NEW MEXICO
Department of Health
Division of Health Improvement

MICHELLE LUJAN GRISHAM
Governor

PATRICK M. ALLEN
Cabinet Secretary

Date:	November 29, 2023
To:	Emad Elmaoued, Director
Provider: Address: State/Zip:	ADID Care INC 5115 Copper Avenue NE Albuquerque, New Mexico 87408
E-mail Address:	emad@adidcare.com
Region: Survey Date:	Metro and Northeast October 16 - 27, 2023
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	Supported Living, Family Living, Customized In-Home Supports, Customized Community Supports
Survey Type:	Routine
Team Leader:	Ashley Gueths, BACJ, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau
Team Members:	Wolf Krusemark, BFA, Healthcare Surveyor Supervisor, Division of Health Improvement/Quality Management Bureau; Monica Valdez, BS, Healthcare Surveyor Advanced / Plan of Correction Coordinator, Division of Health Improvement/Quality Management Bureau; Jamie Pond, BS, Staff Manager, Division of Health Improvement/Quality Management Bureau; Sally Karingada, BS, Healthcare Surveyor Supervisor, Division of Health Improvement/Quality Management Bureau; Kaitlyn Taylor, BSW, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; Koren Chandler, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Dear Mr. Emad Elmaoued,

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:

Non-Compliance: This determination is based on noncompliance with 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 or any amount of Standard Level Tags with 6 or more Condition of Participation Level Tags *(refer to Attachment D for details)*. The attached QMB Report of Findings indicates Standard Level and Condition of Participation Level deficiencies identified and requires completion and implementation of a Plan of Correction.

The following tags are identified as Condition of Participation Level:

- Tag # 1A08.3 Administrative Case File: Individual Service Plan / ISP Components
- Tag # LS14 Residential Service Delivery Site Case File (ISP and Healthcare Requirements)
- Tag # 1A20 Direct Support Professional Training
- Tag # 1A22 Agency Personnel Competency
- Tag # 1A05 General Requirements / Agency Policy and Procedure Requirements
- Tag # 1A08.2 Administrative Case File: Healthcare Requirements & Follow-up
- 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 Administrative Case File (Other Required Documents)
- 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 # 1A38 Living Care Arrangement / Community Inclusion Reporting Requirements
- Tag # IS12 Person Centered Assessment (Community Inclusion)
- Tag # LS14.1 Residential Service Delivery Site Case File (Other Req. Documentation)
- Tag # 1A37 Individual Specific Training
- Tag # 1A09.0 Medication Delivery Routine Medication Administration
- Tag # 1A09.1.0 Medication Delivery PRN Medication Administration
- Tag # 1A29 Complaints / Grievances Acknowledgement
- Tag # 1A31.2 Human Right Committee Composition
- Tag # 1A33 Board of Pharmacy: Med. Storage
- Tag # 1A33.1 Board of Pharmacy – License
- Tag # LS06 Family Living Requirements
- Tag # LS25 Residential Health & Safety (Supported Living / Family Living / Intensive Medical Living)
- Tag # IS30 Customized Community Supports Reimbursement
- Tag # LS26 Supported Living Reimbursement
- Tag # IH32 Customized In-Home Supports 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,

Ashley Gueths, BACJ

Ashley Gueths, BACJ
Team Lead/Healthcare Surveyor
Division of Health Improvement
Quality Management Bureau

Survey Process Employed:

Administrative Review Start Date:	October 16, 2023
Contact:	**ADID Care INC.** Emad Elmaoued, Director **DOH/DHI/QMB** Ashley Gueths, BACJ, Team Lead/Healthcare Surveyor
Entrance Conference Date:	October 16, 2023
Present:	**ADID Care INC.** Emad Elmaoued, Director Ana Hill, DDW Nurse **DOH/DHI/QMB** Ashley Gueths, BACJ, Team Lead/Healthcare Surveyor Wolf Krusemark, BFA, Healthcare Surveyor Supervisor Jamie Pond, BS, Staff Manager
Exit Conference Date:	October 27, 2023
Present:	**ADID Care INC.** Emad Elmaoued, Director **DOH/DHI/QMB** Ashley Gueths, BACJ, Team Lead/Healthcare Surveyor Wolf Krusemark, BFA, Healthcare Surveyor Supervisor Monica Valdez, BS, Healthcare Surveyor Advanced / Plan of Correction Coordinator Jamie Pond, BS, Staff Manager Sally Karingada, BS, Healthcare Surveyor Supervisor **DDSD - Metro Regional Office** Marie Velasco, DDW Program Manager Maura L. Emerine-Danbury, Social and Community Services Coordinator
Administrative Locations Visited:	0 *(Administrative portion of survey completed remotely)*
Total Survey Sample Size:	24 0 – *Former Jackson Class Members* 24 - Non-*Jackson* Class Members 14 - Supported Living 8 - Family Living 2 - Customized In-Home Supports 11 - Customized Community Supports
Total Homes Visits	17
❖ Supported Living Homes Visited	8 *Note: The following Individuals share a SL residence:* • #12, 24

- #6, 21
- #2, 18
- #4, 22
- #1, 25
- #5, 14

❖ Family Living Homes Visited	8
❖ Customized In-Home Support Homes Visited	1
Total Wellness Visits Completed:	33
Persons Served Records Reviewed	24
Persons Served Interviewed	23
Persons Served Not Seen and/or Not Available	1 *(Note: 1 Individual was not available during the on-site survey)*
Direct Support Professional Records Reviewed	96
Direct Support Professional Interviewed	24
Substitute Care/Respite Personnel Records Reviewed	4
Service Coordinator Records Reviewed	3
Administrative Interview	1
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 <u>Living Care Arrangements and Community Inclusion</u> are as follows:

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

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

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

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

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

- **1A20 -** Direct Support 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: Microsoft Word - IRF-QMB-Form.doc (nmhealth.org)
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 Administrative Case File (Other Required Documents)	**Standard Level Deficiency**		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 20: Provider Documentation and Client Records: 20.1 HIPAA:** DD Waiver Provider Agencies shall comply with all applicable requirements of the Health Insurance Portability and Accountability Act of 1996 (HIPAA) and the Health Information Technology for Economic and Clinical Health Act of 2009 (HITECH). All DD Waiver Provider Agencies are required to store information and have adequate procedures for maintaining the privacy and the security of individually identifiable health information. HIPPA compliance extends to electronic and virtual platforms. **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	Based on record review, the Agency did not maintain a complete and confidential case file at the administrative office for 10 of 24 individuals. Review of the Agency administrative individual case files revealed the following items were not found, incomplete, and/or not current: **Behavior Crisis Intervention Plan:** • Not Found (#6) **Speech Therapy Plan (Therapy Intervention Plan TIP):** • Not Found (#18) **Occupational Therapy Plan (Therapy Intervention Plan TIP):** • Not Found (#3, 4, 16) **Physical Therapy Plan (Therapy Intervention Plan TIP)**: • Not Found (#2, 4, 21) **Documentation of Guardianship/Power of Attorney:** • Not Found (#6, 8, 19) **IDT meeting Minutes:** • Individual #13 - Not Found for Hospital Discharge on 3/23/2023.	**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?)***:** →	

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. 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.	• Individual #19 - Not Found for Hospital Discharge on 6/4/2023.		

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, 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	Based on record review, the Agency did not maintain progress notes and other service delivery documentation for 9 of 24 Individuals. Review of the Agency individual case files revealed the following items were not found: **Administrative Case File:** **Family Living Progress Notes/Daily Contact Logs:** • Individual #10 - None found for 7/29/2023. **Residential Case File:** **Supported Living Progress Notes/Daily Contact Logs:** • Individual #2 - None found for 10/6 – 8, 2023. *(Date of home visit: 10/16/2023)* • Individual #12 - None found for 10/14/2023. *(Date of home visit: 10/18/2023)* • Individual #18 - None found for 10/1, 6 - 8, 14, 15, 2023. *(Date of home visit: 10/19/2023)* • Individual #23 - None found for 10/7/2023 *(Date of home visit: 10/17/2023)* **Family Living Progress Notes/Daily Contact Logs:** • Individual #8 - None found for 10/13 -17, 2023. *(Date of home visit: 10/18/2023)* • Individual #9 - None found for 10/9 - 19, 2023. *(Date of home visit: 10/20/2023)* • Individual #10 - None found for 10/1 - 7, 9 - 18, 2023. *(Date of home visit: 10/19/2023)*	**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?)*: →	

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.	• Individual #17 - None found for 10/11 - 18, 2023. *(Date of home visit: 10/19/2023)* • Individual #19 - None found for 10/1, 7, 8, 13 - 17, 2023. *(Date of home visit: 10/18/2023)*		

Tag # 1A08.3 Administrative Case File: Individual Service Plan / ISP Components	Condition of Participation Level Deficiency		
NMAC 7.26.5 SERVICE PLANS FOR INDIVIDUALS WITH DEVELOPMENTAL DISABILITIES LIVING IN THE COMMUNITY. **NMAC 7.26.5.12 DEVELOPMENT OF THE INDIVIDUAL SERVICE PLAN (ISP) - PARTICIPATION IN AND SCHEDULING OF INTERDISCIPLINARY TEAM MEETINGS.** **NMAC 7.26.5.14 DEVELOPMENT OF THE INDIVIDUAL SERVICE PLAN (ISP) - CONTENT OF INDIVIDUAL SERVICE PLANS.** 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. **6.6 DDSD ISP Template:** The ISP must be written according to templates provided by the DDSD. Both children and adults have designated ISP templates. The ISP template includes Vision Statements, Desired Outcomes, a meeting participant signature page, an Addendum A (i.e., an acknowledgement of receipt of specific information) and other elements depending on the age and status of the individual. The ISP templates may be revised and reissued by DDSD to incorporate initiatives that improve person - centered planning practices. Companion documents may also be issued by DDSD and be required for use to better demonstrate required elements of the PCP process and ISP development. **6.6.1 Vision Statements: The long-term vision statement describes the person's major long-term (e.g., within one to three**	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 at the administrative office for 5 of 24 individuals. Review of the Agency administrative individual case files revealed the following items were not found, incomplete, and/or not current: **Addendum A**: • Not Found (#1, 9, 12, 16, 21)	**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?)*: →	

years) life dreams and aspirations in the following areas: 1. Live, 2. Work/Education/Volunteer, 3. Develop Relationships/Have Fun, and 4. Health and/or Other (Optional). **6.6.2 Desired Outcomes:** A Desired Outcome is required for each life area (Live, Work, Fun) for which the person receives paid supports through the DD Waiver. Each service does not need its own, separate outcome, but should be connected to at least one Desired Outcome. **6.6.3.1 Action Plan:** Each Desired Outcome requires an Action Plan. The Action Plan addresses individual strengths and capabilities in reaching Desired Outcomes. **6.6.3.2 Teaching and Supports Strategies (TSS) and Written Direct Support Instructions (WDSI):** After the ISP meeting, IDT members conduct a task analysis and assessments necessary to create effective TSS and WDSI to support those Action Plans that require this extra detail. **6.6.3.3 Individual Specific Training in the ISP:** The CM, with input from each DD Waiver Provider Agency at the annual ISP meeting, completes the IST requirements section of the ISP form listing all training needs specific to the individual. **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.			

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 2 of 24 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 #14 • None found regarding: Live Outcome/Action Step: "… will organize one area of her room" for 9/2023. The action step is to be completed 1 time per week. **Customized Community Supports Data Collection / Data Tracking/Progress with regards to ISP Outcomes**: Individual #17 • None found regarding: Work/learn Outcome/Action Step: "With assistance, … will schedule his monthly activities" for 9/2023. The action step is to be completed 1 time per month. • None found regarding: Work/Learn Outcome/Action Step: "…will participate in the activity of his choice" for 9/2023. The action step is to be completed 1 time per month.	**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	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 7 of 24 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 #12 • According to the Live Outcome; Action Step for "… will complete his rehabilitation exercises" is to be completed 2 times per day. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 7/2023 - 9/2023. • According to the Live Outcome; Action Step for "… will participate in hygiene and other ADLs" is to be completed 1 time per day. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 7/2023 - 9/2023. Individual #24 • According to the Live Outcome; Action Step for "I will gather my needed supplies" is to be completed 2 times per day, 7 days a week. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 7/2023 – 9/2023. • According to the Live Outcome; Action Step for "I will complete the steps in my oral care	**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. 5. Each Provider Agency is responsible for maintaining the daily or other contact notes documenting the nature and frequency of	routine" is to be completed 2 times per day, 7 days a week. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 7/2023 - 9/2023. **Family Living Data Collection / Data Tracking/Progress with regards to ISP Outcomes:** Individual #3 • According to the Live Outcome; Action Step for "with assistance … will choose a recipe for baking" is to be completed 3 times per month. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 8/2023. • According to the Live Outcome; Action step for "with assistance … will bake the chosen item" is to be completed 3 times per month. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 8/2023. Individual #10 • According to the Health Outcome; Action Step for "… will work out at gym" 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 7/2023 - 9/2023. **Customized Community Supports Data Collection/Data Tracking/Progress with regards to ISP Outcomes**: Individual #1 • According to the Fun Outcome; Action Step for "… will plan and participate in a community event" is to be completed 2 times per week. Evidence found indicated it was not being completed at the required		

service delivery, as well as data tracking only for the services provided by their agency.	frequency as indicated in the ISP for 7/2023 – 9/2023. • According to the Fun Outcome; Action Step for "…will gather memories from the event and put them in her photo Album" is to be completed 2 times per week. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 7/2023 - 9/2023. Individual #17 • According to the Work/Learn Outcome; Action step for "…will participate in the activity of his choice" is to be completed 1 time per month. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 7/2023 - 8/2023. Individual #19 • According to the Fun Outcome; Action Step for "… will go to the gym" is to be completed 4 times per month. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 9/2023. • According to the Fun Outcome; Action Step for "… will do 30 minutes of weight bearing exercises" is to be completed 4 times per month. Evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 9/2023. • According to the Fun Outcome; Action Step for "… will swim for at least 30 minutes" is to be completed 4 times per month. Evidence found indicated it was not at frequency as indicated in the ISP for 9/2023.		

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 1 of 24 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 #12 • None found regarding: Live Outcome/Action Step: "… will complete his rehabilitation exercises" for 10/8 - 17, 2023. Action step is to be completed 2 times per day. Document maintained by the provider was blank. *(Date of home visit: 10/18/2023)* • None found regarding: Live Outcome/Action Step: "… will participate in hygiene and other ADLs" for 10/8 - 17, 2023. Action step is to be completed 1 time per day. Document maintained by the provider was blank. *(Date of home visit: 10/18/2023)*	**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?)*: →	

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			

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 # 1A38 Living Care Arrangement / Community Inclusion Reporting Requirements	Standard Level Deficiency		
7.26.5.17 DEVELOPMENT OF THE INDIVIDUAL SERVICE PLAN (ISP) - DISSEMINATION OF THE ISP, DOCUMENTATION AND COMPLIANCE: C. Objective quantifiable data reporting progress or lack of progress towards stated outcomes, and action plans shall be maintained in the individual's records at each provider agency implementing the ISP. Provider agencies shall use this data to evaluate the effectiveness of services provided. Provider agencies shall submit to the case manager data reports and individual progress summaries quarterly, or more frequently, as decided by the IDT. These reports shall be included in the individual's case management record and used by the team to determine the ongoing effectiveness of the supports and services being provided. Determination of effectiveness shall result in timely modification of supports and services as needed. Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 19 Provider Reporting Requirements: 19.5 Se**mi-Annual Reporting: The semi-annual report provides status updates to life circumstances, health, and progress toward ISP goals and/or goals related to professional and clinical services provided through the DD Waiver. This report is submitted to the CM for review and may guide actions taken by the person's IDT if necessary. Semi-annual reports may be requested by DDSD for QA activities. Semi-annual reports are required as follows: 1. DD Waiver Provider Agencies, except AT, EMSP, PRSC, SSE and Crisis Supports, must complete semi-annual.	Based on record review, the Agency did not complete written status reports as required for 5 of 24 individuals receiving Living Care Arrangements and Community Inclusion. **Supported Living Semi-Annual Reports:** • Individual #21 - None found for 3/2023 – 9/2023. *(Term of ISP 3/2023 - 3/2024).* **Family Living Semi- Annual Reports:** Individual #17 - None found for 3/2023 - 9/2023. *(Term of ISP 3/2023 - 3/2024).* **Customized Community Supports Semi-Annual Reports:** • Individual #8 - None found for 1/2023 - 7/2023. *(Term of ISP 1/2023 - 1/2024).* • Individual #17 - None found for 3/2023- 9/2023. *(Term of ISP 3/2023 - 3/2024.).* **Nursing Semi-Annual:** • Individual #12 - None found for 10/2022 - 4/2023. *(Term of ISP 10/2022 - 10/2023).* • Individual #18 – None found for 3/2023- 9/2023. *(Term of ISP 3/2023 - 3/2024).*	**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?)*: →	

2. The first semi-annual report will cover the time from the start of the person's ISP year until the end of the subsequent six-month period (180 calendar days) and is due ten calendar days after the period ends (190 calendar days). 3. The second semi-annual report is integrated into the annual report or professional assessment/annual re-evaluation when applicable and is due 14 calendar days prior to the annual ISP meeting. 4. Semi-annual reports must contain at a minimum written documentation of: a. the name of the person and date on each page; b. the timeframe that the report covers; c. timely completion of relevant activities from ISP Action Plans or clinical service goals during timeframe the report is covering; d. a description of progress towards Desired Outcomes in the ISP related to the service provided; e. a description of progress toward any service specific or treatment goals when applicable (e.g. health related goals for nursing); f. significant changes in routine or staffing if applicable; g. unusual or significant life events, including significant change of health or behavioral health condition; h. the signature of the agency staff responsible for preparing the report; and i. any other required elements by service type that are detailed in these standards. 5. Semi-annual reports must be distributed to the IDT members when due by SComm. 6. Semi-annual reports can be stored in individual document storage. **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. 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 # IS12 Person Centered Assessment (Community Inclusion)	Standard Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 11: Community Inclusion: 11.4 Person Centered Assessments (PCA) and Career Development Plans (CDP)** Agencies who are providing CCS and/or CIE are required to complete a person-centered assessment (PCA). A PCA is a person-centered planning tool that is intended to be used for the service agency to get to know the person whom they are supporting and to help identify the individual needs and strengths to be addressed in the ISP. The PCA should provide the reader with a good sense of who the person is and is a means of sharing what makes an individual unique. The information gathered in a PCA should be used to guide community inclusion services for the individual. Recommended methods for gathering information include paper reviews, interviews with the individual, guardian or anyone who knows the individual well including staff, family members, friends, BSC therapist, school personnel, employers, and providers. Observations in the community, home visits, neighborhood/environmental observations research on community resources, and team input are also reliable means of gathering valuable information. A Career Development Plan (CDP), developed by the CIE Provider Agency with input from the CCS Provider, must be in place for job seekers or those already working to outline the tasks needed to obtain, maintain, or seek advanced opportunities in employment. For those who are employed, the career development plan addresses topics such as a plan to fade paid supports from the worksite or strategies to improve opportunities for career advancement. CCS and CIE Provider Agencies must adhere to the following requirements related to a PCA and Career Development Plan:	Based on record review, the Agency did not maintain a confidential case file for Individuals receiving Inclusion Services for 3 of 24 individuals. Review of the Agency individual case files revealed the following items were not found, incomplete, and/or not current: • Annual Review - Person Centered Assessment (Individual #17, 21, 24)	**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?)*: →	

1. A PCA should contain, the following major topics, at a minimum: a. information about the person's background and current status; b. the person's strengths and interests and how they are known; c. conditions for success to integrate into the community, including conditions for job success (for those who are working or wish to work); and d. support needs for the individual. 2. The agency must involve the individual and describe how they were involved in development of the PCA. A guardian and those who know the person best must also be included in the development of the PCA, as applicable. 3. Timelines for completion: The initial PCA must be completed within the first 90 calendar days of the person receiving services. Thereafter, the Provider Agency must ensure that the PCA is reviewed and updated with the most current information, annually. A more extensive update of a PCA must be completed every five years. PCAs completed at the 5-year mark should include a narrative summary of progress toward outcomes from initial development, changes in support needs, major life changes, etc. If there is a significant change in a person's circumstance, a new PCA should be considered because the information in the PCA may no longer be relevant. A significant change may include but is not limited to losing a job, changing a residence or provider, and/or moving to a new region of the state. 4. If a person is receiving more than one type of service from the same provider, one PCA with information about each service is acceptable. 5. PCA's should be signed and dated to demonstrate that the assessment was reviewed and updated with the most current			

information, at least annually. 6. A career development plan is developed by the CIE provider with input from the CCS provider, as appropriate, and can be a separate document or be added as an addendum to a PCA. The career development plan should have specific action steps that identify who does what and by when.			

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 18 of 22 Individuals receiving Living Care Arrangements. Review of the residential individual case files revealed the following items were not found, incomplete, and/or not current: **Annual ISP:** • Not Found (#10) **ISP Teaching and Support Strategies:** ***Individual #1:*** *TSS not found for the following Live Outcome Statement / Action Steps:* • "...will establish an AM & PM routine." • "...will complete her daily AM & PM to do list on iPad." ***Individual #3:*** *TSS not found for the following Live Outcome Statement / Action Steps:* • "With assistance...will choose a recipe for baking." • " With assistance...will bake the chosen item." ***Individual #4:*** *TSS not found for the following Live Outcome Statement / Action Steps:*	**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.	• "With staff support …will supply Cooper with fresh food, fresh water, and medications." • " With staff support …to clean litter box." • " With staff support … and make a list to shop for needed cat supplies." ***Individual #5:*** *TSS not found for the following Live Outcome Statement / Action Steps:* • "I will show the house lead the verified account." ***Individual #6:*** *TSS not found for the following Live Outcome Statement / Action Steps:* • "… will assist in the meal prep of her choice." ***Individual #8:*** *TSS not found for the following Live Outcome Statement / Action Steps:* • "… will fold and put her clothing away." ***Individual #9:*** *TSS not found for the following Live Outcome Statement / Action Steps:* • "... will choose a snack or a meal that she wants to prepare monthly." • "Will prepare a snack or meal." *TSS not found for the following Fun Outcome Statement / Action Steps:* • "…will find a concert that she would like to attend." • "Will attend the concert once." ***Individual #12:***		

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 <u>conditions or illnesses that present a likely potential to become a life-threatening situation</u>.	*TSS not found for the following Live Outcome Statement / Action Steps:* • "… will complete his rehabilitation exercises." • "… will participate in hygiene and other ADLs." ***Individual #14:*** *TSS not found for the following Live Outcome Statement / Action Steps:* • "… will develop a system of organization." • "… will organize on area of her room." ***Individual #18:*** *TSS not found for the following Live Outcome Statement / Action Steps:* • "I will use equipment with staff support to transfer to the toilet." ***Individual #21:*** *TSS not found for the following Live Outcome Statement / Action Steps:* • "… will follow her recommended nutritional diet. ... will be following a 1500 calorie intake." *TSS not found for the following Fun Outcome Statement / Action Steps:* • "… will choose and participate in a community activity." ***Individual #24:*** *TSS not found for the following Live Outcome Statement / Action Steps:* • "I will gather my needed supplies." • "I will complete the steps in my oral care routine." ***Individual #25:***		

	TSS not found for the following Live Outcome Statement / Action Steps: • "...will purchase items for the garden." *TSS not found for the following Fun Outcome Statement / Action Steps:* • "...will research meals to make." • "...will create a shopping list of the items that he needs." • "...will call to confirm he has money on his true link card." • "...will prepare a meal." **Healthcare Passport:** • Not Found (#9, 19) • Not Current (#17) **Comprehensive Aspiration Risk Management Plan:** • Not Current (#25) **Health Care Plans:** • Falls (#22) • Respiratory (#25) • Seizures (#24) **Medical Emergency Response Plans:** • Allergies (#7, 18) • Aspiration (#7, 25) • Falls (#7, 22) • Respiratory (#25) • Seizures (#24)		

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 22 Individuals receiving Living Care Arrangements. Review of the residential individual case files revealed the following items were not found, incomplete, and/or not current: **Positive Behavioral Supports Plan:** • Not Current (#23)	**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	**Condition of Participation 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	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 ensure Orientation and Training requirements were met for 54 of 99 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 (#518, 523, 534, 544, 558, 567, 568, 569, 573, 574, 576, 578, 586, 588, 590, 591, 592) • Expired (#575, 580, 589) **CPR:** • Not Found (#518, 523, 533, 544, 558, 564, 567, 568, 569, 572, 573, 574, 576, 578, 586, 588, 590, 591, 592) • Expired (#575, 580, 589,) **Assisting with Medication Delivery:** • Not Found (#513, 515, 517, 518, 520, 522, 523, 524, 527, 532, 534, 536, 539, 541, 543, 547, 555, 558, 562, 567, 575, 581, 589, 599) • Expired (#504, 507, 529, 533, 535, 550, 554, 564, 565, 568, 569, 572, 573, 574, 576, 577, 579, 585, 586, 588, 591, 600, 601,)	**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	Condition of Participation 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	After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur. Based on interview, the Agency did not ensure training competencies were met for 15 of 24 Direct Support Professional. **When DSP were asked, what State Agency do you report suspected Abuse, Neglect or Exploitation to, the following was reported:** • DSP #514 stated, "DDSD." Staff was not able to identify the State Agency as the Division of Health Improvement or Adult Protective Services. • DSP #594 stated, "I never thought about that, because it has never happened to me before. I would contact ADID Care." Staff was not able to identify the State Agency as the Division of Health Improvement or Adult Protective Services. • DSP #602 stated, "NM Health for ANE." Staff was not able to identify the State Agency as the Division of Health Improvement or Adult Protective Services. **When DSP were asked to give examples of Abuse, Neglect and Exploitation, the following was reported:** • DSP #514 stated, "Disclosing information to another party." DSP's response with regards to exploitation. • DSP #549 stated, "I'm sorry I can't think of one right now." DSP's response with regards to exploitation.	**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.	• DSP #532 stated, "Not knowing if someone is hurt." DSP's response with regards to Abuse. **When DSP were asked, if they were provided with Individual Specific Training for the Individual they are supporting, the following was reported:** • DSP #532 stated, "They just told me what she likes. Like she is a late sleeper. I get her up early and we go out in the community. She is not a late sleeper anymore. I enjoy working with her. We laugh all day long." The surveyor asked if there was any other training they received to work with the Individual and the DSP stated, "No other training." (Individual #1) **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 #532 stated, "I don't know." According to the Individual Specific Training Section of the ISP, the Individual requires a Positive Behavioral Supports Plan. (Individual #1) **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 #532 stated, "I know she has a lady that visits with her, a behavior something." According to the Individual Specific Training Section of the ISP, the individual has a Behavioral Crisis Intervention Plan. (Individual #1)		

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.	• DSP #557 stated, "She does not." According to the Individual Specific Training Section of the ISP, the individual has a Behavioral Crisis Intervention Plan. (Individual #5) • DSP #541 stated, "I don't think she has one in there; we just call the BSC if we need to." According to the Individual Specific Training Section of the ISP, the individual has Behavioral Crisis Intervention Plan. (Individual #14) • DSP #588 stated, "I believe so, I am supposed to call his BSC and work on ways to deescalate him." According to the Individual Specific Training Section of the ISP, the individual does not have a Behavioral Crisis Intervention Plan. (Individual #17) • DSP #514 stated, "I don't think so, I'm not seeing one." According to the Individual Specific Training Section of the ISP, the individual has Behavioral Crisis Intervention Plan. (Individual #23) **When DSP were asked, if they knew what the Individual's health condition / diagnosis or where the information could be found, the following was reported:** • DSP #532 stated, "Like I said, she is a great little person. I don't know if she has any. She has not shown any to me." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual has a diagnosis of unspecified mood disorder, post-traumatic stress disorder, mild intellectual disabilities. (Individual #1) • DSP #515 stated, "Nothing really. Nothing stands out." As indicated by the Electronic		

	Comprehensive Health Assessment Tool, the Individual has a diagnosis of Moderate intellectual disabilities, Autistic Disorder, Unspecified hearing loss. (Individual #12) **When DSP were asked, if the Individual had a Comprehensive Aspiration Risk Management Plan (CARMP) and if they had been trained on the CARMP, the following was reported:** • DSP #532 stated, "I don't know." As indicated by the Individual Specific Training section of the ISP the individual requires a Comprehensive Aspiration Risk Management Plan (CARMP). (Individual #1) • DSP #508 stated, "No." As indicated by the Individual Specific Training section of the ISP the individual requires a Comprehensive Aspiration Risk Management Plan (CARMP). (Individual #18) **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 #574 stated, "Seizures, falls." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual additionally requires Health Care Plans for Constipation, Skin and Wound. (Individual #8) • DSP #508 stated, "No, I don't think so." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual requires Health Care Plans for Aspiration Risk, Spasticity, or Contractures, Skin and Wound. (Individual #18)		

	• DSP #578 stated, "It should be in my book, but I don't have any book. The agency has not given me a book." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual requires Health Care Plans for A1C Levels, Body Mass Index, Endocrine, Own Blood Glucose Monitoring, and Respiratory. (Individual #19) • DSP #507 stated, "No." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual requires Health Care Plans for Seizures. (Individual #24) **When DSP were asked, if the Individual had Medical Emergency Response Plans where could they be located and if they had been trained, the following was reported, the following was reported:** • DSP #536 stated, "No, I can't find them." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual requires Medical Emergency Response Plans for Aspiration Risk, Falls, Neuro (Devices and implants: cerebral shunt, Baclofen Pump, VNS) and Seizure Disorder. (Individual #2) • DSP #574 stated, "Not sure." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual requires Medical Emergency Response Plans for Aspiration Risk, Falls, and Seizure Disorder. (Individual #8) • DSP #515 stated, "I don't believe he has any MERPS." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual requires Medical Emergency		

	Response Plans for Aspiration Risk (Individual #12) • DSP #598 stated, “I don’t have that in front of me. The documents are in the room with the care giver, and she has covid. I don’t want to go into that area and Therap will not let me in.” As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual requires Medical Emergency Response Plans for Aspiration. (Individual #13) • DSP #578 stated, “I don't know, I was not given a book. I'm transferring to a new agency.” As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual requires Medical Emergency Response Plans for A1C Levels, Endocrine, Own Blood Glucose Monitoring, and Respiratory/Asthma. (Individual #19) **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 #594 stated, “None that I am aware of.” As indicated by the Health Passport, the individual is allergic to Penicillin. (Individual #17) • DSP #514 stated, “Not to my knowledge.” As indicated by the Health Passport, the individual is allergic to Sulfa. (Individual #23)		

Tag # 1A37 Individual Specific 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 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.	Based on record review, the Agency did not ensure that Individual Specific Training requirements were met for 13 of 99 Agency Personnel. Review of personnel records found no evidence of the following: **Direct Support Professional (DSP):** • Individual Specific Training (#502, 504, 506, 519, 534, 556, 558, 563, 590, 591, 593) **Service Coordination Personnel (SC):** • Individual Specific Training (#599, 600)	**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?)*: →	

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. 2. 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.			

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 # 1A05 General Requirements / Agency Policy and Procedure Requirements	**Condition of Participation Level Deficiency**		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 16 Qualified Provider Agencies:** Qualified DD Waiver Provider Agencies must deliver DD Waiver services. DD Waiver Provider Agencies must have a current Provider Agreement and continually meet required screening, licensure, accreditation, and training requirements as well as continually adhere to the DD Waiver Service Standards and relevant NMAC All Provider Agencies must comply with contract management activities to include any type of quality assurance review and/or compliance review completed by DDSD, the Division of Health Improvement (DHI) or other state agencies. **16.7 Compliance with Federal and State Rules and DD Waiver Service Standards** DD Waiver Provider agencies must comply with all applicable federal and state rules and DD Waiver Service Standards. Agencies are required to submit polices or procedural descriptions in their initial and renewal application which address applicable requirements. **16.7.1 Exception to the Standards:** In extraordinary circumstances, a Provider Agency may need to request an exception to the standards. An exception may be based on individual circumstances or extenuating circumstances at the agency. Any exception to the standards needs prior approval from DDSD according to the following: 1. For exceptions to standards that directly impact a person in service, the exception	After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur. Based on interview, the Agency did not develop, implement and / or comply with written policies and procedures to protect the physical / mental health of individuals that complies with all DDSD requirements. **When DSP were asked, what is the agency's on-call process, how on-call works, and how long does it take them to respond to you if you call the following was reported:** • DSP #585 stated, "No more than 24 hours." Per DDW Standards 13.2.6 On Call Nursing "An on-call nurse is required to be available to DSP in a timely manner. They must be able to respond within 15 minutes by phone and within 60 minutes in-person to assess the person if deemed necessary per prudent nursing practice." (Individual #15)	**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?)***:** →	

may be granted using the Exception Authorization Process, formerly known as the H Authorization Process, which requires the CM to submit the request on required forms along with supporting documentation to the respective DDSD Regional Office Director or designee for review and determination. 2. For exceptions to the standards related to service and/or agency requirements, the exception may be granted through a review of specific circumstances by designated DDSD staff, which requires the agency to submit the request to the local Regional Office. The local Regional Office forwards the request to the appropriate DDSD Management staff for review and determination. 3. All exceptions must be approved prior to implementing. 4. Federal and state requirements are considered when reviewing any requests for exceptions. 5. Any Provider Agency operating under an approved exception must have supporting documentation on file for quality review activities. 6. Exceptions may be time limited or revoked based on individual and/or agency circumstances. **NEW MEXICO DEPARTMENT OF HEALTH DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION: Provider Application** • Emergency and on-call procedures; • On-call nursing services that specifically state the nurse must be available to DSP during periods when a nurse is not present. The on-call nurse must be available to make an on-site visit when information provided by the DSP over the phone indicate, in the nurse's professional judgment, a need for a face to face assessment to determine			

appropriate action; • Incident Management Procedures that comply with the current NM Department of Health Improvement Incident Management Guide • Medication Assessment and Delivery Policy and Procedure; • Policy and procedures regarding delegation of specific nursing functions • Policies and procedures regarding the safe transportation of individuals in the community and how you will comply with the New Mexico regulations governing the operation of motor vehicles **STATE OF NEW MEXICO DEPARTMENT OF HEALTH DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION PROVIDER AGREEMENT: ARTICLE 39. POLICIES AND REGULATIONS** Provider Agreements and amendments reference and incorporate laws, regulations, policies, procedures, directives, and contract provisions not only of DOH, but of HSD. Additionally, the PROVIDER agrees to abide by all the following, whenever relevant to the delivery of services specified under this Provider Agreement: a. DD Waiver Service Standards and MF Waiver Service Standards. b. DEPARTMENT/DDSD Accreditation Mandate Policies. c. Policies and Procedures for Centralized Admission and Discharge Process for New Mexicans with Disabilities. d. Policies for Behavior Support Service Provisions. e. Rights of Individuals with Developmental Disabilities living in the Community, 7.26.3 NMAC. f. Service Plans for Individuals with Developmental Disability Community Programs, 7.26.5 NMAC.			

g. Requirement for Developmental Disability Community Programs, 7.26.6 NMAC. h. DEPARTMENT Client Complaint Procedures, 7.26.4 NMAC. i. Individual Transition Planning Process, 7.26.7 NMAC. j. Dispute Resolution Process, 7.26.8 NMAC. k. DEPARTMENT/DDSD Training Policies and Procedures. l. Fair Labor Standards Act. m. New Mexico Nursing Practice Act and New Mexico Board of Nursing requirements governing certified medication aides and administration of medications, 16.12.5 NMAC. n. Incident Reporting and Investigation Requirements for Providers of Community Based Services, 7.14.3 NMAC, and DHI/DEPARTMENT Incident Management System Policies and Procedures. o. DHI/DEPARTMENT Statewide Mortality Review Policy and Procedures. p. Caregivers Criminal History Screening Requirements, 7.1.9 NMAC. q. Quality Management System and Review Requirements for Providers of Community Based Services, 7.1.13 NMAC. r. All Medicaid Regulations of the Medical Assistance Division of the HS D. s. Health Insurance Portability and Accountability Act (HIPAA). t. DEPARTMENT Sanctions Policy. u. All other regulations, standards, policies and procedures, guidelines and interpretive memoranda of the DDSD and the DHI of the DEPARTMENT.			

Tag #1A08.2 Administrative Case File: Healthcare Requirements & Follow-up	Condition of Participation Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 3 Safeguards: 3.1 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 according to the following: 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,	After an analysis of the evidence, it has been determined there is a significant potential for a negative outcome to occur. Based on record review, the Agency did not provide documentation of annual physical examinations and/or other examinations as specified by a licensed physician for 7 of 24 individuals receiving Living Care Arrangements and Community Inclusion. Review of the administrative individual case files revealed the following items were not found, incomplete, and/or not current: ***Living Care Arrangements / Community Inclusion (Individuals Receiving Multiple Services):*** **Annual Physical:** • Not Found (#15, 20) • Not Current (#9) **Annual Physical *(LCA Only)*:** • Not Found (#4, 18) **Annual Dental Exam:** • Individual #14 - As indicated by collateral documentation reviewed, the exam was not found. Per the DDSD file matrix, Dental Exams are to be conducted annually. **Psychiatry:** • Individual #1 - As indicated by collateral documentation reviewed, exam was completed on 6/13/2023. Follow-up was to be completed in 3 months. No evidence of follow-up 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?)*: →	

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 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 20 Provider Documentation and Client Records: 20.2 Client Record 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. 7. All records pertaining to JCMs must be retained permanently and must be made available to DDSD upon request, upon the termination or expiration of a provider agreement, or upon provider withdrawal from services. **20.5.4 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. Requirements for the *Health Passport* and *Physician Consultation* form are: 1. The Case Manager and Primary and Secondary Provider Agencies must communicate critical information to each other and will keep all required sections of Therap updated in order to have a current and thorough *Health Passport* and *Physician Consultation* Form available at all times. Required sections of Therap include the IDF, Diagnoses, and Medication History. 2. The Primary and Secondary Provider Agencies must ensure that a current copy of the *Health Passport* and *Physician Consultation* forms are printed and available at all service delivery sites. Both forms must be reprinted and placed at all service delivery sites each time the e-CHAT is updated for any reason and whenever there is a change to contact information contained in the IDF. 3. Primary and Secondary Provider Agencies must assure that the current *Health Passport* and *Physician Consultation* form accompany each person when taken by the provider to a medical appointment, urgent care, emergency room, or are admitted to a hospital or nursing home. (If the person is taken by a family member or guardian, the *Health Passport* and *Physician Consultation* form must be provided to them.)			

4. The Physician Consultation form must be reviewed, and any orders or changes must be noted and processed as needed by the provider within 24 hours. 5. Provider Agencies must document that the *Health Passport* and *Physician Consultation* form and Advanced Healthcare Directives were delivered to the treating healthcare professional by one of the following means: a. document delivery using the *Appointments Results* section in *Therap Health Tracking Appointments*; and b. scan the signed *Physician Consultation Form* and any provided follow-up documentation into Therap after the person returns from the healthcare visit. **Chapter 13 Nursing Services: 13.2.3 General Requirements Related to Orders, Implementation, and Oversight** 1. Each person has a licensed primary care practitioner and receives an annual physical examination, dental care and specialized medical/behavioral care as needed. PPN communicate with providers regarding the person as needed. 2. Orders from licensed healthcare providers are implemented promptly and carried out until discontinued. a. The nurse will contact the ordering or on call practitioner as soon as possible, or within three business days, if the order cannot be implemented due to the person's or guardian's refusal or due to other issues delaying implementation of the order. The nurse must clearly document the issues and all attempts to resolve the problems with all involved parties. b. Based on prudent nursing practice, if a nurse determines to hold a practitioner's order, they are required to immediately document the circumstances and rationale for this decision and to notify			

the ordering or on call practitioner as soon as possible, but no later than the next business day. c. If the person resides with their biological family, and there are no nursing services budgeted, the family is responsible for implementation or follow up on all orders from all providers. Refer to Chapter 13.3 Adult Nursing Services.			

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, 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 September and October 2023. Based on record review, 9 of 15 individuals had Medication Administration Records (MAR), which contained missing medications entries and/or other errors: Individual #2 October 2023 No Physician's Orders were found for medications listed on the Medication Administration Records for the following medications: • Cyanocobalamin 1000mg • Trintellix 10 mg • Vitamin B12 1000mg • Zinc 50 mg Individual #4 September 2023 As indicated by the Medication Administration Records the individual is to take Mirtazapine 7.5 mg (1 time daily). According to the Physician's Orders, Mirtazapine 15mg is to be taken 1 time daily Medication Administration Record and Physician's Orders do not match. Individual #6 September 2023 Medication Administration Records contained missing entries. No	**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	documentation found indicating reason for missing entries: • Adapalene 0.1% (1 time daily) – Blank 9/30 (8:00 PM) • Fungi-nail pen (1 time daily) – Blank 9/30 (8:00 PM) • Panoxyl 10% (1 time daily) – Blank 9/30 (8:00 PM) • Risperidone 0.25mg (1 time daily) – Blank 9/30 (8:00 PM) • Risperidone 0.5mg (1 time daily) – Blank 9/30 (8:00 PM) Individual #7 September 2023 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Docusate Sodium 100 mg (2 times daily) – Blank 9/29 (8:00 PM) • Famotidine 20mg (2 times daily) – Blank 9/29 (8:00 PM) • Topiramate 25mg (1 time daily) - Blank 9/29 (8:00 PM) • Trazodone HCL 100mg (1 time daily) – Blank 9/29 (8:00 PM) As indicated by the Medication Administration Records the individual is to take Alive Women's Gummy Vitamin 120mcg-37.5mg (1 time daily). According to the Physician's Orders, Alive Women's Gummy Vitamin 200mcg-37.5mg is to be taken 1 time daily. Medication Administration		

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:	Record and Physician's Orders do not match. As indicated by the Medication Administration Records the individual is to take Levonorgestrel-Eth Estradiol 200mg (1 time daily). According to the Physician's Orders, Levonorgestrel-Eth Estradiol 0.1-20mg-mcg tablet to be taken 1 time daily. Medication Administration Record and Physician's Orders do not match. October 2023 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Propranolol HCL 10 mg Tablet (3 times daily) – Blank 10/15, 16 (12:00 pm) Individual #8 September 2023 As indicated by the Medication Administration Records the individual is to take Levothyroxine Sodium 50mcg (1 time daily on Monday, Tuesday, Wednesday, Thursday, Friday, and Saturday). According to the Physician's Orders, Levothyroxine Sodium 50mcg is to be taken 1 time daily. Medication Administration Record and Physician's Orders do not match. No Physician's Orders were found for medications listed on the Medication Administration Records for the following medications: • Levothyroxine Sodium 25 mcg Tablet (Every Sunday) Individual #12 September 2023 No Physician's Orders were found for medications listed on the Medication		

➢ 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.	Administration Records for the following medications: • Digest Plus • Multivitamin Powder • Natural Calm Powder October 2023 As indicated by the Medication Administration Records the individual is to take Fish Oil 1,000 mg Capsule (2 times daily). According to the Medication Label / Package, Fish Oil 700mg is to be taken 2 times daily. Medication Administration Record and the Medication Label / Package do not match. Individual #18 September 2023 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Baclofen 20mg (1 time daily) – Blank 9/25, 30 (8:00 AM) • Cyclobenzaprine HCL 10mg, (1 time daily) – Blank 9/29, 30 (8:00 PM) • Vitamin D3 50mcg, (1 time daily) - Blank 9/30 (8:00 AM) Individual #21 September 2023 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Clotrimazole 1% (2 times daily) – Blank 9/30 (8:00 PM)		

	• Diaper Rash Cream (3 times daily) – Blank 9/30 (8:00 PM) • Gabapentin (Neurontin) 600mg (1 time daily) – Blank 9/30 (8:00 PM) • Hydrocortisone 2.5% (2 times daily) – Blank 9/30 (8:00 PM) • Midodrine HCL 5mg (3 times daily) – Blank 9/30 (8:00 PM) • Nystop Topical Powder (4 times daily) – Blank 9/28, 30 (8:00 PM) • Potassium Chloride 10 MEQ (2 times daily) – Blank 9/30 (8:00 PM) • Trazodone HCL 100 MG (1 time daily) – Blank 9/30 (8:00 PM) No Physician's Orders were found for medications listed on the Medication Administration Records for the following medications: • Furosemide 40mg (1 time daily) Individual #24 September 2023 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Lamotrigine 25mg tablet (2 times daily) – Blank 9/29 (8:00 PM) • Oxcarbazepine 300mg/5 ml suspension (2 times daily) – Blank 9/29 (8:00 PM) • Quetiapine Fumarate 50mg tablet (1 time daily) – Blank 9/29 (8:00 PM)		

Tag # 1A09.0 Medication Delivery Routine 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. 4. Provider Agencies must configure and use the MAR when assisting with medication.	Medication Administration Records (MAR) were reviewed for the months of September and October 2023. Based on record review, 2 of 15 individuals had Medication Administration Records (MAR), which contained missing medications entries and/or other errors: Individual #12 October 2023 Medication Administration Records did not contain the strength of the medication which is to be given: • Vitamin D3 (1 time daily) Individual#14 September 2023 Medication Administration Records did not contain the diagnosis for which the medication is prescribed: • Biotin 5mg (1 time daily) • Dycylomine 10 mg (3 times daily) Medication Administration Records did not contain the dosage for the following medications: • Ferrous Sulfate 10mg	**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?)*: →	

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 # 1A09.1 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 September and October 2023. Based on record review, 11 of 15 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #1 October 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. • Bismatrol 262mg/15mL (PRN) • Deep Sea Nasal Spray 0.65% (PRN) • Geri-Tussin 100.0mg/5mL (PRN) • Loperamide 2.0mg (PRN) • Loratadine 10.mg (PRN) • MI Acid oral suspension (PRN) • Milk of Magnesia 400.0mg/mL (PRN) • Off Deep Woods Dry 25% (PRN) • Risperidone 2mg (PRN) • Sunscreen SPF 30 (PRN) • Vaseline (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	• Vicks VapoRub (PRN) Individual #2 September 2023 No Physician's Orders were found for medications listed on the Medication Administration Records for the following medications: • Tylenol Sinus 325/200/5mg (PRN) October 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 325mg (PRN) • Artificial Tears (PRN) • Deep Sea Nasal Spray 0.65% (PRN) • Geri-Tussin 100mg/5mL (PRN) • Ibuprofen 200mg (PRN) • Loperamide 2mg (PRN) • MI Acid oral suspension (PRN) • Mucus Relief ER 600mg (PRN) • Off Deep Woods Dry 25% (PRN) • Sunscreen SPF 30 (PRN) • Vicks VapoRub (PRN) Individual #4 September 2023		

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:	No Physician's Orders were found for medications listed on the Medication Administration Records for the following medications: • Fluocinonide Cream 0.05% (PRN) October 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 7.5mg (PRN) • Diphenhydramine 25mg (PRN) • Eye Drops (PRN) • Fluocinonide Cream 0.05% (PRN) • Hydrocortisone cream 1% (PRN) • Icy Hot Cream 30-10% (PRN) • Milk of Magnesia 400mg/mL (PRN) • Nystatin Cream/oral 1000 (PRN) • Tums Chewable 200/500mg (PRN) • Tussin DM Syrup 10-100mg/5mL (PRN) Individual #5 October 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. • Artificial Tears (PRN) • Saline Mist Nose Spray 0.65% (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.	• Geri-Tussin 100.0mg/5mL (PRN) • Loperamide 2.0mg (PRN) • Loratadine 10mg (PRN) • MI Acid oral suspension (PRN) • Off Deep Woods Dry 25% (PRN) • Aloe Vera Gel (PRN) • Sunscreen SPF 30 (PRN) • Acetaminophen 325 mg (PRN) • Vicks VapoRub (PRN) Individual #7 September 2023 Physician's Orders indicated the following medication were to be given. The following Medications were not documented on the Medication Administration Records: • Guaifenesin 600mg (PRN) No Physician's Orders were found for medications listed on the Medication Administration Records for the following medications: • Antacid Liquid 200-200-20mg/5mL (PRN) • Geri-Lanta (PRN) • Lorazepam 2 mg (PRN) • Meclizine HCL 12.5 mg (PRN) • Mucinex 600 mg (PRN) • NYAMYC Powder 100000 unit/gram (PRN) • Propranolol HCL 20mg (PRN)		

	October 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. • Aloe Vera Gel (PRN) • Acetaminophen 325 mg (PRN) Individual #8 September 2023 As indicated by the Medication Administration Records the individual is to take Acetaminophen 500 mg 1-2 tablets (PRN). According to the Physician's Orders, Acetaminophen 325mg 2 tablets is to be taken as needed Medication Administration Record and Physician's Orders do not match. Physician's Orders indicated the following medication were to be given. The following Medications were not documented on the Medication Administration Records: • Ibuprofen 200mg (PRN) • Artificial Tears (PRN) • Loratadine 10mg (PRN) No Physician's Orders were found for medications listed on the Medication Administration Records for the following medications: • Abreva 10% (PRN) • Diphenhist 25 mg (PRN) Individual #12 September 2023		

	No Physician's Orders were found for medications listed on the Medication Administration Records for the following medications: • Debrox 6.5% (PRN) October 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. • Artificial Tears (PRN) • Bismuth 262mg/15mL suspension (PRN) • Deep Sea Nasal Spray 0.65% (PRN) • Geri-Tussin 100mg/5mL (PRN) • Ibuprofen 200mg (PRN) • Loperamide 2mg (PRN) • Loratadine 10mg (PRN) • MI Acid oral suspension (PRN) • Milk of Magnesia 400mg/mL (PRN) • Aloe Vera Gel (PRN) • Petroleum Jelly (PRN) • Children's Allergy Relief 12.5mg/ml (PRN) Individual #14 October 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 325 mg (PRN) • Aloe Vera Gel (PRN) • Artificial Tears (PRN) • Geri-Tussin 100.0mg/5mL (PRN) • Loratadine 10mg (PRN) • MI Acid oral suspension (PRN) • Off Deep Woods Dry 25% (PRN) • Saline Mist Nose Spray 0.65% (PRN) • Sunscreen SPF 30 (PRN) • Vicks VapoRub (PRN) Individual #18 October 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 325mg (PRN) • Artificial Tears (PRN) • Deep Sea Nasal Spray 0.65% (PRN) • Geri-Tussin 100mg/5mL (PRN) • Loperamide 2mg (PRN) • Mucus Relief ER 600 mg (PRN) • Off Deep Woods Dry 25% (PRN) • Sunscreen SPF 30 (PRN)		

	• Vicks VapoRub (PRN) Individual #23 October 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 325mg (PRN) • Artificial Tears (PRN) • Deep Sea Nasal Spray 0.65% (PRN) • Ketotifen Fum 0.025% (PRN) • Loratadine 10 mg (PRN) • Insect Repellent (PRN) • Senna Laxative 8.6 mg (PRN) • Sunscreen SPF 30 (PRN) • Vicks VapoRub (PRN) Individual #24 October 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 325mg (PRN) • Aloe Vera Gel (PRN) • Artificial Tears (PRN) • Bismuth 262mg/15mL suspension (PRN) • Deep Sea Nasal Spray 0.65% (PRN)		

	• Geri-Tussin 100mg/5mL (PRN) • Loperamide 2mg (PRN) • Loratadine 10mg (PRN) • MI Acid oral suspension (PRN) • Milk of Magnesia 400mg/mL (PRN) • Petroleum Jelly (PRN) Individual #25 October 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. • Aloe Vera Gel (PRN) • Artificial Tears (PRN) • Insect Repellent (PRN) • Loperamide 2mg (PRN) • Loratadine 10mg (PRN) • MI Acid oral suspension (PRN) • Milk of Magnesia 400mg/mL (PRN) • Petroleum Jelly (PRN) • Pink Bismuth 262mg/15mL suspension (PRN) • Saline Mist Nose Spray 0.65% (PRN)		

	As indicated by the Medication Administration Records the individual is to take Albuterol Sulfate HFA 108mcg (1 - 2 puff as needed). According to the Medication Label / Package, Albuterol Sulfate HFA 90mcg is to be taken 1 - 2 times as needed Medication Administration Record and the Medication Label / Package do not match.		

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. 4. Provider Agencies must configure and use the MAR when assisting with medication.	Medication Administration Records (MAR) were reviewed for the months of September and October 2023. Based on record review, 4 of 15 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #1 October 2023 Medication Administration Records did not contain the number of doses that may be used in a 24-hour period: • Hydroxyzine Pam 50mg (PRN) Individual #7 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Propranolol HCL 20mg – PRN – 9/25 (given 1 time) Individual #12 September 2023 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Bismuth 262mg/15ml– PRN – 9/26 (given 1 time) • Children's Allergy Relief 12.5mg/5ml – PRN – 9/22, 24, 25, 27 (given 1 time) October 2023 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Children's Allergy Relief 12.5mg/5ml – PRN – 10/8 - 13, 15 - 18 (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?)*: →	

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	Individual #25 October 2023 Medication Administration Records did not contain the number of doses that may be used in a 24-hour period: • Albuterol Sulfate 0.083% (PRN) • Albuterol Sulfate HFA 108mcg (PRN)		

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 and interview, the Agency did not maintain documentation of PRN authorization as required by standard for 3 of 15 Individuals. Individual #7 September 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: • Propranolol HCL – PRN – 9/25 (given 1 time) Individual #12 September 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: • Children's Allergy Relief 12.5mg/5ml – PRN – 9/22, 24 (given 1 time) Individual #14 September 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: • Ibuprofen 200mg – PRN – 9/20 (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 2. 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,	After an analysis of the evidence, it has been determined the following finding resulted in a negative outcome. Based on record review, the Agency did not maintain the required documentation in the Individuals Agency Record as required by standard for 13 of 24 individuals. 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, 3, 7, 8, 10, 14, 15, 22) • Did not contain Emergency Contact Information (#1, 3, 5, 7, 14, 15, 22) • Did not contain Health and Safety Risk Factors (#1, 3) • Did not Contain Medical Diagnosis (#2, 3) • Did not contain Guardianship/Healthcare Decision Maker (#1, 2, 3, 8, 22) **Electronic Comprehensive Health Assessment Tool (eCHAT):** • Not approved within 3-days of being completed (#23) **Comprehensive Aspiration Risk Management Plan*:*** • Not Found (#2, 13) **Health Care Plans:** ***Paralysis Present:***	**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?)*: →	

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 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.	• Individual #18 – Per the Electronic Comprehensive Health Assessment Tool, the individual is required to have a plan. No evidence of a plan found. **Medical Emergency Response Plans:** ***Allergies:*** • Individual #7 – As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found. • Individual #17 – As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found. ***Paralysis Present:*** • Individual #18 – Per the Electronic Comprehensive Health Assessment Tool, the individual is required to have a plan. No evidence of a plan found. ***Risk for Falls:*** • Individual #22 – As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found.		

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 # 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 18 of 24 individuals. Review of the Agency individual case files revealed the following items were not found and/or incomplete: **Grievance/Complaint Procedure Acknowledgement:** • Not found (#1, 2, 3, 4, 6, 7, 8, 9, 12, 13, 15, 16, 17, 18, 20, 21, 23, 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?)*: →	

Tag # 1A31.2 Human Right Committee Composition	Standard Level Deficiency		
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 3 Safeguards: 3.3 *Human Rights Committee:*** Human Rights Committees (HRC) exist to protect the rights and freedoms of all waiver participants through the review of proposed restrictions to a person's rights based on a documented health and safety concern of a severe nature (e.g., a serious, significant, credible threat or act of harm against self, others, or property) . HRCs monitor the implementation of certain time-limited restrictive interventions designed to protect a waiver participant and/or the community from harm. An HRC may also serve other functions as appropriate, such as the review of agency policies on the use of emergency physical restraint or sexuality if desired. HRCs are required for all Living Supports (Supported Living, Family Living, Intensive Medical Living Services), Customized Community Supports (CCS) and Community Integrated Employment (CIE) Provider Agencies. 1. HRC membership must include: a. at least one member with a diagnosis of I/DD; b. a parent or guardian of a person with I/DD; c. a health care services professional (e.g., a physician or nurse); and d. a member from the community at large that is not associated (past or present) with DD Waiver services. 2. Committee members must abide by HIPAA; 3. All committee members will receive training on Abuse, Neglect and Exploitation (ANE) Awareness, Human Rights, HRC requirements, and other pertinent DD Waiver Service Standards prior to their voting participation on the HRC. A committee member trained by the Bureau of	Based on record review the Agency did not ensure the correct composition of the human rights committee. **Review of Agency's HRC committee found the following were not members of the HRC:** • *a member from the community at large that is not associated (past or present) with DD Waiver services.*	**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?)*: →	

Behavioral Supports (BBS) may conduct training for other HRC members, with prior approval from BBS; 4. HRCs will appoint an HRC chair. Each committee chair shall be appointed to a two-year term. Each chair may serve only two consecutive two-year terms at a time; 5. While agencies may have an intra-agency HRC, meeting the HRC requirement by being a part of an interagency committee is also highly encouraged.			

Tag # 1A33 Board of Pharmacy: Med. Storage	Standard Level Deficiency		
New Mexico Board of Pharmacy Model Custodial Drug Procedures Manual **E. Medication Storage:** 1. Prescription drugs will be stored in a locked cabinet and the key will be in the care of the administrator or designee. 2. Drugs to be taken by mouth will be separate from all other dosage forms. 3. A locked compartment will be available in the refrigerator for those items labeled "Keep in Refrigerator." The temperature will be kept in the 36°F - 46°F range. An accurate thermometer will be kept in the refrigerator to verify temperature. 4. Separate compartments are required for each resident's medication. 5. All medication will be stored according to their individual requirement or in the absence of temperature and humidity requirements, controlled room temperature (68-77°F) and protected from light. Storage requirements are in effect 24 hours a day. 6. Medication no longer in use, unwanted, outdated, or adulterated will be placed in a quarantine area in the locked medication cabinet and held for destruction by the consultant pharmacist. **8. References** A. Adequate drug references shall be available for facility staff **H. Controlled Substances (Perpetual Count Requirement)** 1. Separate accountability or proof-of-use sheets shall be maintained, for each controlled substance, indicating the following information: a. date b. time administered c. name of patient d. dose	Based on record review and observation, the Agency did not ensure proper storage of medication for 6 of 15 individuals. Observation included: Separate compartments were NOT kept for each individual living in the home. (Individual #2, 5, 14, 18, and 23) Individual #1 • Risperidone 2mg tablet: expired 8/2023. Expired medication was not kept separate from other medications as required by the Board of Pharmacy Procedures. .	**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?)*: →	

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

(iii) Name, address and phone number of pharmacy; **(iv)** Prescription number; **(v)** Name of the drug and quantity; **(vi)** Strength of drug and quantity; **(vii)** Directions for use, route of administration; **(viii)** Date of prescription (date of refill in case of a prescription renewal); **(ix)** Expiration date where applicable: The dispenser shall place on the label a suitable beyond-use date to limit the patient's use of the medication. Such beyond-use date shall be not later than (a) the expiration date on the manufacturer's container, or (b) one year from the date the drug is dispensed, whichever is earlier; **(x)** Auxiliary labels where applicable; **(xi)** The Manufacturer's name; **(xii)** State of the art drug delivery systems using unit of use packaging require items i and ii above, provided that any additional information is readily available at the nursing station. Developmental Disabilities Waiver Service Standards 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: 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;			

Tag # 1A33.1 Board of Pharmacy - License	Standard Level Deficiency		
New Mexico Board of Pharmacy Model Custodial Drug Procedures Manual **Display of License and Inspection Reports** The following are required to be publicly displayed: • Current Custodial Drug Permit from the NM Board of Pharmacy • Current registration from the consultant pharmacist • Current NM Board of Pharmacy Inspection Report Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 16 Qualified Provider Agencies:** **16.5 Board of Pharmacy:** All DD Waiver Provider Agencies with service settings where medication administration / assistance to two or more unrelated individuals occurs must be licensed by the Board of Pharmacy and must follow all Board of Pharmacy regulations related to medication delivery including but not limited to: 1. pharmacy licensing; 2. medication delivery; 3. proper documentation and storage of medication; 4. use of a pharmacy policy manual; and 5. holding an active contract with a Pharmacy Consultant.	Based on observation, the Agency did not provide the current Custodial Drug Permit from the New Mexico Board of Pharmacy, the current registration from the Consultant Pharmacist, or the current New Mexico Board of Pharmacy Inspection Report for 2 of 17 residences: **Individual Residence:** • Current Custodial Drug Permit from the NM Board of Pharmacy with the current address of the residence (#7, 12, 24) *Note: The following Individuals share a residence:* • #1, 25 • #2, 18 • #4, 22 • #5, 14 • #6, 21 • #12, 24	**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 6 of 8 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 #9 - Not Found. • Individual #13 - Not Found. • Individual #15 - Not Found. • Individual #17 - Not Found. **Monthly Consultation with the Direct Support Provider and the person receiving services:** • Individual #3 - None found for 10/2022 - 9/2023. • Individual #8 - None found for 2/2023 - 5/2023. • Individual #9 - None found for 11/2022 - 3/2023 and 6/2023. • Individual #13 - None found for 10/2022 - 2/2023, and 5/2023. • Individual #15 - None found for 12/2022, 2/2023, and 3/2023. • Individual #17 - None found for 10/2022 - 3/2023.	**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 record review and / or observation, the Agency did not ensure that each individuals' residence met all requirements within the standard for 11 of 17 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:
	
	• Carbon monoxide detectors (#12, 24)
	
	• General-purpose first aid kit (#12, 24)
	
	• Poison Control Phone Number (#4, 22)
	
	• Water temperature in home exceeds safe temperature (110^0 F):
	  • Water temperature in home measured 146.7^0 F (#1, 25)
	
	  • Water temperature in home measured 124.5^0 F (#2, 18)
	
	  • Water temperature in home measured 133.3^0 F (#4, 22, 23)
	
	  • Water temperature in home measured 114.6^0 F (#5, 14)
	
	  • Water temperature in home measured 152.7^0 F (#6, 21)
	
	  • Water temperature in home measured 129.9^0 F (#7)
		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 113.9^0 F (#12, 24) *Note: The following Individuals share a residence:* • #1, 25 • #2, 18 • #4, 22 • #5, 14 • #6, 21 • #12, 24 **Family Living Requirements:** • Battery operated or electric smoke detectors or a sprinkler system (#15) • Carbon monoxide detectors (#4, 22) • Poison Control Phone Number (#15) • Water temperature in home exceeds safe temperature (110^0 F) • Water temperature in home measured 119.5^0 F (#4, 22) • Water temperature in home measured 133^0 F (#15) • Water temperature in home measured 141.3^0 F (#19) • Water temperature in home measured 117.5^0 F (#17)		

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 1 of 11 individuals. Individual #17 July 2023 • The Agency billed 26 units of Customized Community Supports (H2021 HB-U1 from on 7/2/2023. Documentation did not contain the required element(s) on 7/2/2023. Documentation received accounted for 0 units. The required element(s) were not met: • Services were provided concurrently with another service.	**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.			

Tag # LS26 Supported 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 Supported Living Services for 4 of 14 individuals. Individual #4 July 2023 • The Agency billed 1 unit of Supported Living (T2016 HB - U7) on 7/17/2023. Documentation received accounted for .5 units. As indicated by the DDW Standards, at least 12 hours in a 24-hour period must be provided in order to bill a complete unit. Documentation received accounted for 5 hours, which is less than the required amount. Individual #5 September 2023 • The Agency billed 1 unit of Supported Living (T2016 HB-U6) on 9/14/2023. Documentation received accounted for .5 units. As indicated by the DDW Standards, at least 12 hours in a 24-hour period must be provided in order to bill a complete unit. Documentation received accounted for 3 hours, which is less than the required amount. Individual #6 September 2023 • The Agency billed 1 unit of Supported Living (T2016 HB-U5) on 9/11/2023. Documentation received accounted for .5 units. As indicated by the DDW Standards, at least 12 hours in a 24-hour period must be provided in order to bill a complete unit. Documentation received accounted for 9 hours, which is less than the required amount.	**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.	Individual #18 August 2023 • The Agency billed 1 unit of Supported Living (T2016 HB-U7) on 8/23/2023. Documentation received accounted for .5 units. As indicated by the DDW Standards, at least 12 hours in a 24-hour period must be provided in order to bill a complete unit. Documentation received accounted for 9 hours, which is less than the required amount.		

Tag #IH32 Customized In-Home 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 Intensive Medical Living Services for 1 of 2 individuals. Individual #20 August 2023 • The Agency billed 15 units of Customized In-Home Supports (S5125 HB - UA) on 8/14/2023. Documentation did not contain the required element(s) on 8/14/2023. Documentation received accounted for 0 units. The required element was not met: • A description of what occurred during the encounter or service interval. • The Agency billed 16 units of Customized In-Home Supports (S5125 HB - UA) on 8/18/2023. Documentation did not contain the required element(s) on 8/18/2023. Documentation received accounted for 0 units. The required element was not met: • A description of what occurred during the encounter or service interval. • The Agency billed 16 units of Customized In-Home Supports (S5125 HB-UA) on 8/19/2023. Documentation did not contain the required element(s) on 8/19/2023. Documentation received accounted for 0 units. The required element was not met: • A description of what occurred during the encounter or service interval.	**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. **21.4 Electronic Visit Verification:** Section 12006(a) of the 21st Century Cures Act (the Cures Act) requires that states implement Electronic Visit Verification (EVV) for all Medicaid services under the umbrella of personal care and home health care that require an in-home visit by a provider. EVV is a technological solution used to electronically verify whether providers delivered or rendered services as billed. Personal Care Services are services supporting Activities of Daily Living (ADLs) or services supporting both ADLs and Instrumental Activities of Daily Living (IADLs). Home Health Care Services (HHCS) are services providing nursing services and/or home health aide services. The Cures Act allows states to implement EVV in a phased approach starting with the services meeting federal guidelines for PCS and later HHCS. The use of the state approved EVV system does not replace other standards requirements. EVV system has potential for benefits that may include: a. Improved practices inherent in the use of EVV. b. Centralized, real-time monitoring and comprehensive reporting on services provided. c. Use of EVV data to identify delivery issues and make care delivery more efficient. d. Improving program integrity and higher quality of services. e. Improving risk management and fraud protection. f. Secure, HIPAA compliant automated claims. The EVV system verifies the:			

a. **Type** of service performed. b. **Individual receiving** the service. c. **Date** of service. d. **Location** of service delivery. e. **Individual providing** the service. f. **Time** the service begins and ends. The state supplies agencies with a single approved EVV system that must be used. Effective January 1, 2021, DD Waiver providers of CIHS and Respite are required to implement the use of state approved EVV system. As home health care services are phased in according to federal and state requirements, additional services may require the use of EVV.			

NEW MEXICO
Department of Health
Division of Health Improvement

MICHELLE LUJAN GRISHAM
Governor

PATRICK M. ALLEN
Cabinet Secretary

Date: January 23, 2024

To: Emad Elmaoued, Director

Provider: ADID Care INC
Address: 5115 Copper Avenue NE
State/Zip: Albuquerque, New Mexico 87408

E-mail Address: emad@adidcare.com

Region: Metro and Northeast
Survey Date: October 16 - 27, 2023

Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: Supported Living, Family Living, Customized In-Home Supports, Customized Community Supports

Survey Type: Routine

Dear Mr. Elmaoued,

The Division of Health Improvement Quality Management Bureau received and reviewed the documents you submitted for your Plan of Correction. Your Plan of Correction is not closed.

Your Plan of Correction will be considered for closure when a Verification survey confirms that you have corrected all survey deficiencies and sustained all corrections.

The Quality Management Bureau will need to conduct a verification survey to ensure previously cited deficiencies have been corrected and that systemic Quality Improvement and Quality Assurance processes have been effective at sustaining corrections.

If the Verification survey determines survey deficiencies have been corrected and corrective measures have effectively maintained compliance with DDW Standards, your Plan of Correction will be considered for closure.

If the Verification survey identifies repeat deficiencies, the Plan of Correction process will continue and your case may be referred to the Internal Review Committee for discussion of possible civil monetary penalties possible monetary fines and/or other sanctions.

Thank you for your cooperation with the Plan of Correction process.

Sincerely,

Monica Valdez, BS

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

Q.24.2.DDW.D4455.2/5.001.RTN.07.23.024