does not exceed a safe temperature (110^0 F). Anyone with a history of being unsafe in or around water while bathing, grooming, etc. or with a history of at least one scalding incident will have a regulated temperature control valve or device installed in the home.
7. 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;
8. has an emergency placement plan for relocation of people in the event of an emergency evacuation that makes the residence unsuitable for occupancy;
	Based on observation, the Agency did not ensure that each individuals' residence met all requirements within the standard for 10 of 20 Living Care Arrangement residences.
	
	Review of the residential records and observation of the residence revealed the following items were not found, not functioning or incomplete:
	
	Supported Living Requirements:
	
	• Poison Control Phone Number (#3)
	
	• Water temperature in home exceeds safe temperature (110^0 F):
	  • Water temperature in home measured 112^0 F (#3)
	  • Water temperature in home measured 127^0 F (#4)
	  • Water temperature in home measured 132.4^0 F (#17)
	  • Water temperature in home measured 114^0 F (#10, 18)
	  • Water temperature in home measured 129^0 F (#29)
	
	Note: The following Individuals share a residence:
	• #6, 15
	
	Family Living Requirements:
	
	• Water temperature in home exceeds safe temperature (110^0 F)
		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?)*: →	

9. has emergency evacuation procedures that address, but are not limited to, fire, chemical and/or hazardous waste spills, and flooding; 10. supports environmental modifications, remote personal support technology (RPST), and assistive technology devices, including modifications to the bathroom (i.e., shower chairs, grab bars, walk in shower, raised toilets, etc.) based on the unique needs of the individual in consultation with the IDT; 11. has or arranges for necessary equipment for bathing and transfers to support health and safety with consultation from therapists as needed; 12. has the phone number for poison control within line of site of the telephone; 13. has general household appliances, and kitchen and dining utensils; 14. has proper food storage and cleaning supplies; 15. has adequate food for three meals a day and individual preferences; and 16. has at least two bathrooms for residences with more than two residents. 17. Training in and assistance with community integration that include access to and participation in preferred activities to include providing or arranging for transportation needs or training to access public transportation. 18. Has Personal Protective Equipment available, when needed	• Water temperature in home measured 118^0 F (#19) • Water temperature in home measured 115^0 F (#9) • Water temperature in home measured 133^0 F (#13) • Water temperature in home measured 130.8^0 F (#22)		

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 # IS30 Customized Community Supports Reimbursement	**Standard Level Deficiency**		
NMAC 8.302.2 Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **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. 4. A Provider Agency that receives payment for treatment, services or goods must retain all medical and business records relating to any of the following for a period of at least six years from the payment date: a. treatment or care of any eligible recipient;	Based on record review, the Agency did not provide written or electronic documentation as evidence for each unit billed for Customized Community Supports services for 3 of 15 individuals. Individual #4 May 2023 • The Agency billed 108 units of Customized Community Supports (H2021 HB U1) from 5/1/2023 through 5/15/2023. Documentation received accounted for 102 units. June 2023 • The Agency billed 48 units of Customized Community Supports (H2021 HB U1) from 6/16/2023 through 6/30/2023. Documentation received accounted for 44 units. Individual #17 June 2023 • The Agency billed 58 units of Customized Community Supports (H2021 HB U1) from 6/1/2023 through 6/15/2023. Documentation received accounted for 46 units. *(Note: Void/Adjust provided on-site during survey. Provider please complete POC for ongoing QA/QI.)* Individual #18 June 2023 • The Agency billed 244 units of Customized Community Supports (H2021 HB U1) from 6/1/2023 through 6/15/2023.	**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?)***:** →	

b. services or goods provided to any eligible recipient; c. amounts paid by MAD on behalf of any eligible recipient; and d. any records required by MAD for the administration of Medicaid. **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. **21.9.2 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.	Documentation received accounted for 176 units.		

Tag # LS27 Family Living Reimbursement	Standard Level Deficiency		
NMAC 8.302.2 Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **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. 4. A Provider Agency that receives payment for treatment, services or goods must retain all medical and business records relating to any of the following for a period of at least six years from the payment date: a. treatment or care of any eligible recipient;	Based on record review, the Agency did not provide written or electronic documentation as evidence for each unit billed for Family Living Services for 1 of 10 individuals. Individual #13 June 2023 • The Agency billed 7 units of Family Living (T2033 HB) from 6/8/2023 through 6/14/2023. Documentation received accounted for 6 units. • The Agency billed 7 units of Family Living (T2033 HB) from 6/15/2023 through 6/21/2023. Documentation received accounted for 1 unit.	**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?)*: →	

b. services or goods provided to any eligible recipient; c. amounts paid by MAD on behalf of any eligible recipient; and d. any records required by MAD for the administration of Medicaid. **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.			

NEW MEXICO
Department of Health
Division of Health Improvement

MICHELLE LUJAN GRISHAM
Governor

PATRICK M. ALLEN
Cabinet Secretary

Date:	December 11, 2023
To:	Kami Silva, Executive Director
Provider:	Lessons of Life LLC
Address:	1720 S. Telshor Blvd
State/Zip:	Las Cruces, New Mexico 88011
E-mail Address:	ksilva@lessonsoflifellc.com
Region:	Southeast and Southwest
Survey Date:	August 21 – September 1, 2023
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	Supported Living, Family Living, Customized In-Home Supports; Customized Community Supports, and Community Integrated Employment Services
Survey Type:	Routine

Dear Ms. Silva:

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

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Quality Management Bureau/DHI

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