

Date: May 10, 2022

To: Melissa McCue, Executive Director

Provider: Mandy's Special Farm dba Mandy's  
Address: 3501 Campus Blvd. NE  
State/Zip: Albuquerque, New Mexico 87106

E-mail Address: [melissa@mandysfarm.org](mailto:melissa@mandysfarm.org)

Board Chair: Ruthie Robbins, Board President  
E-Mail Address: [drmrobb@att.net](mailto:drmrobb@att.net)

Region: Metro  
Survey Date: April 4 - 15, 2022

Program Surveyed: Developmental Disabilities Waiver

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

Survey Type: Routine

Team Leader: Sally Rel, MS, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Team Members: LeiLani Nava, MPH, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; Heather Driscoll, AA, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; Jamie Pond, BS, QMB Staff Manager, Division of Health Improvement/Quality Management Bureau; Alyssa Swisher, IMB/QMB Nurse Investigator/Surveyor, Division of Health Improvement/Quality Management Bureau.

Dear Ms. Melissa McCue;

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.

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

**Partial Compliance with Standard Level Tags and Conditions of Participation Level Tags:** This determination is based on noncompliance with one to five (1 – 5) Condition of Participation Level Tags (*refer to Attachment D for details*). The attached QMB Report of Findings indicates Standard Level and Condition of Participation Level deficiencies identified and requires completion and implementation of a Plan of Correction.

**DIVISION OF HEALTH IMPROVEMENT**

5301 Central Avenue NE, Suite 400 • Albuquerque, New Mexico • 87108  
(505) 222-8623 • FAX: (505) 222-8661 • <https://nmhealth.org/about/dhi>

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Survey Report #: Q 22.4.DDW.32408382.5.RTN.01.22.130



The following tags are identified as Condition of Participation Level:

- Tag # 1A08.2 Administrative Case File: Healthcare Requirements & Follow-up
- Tag # 1A09 Medication Delivery Routine Medication Administration
- 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 # 1A43.1 General Events Reporting: Individual Reporting
- Tag # 1A09.0 Medication Delivery Routine Medication Administration
- Tag # 1A09.1 Medication Delivery PRN Medication Administration
- Tag # 1A09.1.0 Medication Delivery PRN Medication Administration
- Tag # IS30 Customized Community 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, Attention: Monica Valdez, Plan of Correction Coordinator in any of the following ways:**
  - a. Electronically at [MonicaE.Valdez@state.nm.us](mailto:MonicaE.Valdez@state.nm.us) (*preferred method*)
  - b. Fax to 505-222-8661, or
  - c. Mail to POC Coordinator, 5301 Central Ave NE Suite 400, Albuquerque, New Mexico 87108
- 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.

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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  
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@state.nm.us](mailto:Lisa.medina-lujan@state.nm.us))

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  
5301 Central Ave NE Suite #400  
Albuquerque, NM 87108  
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@state.nm.us](mailto:MonicaE.Valdez@state.nm.us) if you have questions about the Report of Findings or Plan of Correction. Thank you for your cooperation and for the work you perform.

Sincerely,

*Sally Rel, MS*

Sally Rel, MS  
Team Lead/Healthcare Surveyor  
Division of Health Improvement  
Quality Management Bureau

**Survey Process Employed:**

Administrative Review Start Date: April 4, 2022

Contact: **Mandy's Special Farm dba Mandy's**  
Melissa McCue, Executive Director

**DOH/DHI/QMB**  
Sally Rel, MS, Team Lead/Healthcare Surveyor

On-site Entrance Conference Date: April 4, 2022

Present: **Mandy's Special Farm dba Mandy's**  
Melissa McCue, Executive Director  
Libby Pacheco, Quality Assurance Coordinator  
Stephanie Aragon, Training Coordinator  
Christopher Love, CIE Program Coordinator  
Marion Biggs Jackson, Operations Coordinator  
Bernadette Garcia, Service Coordinator / Program Director

**DOH/DHI/QMB**  
Sally Rel, MS, Team Lead/ Healthcare Surveyor  
Amanda Castaneda-Holguin, MPA, Healthcare Surveyor Supervisor  
LeiLani Nava, MPH, Healthcare Surveyor  
Heather Driscoll, AA, Healthcare Surveyor  
Jamie Pond, BS, QMB Staff Manager  
Alyssa Swisher, IMB/QMB Nurse Investigator/Surveyor

Exit Conference Date: April 15, 2022

Present: **Mandy's Special Farm dba Mandy's**  
Melissa McCue, Executive Director  
Libby Pacheco, Quality Assurance Coordinator  
Stephanie Aragon, Training Coordinator  
Bernadette Garcia, Service Coordinator / Program Director  
Yvette Trujillo, Healthcare Coordinator

**DOH/DHI/QMB**  
Sally Rel, MS, Team Lead/Healthcare Surveyor  
Amanda Castaneda-Holguin, MPA, Healthcare Surveyor Supervisor  
LeiLani Nava, MPH, Healthcare Surveyor  
Heather Driscoll, AA, Healthcare Surveyor

**DDSD - Metro Regional Office**  
Linda Clark, Assistant Regional Director

Administrative Locations Visited: 0 (Note: No administrative locations visited due to COVID-19 Public Health Emergency)

Total Sample Size: 11  
0 - Jackson Class Members  
11 - Non-Jackson Class Members  
4 - Supported Living  
11 - Customized Community Supports  
3 - Community Integrated Employment

Total Homes Visited	2
❖ Supported Living Homes Visited	2
	<i>Note: The following individuals share a SL residence.</i>
	➤ #9, 11, 12
Persons Served Records Reviewed	11
Persons Served Interviewed	6
Persons Served Observed	4
Persons Served Not Seen and/or Not Available	1 ( <i>Note: 1 Individual was not available during the on-site survey</i> )
Direct Support Personnel Records Reviewed	49
Direct Support Personnel Interviewed	7
Service Coordinator Records Reviewed	2
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
  - Medication Administration Records
  - Medical Emergency Response Plans
  - Therapy Evaluations and Plans
  - Healthcare Documentation Regarding Appointments and Required Follow-Up
  - Other Required Health Information
- Internal Incident Management Reports and System Process / General Events Reports
- Personnel Files, including nursing and subcontracted staff
- Staff Training Records, Including Competency Interviews with Staff
- Agency Policy and Procedure Manual
- Caregiver Criminal History Screening Records
- Consolidated Online Registry/Employee Abuse Registry
- Human Rights Committee Notes and Meeting Minutes
- Evacuation Drills of Residences and Service Locations
- Quality Assurance / Improvement Plan

CC: Distribution List: DOH - Division of Health Improvement  
 DOH - Developmental Disabilities Supports Division  
 DOH - Office of Internal Audit  
 HSD - Medical Assistance Division  
 NM Attorney General's Office

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## 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 DDS 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@state.nm.us](mailto:MonicaE.Valdez@state.nm.us). Requests for technical assistance must be requested through your Regional DDS 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@state.nm.us](mailto:MonicaE.Valdez@state.nm.us) 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 in any of the following ways:
  - a. Electronically at [MonicaE.Valdez@state.nm.us](mailto:MonicaE.Valdez@state.nm.us) (**preferred method**)
  - b. Fax to 505-222-8661, or
  - c. Mail to POC Coordinator, 5301 Central Ave NE Suite 400, Albuquerque, New Mexico 87108
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, as follows.

1. Your internal documents are due within a **maximum** of 45-business days of receipt of your Report of Findings.

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2. It is preferred that you submit your documents via USPS or other carrier (scanned and saved to CD/DVD disc, flash drive, etc.). If documents containing HIPAA Protected Health Information (PHI) documents must be submitted through S-Comm (Therap), Fax or Postal System, do not send PHI directly to NMDOH email accounts. If the documents do not contain protected Health information (PHI) then you may submit your documents electronically scanned and attached to e-mails.
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 completion date and are approved on a case-by-case basis. No changes may be made to your POC or the timeframes for implementation without written approval of the POC Coordinator.**

## Attachment B

### Department of Health, Division of Health Improvement QMB Determination of Compliance Process

The Division of Health Improvement, Quality Management Bureau (QMB) surveys compliance of the Developmental Disabilities Waiver (DDW) standards and other state and federal regulations. For the purpose of the LCA / CI survey the CMS waiver assurances have been grouped into four (4) Service Domains: Plan of Care (ISP Implementation); Qualified Providers; Health, Welfare and Safety; and Administrative Oversight (note that Administrative Oversight listed in this document is not the same as the CMS assurance of Administrative Authority. Used in this context it is related to the agency's operational policies and procedures, Quality Assurance system and Medicaid billing and reimbursement processes.)

The QMB Determination of Compliance process is based on provider compliance or non-compliance with standards and regulations identified during the on-site survey process and as reported in the QMB Report of Findings. All areas reviewed by QMB have been agreed to by DDSD and DHI/QMB and are reflective of CMS requirements. All deficiencies (non-compliance with standards and regulations) are identified and cited as either a Standard level deficiency or a Condition of Participation level deficiency in the QMB Reports of Findings. All deficiencies require corrective action when non-compliance is identified.

Each deficiency in your Report of Findings has been predetermined to be a Standard Level Deficiency, a Condition of Participation Level Deficiency, if below 85% compliance or a non-negotiable Condition of Participation Level Deficiency. Your Agency's overall Compliance Determination is based on a Scope and Severity Scale which takes into account the number of Standard, and Condition Level Tags cited as well as the percentage of Individuals affected in the sample.

#### Conditions of Participation (CoPs)

CoPs are based on the Centers for Medicare and Medicaid Services, Home and Community-Based Waiver required assurances, in addition to the New Mexico Developmental Disability Waiver (DDW) Service Standards. The Division of Health Improvement (DHI), in conjunction with the Developmental Disability Support Division (DDSD), has identified certain deficiencies that have the potential to be a Condition of Participation Level, if the tag falls below 85% compliance based on the number of people affected. Additionally, there are what are called non-negotiable Conditions of Participation, regardless if one person or multiple people are affected. In this context, a CoP is defined as an essential / fundamental regulation or standard, which when out of compliance directly affects the health and welfare of the Individuals served. If no deficiencies within a Tag are at the level of a CoP, it is cited as a Standard Level Deficiency.

***Service Domains and CoPs for Living Care Arrangements and Community Inclusion are as follows:***

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

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

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

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

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

- **1A20** - Direct Support Personnel Training
- **1A22** - Agency Personnel Competency
- **1A37** – Individual Specific Training

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**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 Requirements. (Physical Environment - Supported Living / Family Living / Intensive Medical Living)

## Attachment C

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

#### Introduction:

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

#### Instructions:

1. The Informal Reconsideration of the Finding (IRF) request must be received in writing to the QMB Bureau Chief **within 10 business days** of receipt of the final Report of Findings (**Note: No extensions are granted for the IRF**).
2. The written request for an IRF *must* be completed on the QMB Request for Informal Reconsideration of Finding form available on the QMB website: <https://nmhealth.org/about/dhi/cbp/irf/>
3. The written request for an IRF must specify in detail the request for reconsideration and why the finding is inaccurate.
4. The IRF request must include all supporting documentation or evidence.
5. If you have questions about the IRF process, email the IRF Chairperson, Valerie V. Valdez at [valerie.valdez@state.nm.us](mailto:valerie.valdez@state.nm.us) for assistance.

#### The following limitations apply to the IRF process:

- The written request for an IRF and all supporting evidence must be received within 10 business days.
- Findings based on evidence requested during the survey and not provided may not be subject to reconsideration.
- The supporting documentation must be new evidence not previously reviewed or requested by the survey team.
- Providers must continue to complete their Plan of Correction during the IRF process
- Providers may not request an IRF to challenge the sampling methodology.
- Providers may not request an IRF based on disagreement with the nature of the standard or regulation.
- Providers may not request an IRF to challenge the team composition.
- Providers may not request an IRF to challenge the DHI/QMB determination of compliance or the length of their DDSD provider contract.

A Provider forfeits the right to an IRF if the request is not received within 10 business days of receiving the report and/or does not include all supporting documentation or evidence to show compliance with the standards and regulations.

The IRF Committee will review the request; the Provider will be notified in writing of the ruling; no face-to-face meeting will be conducted.

When a Provider requests that a finding be reconsidered, it does not stop or delay the Plan of Correction process. **Providers must continue to complete the Plan of Correction, including the finding in dispute regardless of the IRF status.** If a finding is removed or modified, it will be noted and removed or modified from the Report of Findings. It should be noted that in some cases a Plan of Correction may be completed prior to the IRF process being completed. The provider will be notified in writing on the decisions of the IRF committee.

## 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%	
<i>“Non-Compliance”</i>						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.
<i>“Partial Compliance with Standard Level tags and Condition of Participation Level Tags”</i>					Any Amount Standard Level Tags, plus 1 to 5 Conditions of Participation Level tags.		
<i>“Partial Compliance with Standard Level tags”</i>			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.			
<i>“Compliance”</i>	Up to 16 Standard Level Tags with 0 to 74% of the individuals in the sample cited in any tag.	17 or more Standard Level Tags with 0 to 49% of the individuals in the sample cited in any tag.					

**Agency:** Mandy's Special Farm dba Mandy's - Metro Region  
**Program:** Developmental Disabilities Waiver  
**Service:** Supported Living, Customized Community Supports, and Community Integrated Employment Services  
**Survey Type:** Routine  
**Survey Date:** April 4 - 15, 2022

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Completion Date
<b>Service Domain: Service Plans: ISP Implementation</b> – Services are delivered in accordance with the service plan, including type, scope, amount, duration and frequency specified in the service plan.			
<b>Tag # 1A08 Administrative Case File (Other Required Documents)</b>	<b>Standard Level Deficiency</b>		
<p>Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019</p> <p><b>Chapter 20: Provider Documentation and Client Records: 20.2</b> 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:</p> <ol style="list-style-type: none"> <li>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.</li> <li>Provider Agencies must have readily accessible records in home and community settings in paper or electronic form. Secure access to electronic records through the Therap web-based system using computers or mobile devices is acceptable.</li> <li>Provider Agencies are responsible for ensuring that all plans created by nurses, RDs, therapists or BSCs are present in all needed</li> </ol>	<p>Based on record review and interview , the Agency did not maintain a complete and confidential case file at the administrative office for 1 of 11 individuals.</p> <p>Review of the Agency administrative individual case files revealed the following items were not found, incomplete, and/or not current:</p> <p><b>Occupational Therapy Plan (Therapy Intervention Plan TIP):</b></p> <ul style="list-style-type: none"> <li>Not Current (#9)</li> </ul>	<p><b>Provider:</b>  <b>State your Plan of Correction for the deficiencies cited in this tag here</b> <i>(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?):</i> →</p> <p><b>Provider:</b>  <b>Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here</b> <i>(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?):</i> →</p>	

<p>settings.</p> <p>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.</p> <p>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.</p> <p>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.</p> <p>7. All records pertaining to JCMs must be retained permanently and must be made available to DDS upon request, upon the termination or expiration of a provider agreement, or upon provider withdrawal from services.</p> <p><b>20.5.1 Individual Data Form (IDF):</b> The Individual Data Form provides an overview of demographic information as well as other key personal, programmatic, insurance, and health related information. It lists medical information; assistive technology or adaptive equipment; diagnoses; allergies; information about whether a guardian or advance directives are in place; information about behavioral and health related needs; contacts of Provider Agencies and team members and other critical information. The IDF automatically loads information into other fields and forms and must be complete and kept current. This form is initiated by the CM. It must be opened and continuously updated by Living Supports,</p>			
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<p>CCS- Group, ANS, CIHS and case management when applicable to the person in order for accurate data to auto populate other documents like the Health Passport and Physician Consultation Form. Although the Primary Provider Agency is ultimately responsible for keeping this form current, each provider collaborates and communicates critical information to update this form.</p> <p><b>Chapter 3: Safeguards 3.1.2 Team Justification Process:</b> DD Waiver participants may receive evaluations or reviews conducted by a variety of professionals or clinicians. These evaluations or reviews typically include recommendations or suggestions for the person/guardian or the team to consider. The team justification process includes:</p> <ol style="list-style-type: none"> <li>1. Discussion and decisions about non-health related recommendations are documented on the Team Justification form.</li> <li>2. The Team Justification form documents that the person/guardian or team has considered the recommendations and has decided: <ol style="list-style-type: none"> <li>a. to implement the recommendation;</li> <li>b. to create an action plan and revise the ISP, if necessary; or</li> <li>c. not to implement the recommendation currently.</li> </ol> </li> <li>3. All DD Waiver Provider Agencies participate in information gathering, IDT meeting attendance, and accessing supplemental resources if needed and desired.</li> <li>4. The CM ensures that the Team Justification Process is followed and complete.</li> </ol>			
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<p>documenting the nature and frequency of service delivery, as well as data tracking only for the services provided by their agency.</p> <p>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.</p> <p>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.</p>			
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purpose in planning for individuals with developmental disabilities. [05/03/94; 01/15/97; Recompiled 10/31/01]

Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019

**Chapter 6: Individual Service Plan (ISP)**

**6.8 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 Chapter 20: Provider Documentation and Client Records.) CMs facilitate and maintain communication with the person, his/her representative, other IDT members, Provider Agencies, and relevant parties to ensure that the person receives the maximum benefit of his/her 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 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

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<p>essential to the service being provided and essential to ensuring the health and safety of the person during the provision of the service.</p> <p>2. Provider Agencies must have readily accessible records in home and community settings in paper or electronic form. Secure access to electronic records through the Therap web-based system using computers or mobile devices is acceptable.</p> <p>3. Provider Agencies are responsible for ensuring that all plans created by nurses, RDs, therapists or BSCs are present in all needed settings.</p> <p>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.</p> <p>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.</p> <p>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.</p> <p>7. All records pertaining to JCMs must be retained permanently and must be made available to DDS upon request, upon the termination or expiration of a provider agreement, or upon provider withdrawal from services.</p>			
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Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Completion Date
<p><b>Service Domain: Qualified Providers</b> – 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.</p>			
<p><b>Tag # 1A43.1 General Events Reporting: Individual Reporting</b></p>	<p><b>Standard Level Deficiency</b></p>		
<p>Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019  <b>Chapter 19: Provider Reporting Requirements: 19.2 General Events Reporting (GER):</b> The purpose of General Events Reporting (GER) is to report, track and analyze events, which pose a risk to adults in the DD Waiver program, but do not meet criteria for ANE or other reportable incidents as defined by the IMB. Analysis of GER is intended to identify emerging patterns so that preventative action can be taken at the individual, Provider Agency, regional and statewide level. On a quarterly and annual basis, DDS analyzes GER data at the provider, regional and statewide levels to identify any patterns that warrant intervention. Provider Agency use of GER in Therap is required as follows:  1. DD Waiver Provider Agencies approved to provide Customized In-Home Supports, Family Living, IMLS, Supported Living, Customized Community Supports, Community Integrated Employment, Adult Nursing and Case Management must use GER in the Therap system.  2. DD Waiver Provider Agencies referenced above are responsible for entering specified information into the GER section of the secure website operated under contract by Therap according to the GER Reporting Requirements in Appendix B GER Requirements.</p>	<p>Based on record review, the Agency did not follow the General Events Reporting requirements as indicated by the policy for 3 of 11 individuals.</p> <p><b>The following General Events Reporting records contained evidence that indicated the General Events Report was not entered and / or approved within the required timeframe:</b></p> <p><b>Individual #9</b></p> <ul style="list-style-type: none"> <li>General Events Report (GER) indicates on 8/25/2021 there was a Medication Error (Medication Error). GER was approved 10/17/2021.</li> <li>General Events Report (GER) indicates on 12/1/2021 received COVID-19 Booster Vaccine (COVID Vaccination). GER was approved 12/8/2021.</li> </ul> <p><b>Individual #10</b></p> <ul style="list-style-type: none"> <li>General Events Report (GER) indicates on 4/19/2021 Medication Error (Medication Error). GER was approved 6/2/2021.</li> <li>General Events Report (GER) indicates on 4/26/2021 there was a Medication Error (Medication Error). GER was approved 6/2/2021.</li> <li>General Events Report (GER) indicates on 4/27/2021 there was a Medication Error</li> </ul>	<p><b>Provider:</b>  <b>State your Plan of Correction for the deficiencies cited in this tag here</b> (<i>How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?</i>): →</p> <p><b>Provider:</b>  <b>Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here</b> (<i>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?</i>): →</p>	

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<p>3. At the Provider Agency's discretion additional events, which are not required by DDS, may also be tracked within the GER section of Therap.</p> <p>4. GER does not replace a Provider Agency's obligations to report ANE or other reportable incidents as described in Chapter 18: Incident Management System.</p> <p>5. GER does not replace a Provider Agency's obligations related to healthcare coordination, modifications to the ISP, or any other risk management and QI activities.</p> <p><b>Appendix B GER Requirements:</b> DDS is pleased to introduce the revised General Events Reporting (GER), requirements. There are two important changes related to medication error reporting:</p> <p>1. <i>Effective immediately</i>, DDS requires ALL medication errors be entered into Therap GER with the exception of those required to be reported to Division of Health Improvement-Incident Management Bureau.</p> <p>2. No alternative methods for reporting are permitted.</p> <p><b><u>The following events need to be reported in the Therap GER:</u></b></p> <ul style="list-style-type: none"> <li>• Emergency Room/Urgent Care/Emergency Medical Services</li> <li>• Falls Without Injury</li> <li>• Injury (including Falls, Choking, Skin Breakdown and Infection)</li> <li>• Law Enforcement Use</li> <li>• Medication Errors</li> <li>• Medication Documentation Errors</li> <li>• Missing Person/Elopement</li> <li>• Out of Home Placement- Medical: Hospitalization, Long Term Care, Skilled Nursing or Rehabilitation Facility Admission</li> <li>• PRN Psychotropic Medication</li> </ul>	<p>(Medication Error). GER was approved 6/2/2021.</p> <ul style="list-style-type: none"> <li>• General Events Report (GER) indicates on 4/28/2021 individual had an injury from a fall (Injury). GER was approved 5/3/2021.</li> <li>• General Events Report (GER) indicates on 7/22/2021 the individual was taken to UNMH Psych. (Law Enforcement Involvement). GER was approved 7/29/2021.</li> <li>• General Events Report (GER) indicates on 12/1/2021 the Individual received a COVID-19 Booster (COVID Vaccination). GER was approved 12/7/2021.</li> <li>• General Events Report (GER) indicates on 1/5/2022 there was a Medication Error (Medication Error). GER was approved 2/28/2022.</li> </ul> <p><b>Individual #11</b></p> <ul style="list-style-type: none"> <li>• General Events Report (GER) indicates on 4/19/2021 there was a Medication Error (Medication Error). GER was approved 7/8/2021.</li> <li>• General Events Report (GER) indicates on 7/2/2021 the individual had a Seizure. (Emergency Services). GER was approved 7/7/2021.</li> <li>• General Events Report (GER) indicates on 8/31/2021 there was a Medication Error (Medication Error). GER was approved 10/17/2021.</li> <li>• General Events Report (GER) indicates on 12/1/2021 individual received the COVID-19 Booster (COVID Vaccine). GER was approved 12/8/2021.</li> </ul>		
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<ul style="list-style-type: none"> <li>• Restraint Related to Behavior</li> <li>• Suicide Attempt or Threat</li> </ul> <p><b>Entry Guidance:</b> Provider Agencies must complete the following sections of the GER with detailed information: profile information, event information, other event information, general information, notification, actions taken or planned, and the review follow up comments section. Please attach any pertinent external documents such as discharge summary, medical consultation form, etc. <u>Provider Agencies must enter and approve GERs within 2 business days with the exception of Medication Errors which must be entered into GER on at least a monthly basis.</u></p>	<p><b>The following events were not reported in the General Events Reporting System as required by policy:</b></p> <p><b>Individual #9</b></p> <ul style="list-style-type: none"> <li>• Documentation reviewed indicates on 5/3/2021 the Individual went to Urgent Care (Urgent Care). No GER was found.</li> <li>• Documentation reviewed indicates on 1/31/2022 the Individual went to the Hospital (Hospital). No GER was found.</li> <li>• Documentation reviewed indicates on 2/7/2022 the Individual went to the ER (Emergency Room). No GER was found.</li> </ul> <p><b>Individual #10</b></p> <ul style="list-style-type: none"> <li>• Documentation reviewed indicates on 7/1/2021 the Individual went to Urgent Care. No GER was found.</li> <li>• Documentation reviewed indicates on 7/10/2021 the Individual went to Urgent Care (Urgent Care). No GER was found.</li> <li>• Documentation reviewed indicates on 8/13/2021 the Individual went to Urgent Care (Urgent Care). No GER was found.</li> <li>• Documentation reviewed indicates on 9/17/2021 the Individual went to Urgent Care (Urgent Care). No GER was found.</li> <li>• Documentation reviewed indicates on 10/5/2021 the Individual went to Urgent Care (Urgent Care). No GER was found.</li> <li>• Documentation reviewed indicates on 11/2/2021 the Individual went to Urgent Care (Urgent Care). No GER was found.</li> </ul>		
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|  | <ul style="list-style-type: none"><li>• Documentation reviewed indicates on 11/17/2021 the Individual went to Urgent Care (Urgent Care). No GER was found.</li><li>• Documentation reviewed indicates on 3/4/2022 the Individual went to Urgent Care (Urgent Care). No GER was found.</li><li>• Documentation reviewed indicates on 3/22/2022 the Individual went to Urgent Care (Urgent Care). No GER was found.</li></ul> |  |  |
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Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Completion Date
<b>Service Domain: Health and Welfare</b> – 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.			
<b>Tag # 1A08.2 Administrative Case File: Healthcare Requirements &amp; Follow-up</b>	<b>Condition of Participation Level Deficiency</b>		
<p>Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019</p> <p><b>Chapter 3 Safeguards: 3.1.1 Decision Consultation Process (DCP):</b> 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 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:</p> <ol style="list-style-type: none"> <li>1. The DCP is used when a person or his/her guardian/healthcare decision maker has concerns, needs more information about health-related issues, or has decided not to follow all or part of an order, recommendation, or suggestion. This includes, but is not limited to: <ol style="list-style-type: none"> <li>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;</li> <li>b. clinical recommendations made by registered/licensed clinicians who are either members of the IDT or clinicians who have performed an evaluation such as a video-fluoroscopy;</li> <li>c. health related recommendations or suggestions from oversight activities such</li> </ol> </li> </ol>	<p>After an analysis of the evidence, it has been determined there is a significant potential for a negative outcome to occur.</p> <p>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 2 of 11 individuals receiving Living Care Arrangements and Community Inclusion.</p> <p>Review of the administrative individual case files revealed the following items were not found, incomplete, and/or not current:</p> <p><b>Community Inclusion Services (Individuals Receiving Inclusion Services Only):</b></p> <p><b>Annual Physical:</b></p> <ul style="list-style-type: none"> <li>• Not Found (#1)</li> </ul> <p><b>Living Care Arrangements / Community Inclusion (Individuals Receiving Multiple Services):</b></p> <p><b>Podiatry:</b></p> <ul style="list-style-type: none"> <li>• Individual #12 - As indicated by collateral documentation reviewed, exam was completed on 9/8/2021. Follow-up was scheduled for 11/23/2021. No evidence of follow-up found.</li> </ul>	<p><b>Provider:</b> 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?): →</p> <p><b>Provider:</b> 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?): →</p>	

<p>as the Individual Quality Review (IQR) or other DOH review or oversight activities; and</p> <p>d. recommendations made through a Healthcare Plan (HCP), including a Comprehensive Aspiration Risk Management Plan (CARMP), or another plan.</p> <p>2. When the person/guardian disagrees with a recommendation or does not agree with the implementation of that recommendation, Provider Agencies follow the DCP and attend the meeting coordinated by the CM. During this meeting:</p> <p>a. Providers inform the person/guardian of the rationale for that recommendation, so that the benefit is made clear. This will be done in layman's terms and will include basic sharing of information designed to assist the person/guardian with understanding the risks and benefits of the recommendation.</p> <p>b. The information will be focused on the specific area of concern by the person/guardian. Alternatives should be presented, when available, if the guardian is interested in considering other options for implementation.</p> <p>c. Providers support the person/guardian to make an informed decision.</p> <p>d. The decision made by the person/guardian during the meeting is accepted; plans are modified; and the IDT honors this health decision in every setting.</p> <p><b>Chapter 20: Provider Documentation and Client Records: 20.2 Client Records Requirements:</b> All DD Waiver Provider Agencies are required to create and maintain</p>			
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<p>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:</p> <ol style="list-style-type: none"> <li>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.</li> <li>2. Provider Agencies must have readily accessible records in home and community settings in paper or electronic form. Secure access to electronic records through the Therap web-based system using computers or mobile devices is acceptable.</li> <li>3. Provider Agencies are responsible for ensuring that all plans created by nurses, RDs, therapists or BSCs are present in all needed settings.</li> <li>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.</li> <li>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.</li> <li>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.</li> </ol>			
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7. All records pertaining to JCMs must be retained permanently and must be made available to DDSD upon request, upon the termination or expiration of a provider agreement, or upon provider withdrawal from services.

**20.5.3 Health Passport and Physician**

**Consultation Form:** All Primary and Secondary Provider Agencies must use the *Health Passport and Physician Consultation* form from the Therap system. This standardized document contains individual, physician and emergency contact information, a complete list of current medical diagnoses, health and safety risk factors, allergies, and information regarding insurance, guardianship, and advance directives. The *Health Passport* also includes a standardized form to use at medical appointments called the *Physician Consultation* form. The *Physician Consultation* form contains a list of all current medications.

**Chapter 10: Living Care Arrangements (LCA) Living Supports-Supported Living: 10.3.9.6.1 Monitoring and Supervision**

4. Ensure and document the following:
- a. The person has a Primary Care Practitioner.
  - b. The person receives an annual physical examination and other examinations as recommended by a Primary Care Practitioner or specialist.
  - c. The person receives annual dental check-ups and other check-ups as recommended by a licensed dentist.
  - d. The person receives a hearing test as recommended by a licensed audiologist.
  - e. The person receives eye examinations as

<p>recommended by a licensed optometrist or ophthalmologist.</p> <p>5. Agency activities occur as required for follow-up activities to medical appointments (e.g., treatment, visits to specialists, and changes in medication or daily routine).</p> <p><b>10.3.10.1 Living Care Arrangements (LCA) Living Supports-IMLS: 10.3.10.2 General Requirements:</b> 9 . Medical services must be ensured (i.e., ensure each person has a licensed Primary Care Practitioner and receives an annual physical examination, specialty medical care as needed, and annual dental checkup by a licensed dentist).</p> <p><b>Chapter 13 Nursing Services: 13.2.3 General Requirements:</b></p> <p>1. Each person has a licensed primary care practitioner and receives an annual physical examination and specialty medical/dental care as needed. Nurses communicate with these providers to share current health information.</p>			
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<p>counter (OTC) or “comfort” medications or treatments and all self-selected herbal or vitamin therapy;</p> <p>c. Documentation of all time limited or discontinued medications or treatments;</p> <p>d. The initials of the individual administering or assisting with the medication delivery and a signature page or electronic record that designates the full name corresponding to the initials;</p> <p>e. Documentation of refused, missed, or held medications or treatments;</p> <p>f. Documentation of any allergic reaction that occurred due to medication or treatments; and</p> <p>g. For PRN medications or treatments:</p> <p>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;</p> <p>ii. clear documentation that the DSP contacted the agency nurse prior to assisting with the medication or treatment, unless the DSP is a Family Living Provider related by affinity of consanguinity; and</p> <p>iii. documentation of the effectiveness of the PRN medication or treatment.</p> <p><b>Chapter 10 Living Care Arrangements</b>  <b>10.3.4 Medication Assessment and Delivery:</b>  Living Supports Provider Agencies must support and comply with:  1. the processes identified in the DDSD AWMD training;</p>			
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<p>2. the nursing and DSP functions identified in the Chapter 13.3 Part 2- Adult Nursing Services;</p> <p>3. all Board of Pharmacy regulations as noted in Chapter 16.5 Board of Pharmacy; and</p> <p>4. documentation requirements in a Medication Administration Record (MAR) as described in Chapter 20.6 Medication Administration Record (MAR).</p> <p><b>NMAC 16.19.11.8 MINIMUM STANDARDS:</b></p> <p>A. MINIMUM STANDARDS FOR THE DISTRIBUTION, STORAGE, HANDLING AND RECORD KEEPING OF DRUGS:</p> <p>(d) The facility shall have a Medication Administration Record (MAR) documenting medication administered to residents, <b>including over-the-counter medications.</b></p> <p>This documentation shall include:</p> <ul style="list-style-type: none"> <li>(i) Name of resident;</li> <li>(ii) Date given;</li> <li>(iii) Drug product name;</li> <li>(iv) Dosage and form;</li> <li>(v) Strength of drug;</li> <li>(vi) Route of administration;</li> <li>(vii) How often medication is to be taken;</li> <li>(viii) Time taken and staff initials;</li> <li>(ix) Dates when the medication is discontinued or changed;</li> <li>(x) The name and initials of all staff administering medications.</li> </ul> <p><b>Model Custodial Procedure Manual</b>  <b>D. Administration of Drugs</b></p> <p>Unless otherwise stated by practitioner, patients will not be allowed to administer their own medications.</p> <p>Document the practitioner's order authorizing the self-administration of medications.</p> <p>All PRN (As needed) medications shall have complete detail instructions regarding the</p>			
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<p>administering of the medication. This shall include:</p> <ul style="list-style-type: none"><li>➤ symptoms that indicate the use of the medication,</li><li>➤ exact dosage to be used, and</li><li>➤ the exact amount to be used in a 24-hour period.</li></ul>			
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<p>counter (OTC) or “comfort” medications or treatments and all self-selected herbal or vitamin therapy;</p> <p>c. Documentation of all time limited or discontinued medications or treatments;</p> <p>d. The initials of the individual administering or assisting with the medication delivery and a signature page or electronic record that designates the full name corresponding to the initials;</p> <p>e. Documentation of refused, missed, or held medications or treatments;</p> <p>f. Documentation of any allergic reaction that occurred due to medication or treatments; and</p> <p>g. For PRN medications or treatments:</p> <p>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;</p> <p>ii. clear documentation that the DSP contacted the agency nurse prior to assisting with the medication or treatment, unless the DSP is a Family Living Provider related by affinity of consanguinity; and</p> <p>iii. documentation of the effectiveness of the PRN medication or treatment.</p> <p><b>Chapter 10 Living Care Arrangements</b>  <b>10.3.4 Medication Assessment and Delivery:</b>  Living Supports Provider Agencies must support and comply with:  1. the processes identified in the DDSD AWMD training;</p>			
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<p>2. the nursing and DSP functions identified in the Chapter 13.3 Part 2- Adult Nursing Services;</p> <p>3. all Board of Pharmacy regulations as noted in Chapter 16.5 Board of Pharmacy; and</p> <p>4. documentation requirements in a Medication Administration Record (MAR) as described in Chapter 20.6 Medication Administration Record (MAR).</p> <p><b>NMAC 16.19.11.8 MINIMUM STANDARDS:</b></p> <p>A. MINIMUM STANDARDS FOR THE DISTRIBUTION, STORAGE, HANDLING AND RECORD KEEPING OF DRUGS:</p> <p>(d) The facility shall have a Medication Administration Record (MAR) documenting medication administered to residents, <b>including over-the-counter medications.</b></p> <p>This documentation shall include:</p> <ul style="list-style-type: none"> <li>(i) Name of resident;</li> <li>(ii) Date given;</li> <li>(iii) Drug product name;</li> <li>(iv) Dosage and form;</li> <li>(v) Strength of drug;</li> <li>(vi) Route of administration;</li> <li>(vii) How often medication is to be taken;</li> <li>(viii) Time taken and staff initials;</li> <li>(ix) Dates when the medication is discontinued or changed;</li> <li>(x) The name and initials of all staff administering medications.</li> </ul> <p><b>Model Custodial Procedure Manual</b>  <b>D. Administration of Drugs</b></p> <p>Unless otherwise stated by practitioner, patients will not be allowed to administer their own medications.</p> <p>Document the practitioner's order authorizing the self-administration of medications.</p> <p>All PRN (As needed) medications shall have complete detail instructions regarding the</p>			
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<p>administering of the medication. This shall include:</p> <ul style="list-style-type: none"><li>➤ symptoms that indicate the use of the medication,</li><li>➤ exact dosage to be used, and</li><li>➤ the exact amount to be used in a 24-hour period.</li></ul>			
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Tag # 1A09.1 Medication Delivery PRN Medication Administration	Standard Level Deficiency		
<p>Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019</p> <p><b>Chapter 20: Provider Documentation and Client Records 20.6 Medication Administration Record (MAR):</b> A current Medication Administration Record (MAR) must be maintained in all settings where medications or treatments are delivered. Family Living Providers may opt not to use MARs if they are the sole provider who supports the person with medications or treatments. However, if there are services provided by unrelated DSP, ANS for Medication Oversight must be budgeted, and a MAR must be created and used by the DSP. Primary and Secondary Provider Agencies are responsible for:</p> <ol style="list-style-type: none"> <li>1. Creating and maintaining either an electronic or paper MAR in their service setting. Provider Agencies may use the MAR in Therap, but are not mandated to do so.</li> <li>2. Continually communicating any changes about medications and treatments between Provider Agencies to assure health and safety. <ol style="list-style-type: none"> <li>7. Including the following on the MAR: <ol style="list-style-type: none"> <li>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;</li> <li>b. The prescribed dosage, frequency and method or route of administration; times and dates of administration for all ordered routine or PRN prescriptions or treatments; over the</li> </ol> </li> </ol> </li> </ol>	<p>Medication Administration Records (MAR) were reviewed for the month of March 2022.</p> <p>Based on record review, 1 of 8 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard:</p> <p>Individual #11 March 2022</p> <p>Medication Administration Records contain the following medications. No Physician's Orders were found for the following medications:</p> <ul style="list-style-type: none"> <li>• Benzonatate 200mg (PRN)</li> <li>• Claritin 10mg (PRN)</li> <li>• Docusate Sodium 100mg (PRN)</li> <li>• Eucerin Lotion (PRN)</li> </ul> <p>Physician's Orders indicated the following medication were to be given. The following Medications were not documented on the Medication Administration Records:</p> <ul style="list-style-type: none"> <li>• Diphenhydramine 25 / 50mg (PRN)</li> </ul>	<p><b>Provider:</b> <b>State your Plan of Correction for the deficiencies cited in this tag here</b> <i>(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?):</i> →</p> <p><b>Provider:</b> <b>Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here</b> <i>(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?):</i> →</p>	

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<p>counter (OTC) or “comfort” medications or treatments and all self-selected herbal or vitamin therapy;</p> <p>c. Documentation of all time limited or discontinued medications or treatments;</p> <p>d. The initials of the individual administering or assisting with the medication delivery and a signature page or electronic record that designates the full name corresponding to the initials;</p> <p>e. Documentation of refused, missed, or held medications or treatments;</p> <p>f. Documentation of any allergic reaction that occurred due to medication or treatments; and</p> <p>g. For PRN medications or treatments:</p> <p>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;</p> <p>ii. clear documentation that the DSP contacted the agency nurse prior to assisting with the medication or treatment, unless the DSP is a Family Living Provider related by affinity of consanguinity; and</p> <p>iii. documentation of the effectiveness of the PRN medication or treatment.</p> <p><b>Chapter 10 Living Care Arrangements</b>  <b>10.3.4 Medication Assessment and Delivery:</b>  Living Supports Provider Agencies must support and comply with:</p> <ol style="list-style-type: none"> <li>1. the processes identified in the DDSD AWMD training;</li> </ol>			
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<p>2. the nursing and DSP functions identified in the Chapter 13.3 Part 2- Adult Nursing Services;</p> <p>3. all Board of Pharmacy regulations as noted in Chapter 16.5 Board of Pharmacy; and</p> <p>4. documentation requirements in a Medication Administration Record (MAR) as described in Chapter 20.6 Medication Administration Record (MAR).</p>			
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<p>counter (OTC) or “comfort” medications or treatments and all self-selected herbal or vitamin therapy;</p> <p>c. Documentation of all time limited or discontinued medications or treatments;</p> <p>d. The initials of the individual administering or assisting with the medication delivery and a signature page or electronic record that designates the full name corresponding to the initials;</p> <p>e. Documentation of refused, missed, or held medications or treatments;</p> <p>f. Documentation of any allergic reaction that occurred due to medication or treatments; and</p> <p>g. For PRN medications or treatments:</p> <p>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;</p> <p>ii. clear documentation that the DSP contacted the agency nurse prior to assisting with the medication or treatment, unless the DSP is a Family Living Provider related by affinity of consanguinity; and</p> <p>iii. documentation of the effectiveness of the PRN medication or treatment.</p> <p><b>Chapter 10 Living Care Arrangements</b>  <b>10.3.4 Medication Assessment and Delivery:</b>  Living Supports Provider Agencies must support and comply with:</p> <ol style="list-style-type: none"> <li>1. the processes identified in the DDSD AWMD training;</li> </ol>			
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<p>2. the nursing and DSP functions identified in the Chapter 13.3 Part 2- Adult Nursing Services;</p> <p>3. all Board of Pharmacy regulations as noted in Chapter 16.5 Board of Pharmacy; and</p> <p>4. documentation requirements in a Medication Administration Record (MAR) as described in Chapter 20.6 Medication Administration Record (MAR).</p>			
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Tag # 1A15.2 Administrative Case File: Healthcare Documentation (Therap and Required Plans)	Condition of Participation Level Deficiency		
<p>Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019</p> <p><b>Chapter 20: Provider Documentation and Client Records: 20.2 Client Records Requirements:</b> 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:</p> <ol style="list-style-type: none"> <li>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.</li> <li>2. Provider Agencies must have readily accessible records in home and community settings in paper or electronic form. Secure access to electronic records through the Therap web-based system using computers or mobile devices is acceptable.</li> <li>3. Provider Agencies are responsible for ensuring that all plans created by nurses, RDs, therapists or BSCs are present in all needed settings.</li> <li>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.</li> <li>5. Each Provider Agency is responsible for</li> </ol>	<p>After an analysis of the evidence, it has been determined there is a significant potential for a negative outcome to occur.</p> <p>Based on record review, the Agency did not maintain the required documentation in the Individuals Agency Record as required by standard for 5 of 11 individuals.</p> <p>Review of the administrative individual case files revealed the following items were not found, incomplete, and/or not current:</p> <p><b>Electronic Comprehensive Health Assessment Tool (eCHAT):</b></p> <ul style="list-style-type: none"> <li>➤ Not Found (#1)</li> </ul> <p><b>eCHAT Summary:</b></p> <ul style="list-style-type: none"> <li>➤ Not Found (#1, 9)</li> </ul> <p><b>Medication Administration Assessment Tool:</b></p> <ul style="list-style-type: none"> <li>➤ Not Found (#1)</li> </ul> <p><b>Aspiration Risk Screening Tool:</b></p> <ul style="list-style-type: none"> <li>➤ Not Found (#1)</li> </ul> <p><b>Healthcare Passport:</b></p> <ul style="list-style-type: none"> <li>➤ Did not contain Name of Physician (#1)</li> <li>➤ Did not contain Emergency Contact Information (#1, 13)</li> <li>➤ Did not contain Did not contain Health &amp; Safety risk factors (#1)</li> <li>➤ Did not contain Guardianship/Healthcare Decision Maker (#1, 13)</li> </ul>	<p><b>Provider:</b>  <b>State your Plan of Correction for the deficiencies cited in this tag here</b> <i>(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?):</i> →</p> <p><b>Provider:</b>  <b>Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here</b> <i>(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?):</i> →</p>	

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<p>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.</p> <p>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.</p> <p>7. All records pertaining to JCMs must be retained permanently and must be made available to DDS upon request, upon the termination or expiration of a provider agreement, or upon provider withdrawal from services.</p> <p><b>Chapter 3 Safeguards: 3.1.1 Decision Consultation Process (DCP):</b> 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 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:</p> <p>2. The DCP is used when a person or his/her guardian/healthcare decision maker has concerns, needs more information about health-related issues, or has decided not to follow all or part of an order, recommendation, or suggestion. This includes, but is not limited to:</p> <p>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;</p>	<p>➤ Did not contain Information Regarding Insurance (#2)</p> <p><b>Medical Emergency Response Plans:</b></p> <p><b>Aspiration</b></p> <p>➤ Individual #5 - As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found.</p> <p><b>Paralysis</b></p> <p>➤ Individual #5 - As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found.</p> <p><b>Seizures</b></p> <p>➤ Individual #5 - As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found</p>		
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<p>b. clinical recommendations made by registered/licensed clinicians who are either members of the IDT or clinicians who have performed an evaluation such as a video-fluoroscopy;</p> <p>c. health related recommendations or suggestions from oversight activities such as the Individual Quality Review (IQR) or other DOH review or oversight activities; and</p> <p>d. recommendations made through a Healthcare Plan (HCP), including a Comprehensive Aspiration Risk Management Plan (CARMP), or another plan.</p> <p>2. When the person/guardian disagrees with a recommendation or does not agree with the implementation of that recommendation, Provider Agencies follow the DCP and attend the meeting coordinated by the CM. During this meeting:</p> <p>a. Providers inform the person/guardian of the rationale for that recommendation, so that the benefit is made clear. This will be done in layman's terms and will include basic sharing of information designed to assist the person/guardian with understanding the risks and benefits of the recommendation.</p> <p>b. The information will be focused on the specific area of concern by the person/guardian. Alternatives should be presented, when available, if the guardian is interested in considering other options for implementation.</p> <p>c. Providers support the person/guardian to make an informed decision.</p> <p>d. The decision made by the person/guardian during the meeting is accepted; plans are modified; and the IDT honors this health decision in every setting.</p>			
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<p><b>Chapter 13 Nursing Services: 13.2.5 Electronic Nursing Assessment and Planning Process:</b> The nursing assessment process includes several DDS mandated tools: the electronic Comprehensive Nursing Assessment Tool (e-CHAT), the Aspiration Risk Screening Tool (ARST) and the Medication Administration Assessment Tool (MAAT) . This process includes developing and training Health Care Plans and Medical Emergency Response Plans.</p> <p>The following hierarchy is based on budgeted services and is used to identify which Provider Agency nurse has primary responsibility for completion of the nursing assessment process and related subsequent planning and training. Additional communication and collaboration for planning specific to CCS or CIE services may be needed.</p> <p>The hierarchy for Nursing Assessment and Planning responsibilities is:</p> <ol style="list-style-type: none"> <li>1. Living Supports: Supported Living, IMLS or Family Living via ANS;</li> <li>2. Customized Community Supports- Group; and</li> <li>3. Adult Nursing Services (ANS):       <ol style="list-style-type: none"> <li>a. for persons in Community Inclusion with health-related needs; or</li> <li>b. if no residential services are budgeted but assessment is desired and health needs may exist.</li> </ol> </li> </ol> <p><b>13.2.6 The Electronic Comprehensive Health Assessment Tool (e-CHAT)</b></p> <ol style="list-style-type: none"> <li>1. The e-CHAT is a nursing assessment. It may not be delegated by a licensed nurse to a non-licensed person.</li> <li>2. The nurse must see the person face-to-face to complete the nursing assessment. Additional information may be gathered from members of the IDT and other sources.</li> <li>3. An e-CHAT is required for persons in FL,</li> </ol>			
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SL, IMLS, or CCS-Group. All other DD Waiver recipients may obtain an e-CHAT if needed or desired by adding ANS hours for assessment and consultation to their budget.

4. When completing the e-CHAT, the nurse is required to review and update the electronic record and consider the diagnoses, medications, treatments, and overall status of the person. Discussion with others may be needed to obtain critical information.

5. The nurse is required to complete all the e-CHAT assessment questions and add additional pertinent information in all comment sections.

**13.2.7 Aspiration Risk Management Screening Tool (ARST)**

**13.2.8 Medication Administration Assessment Tool (MAAT):**

1. A licensed nurse completes the DDS Medication Administration Assessment Tool (MAAT) at least two weeks before the annual ISP meeting.

2. After completion of the MAAT, the nurse will present recommendations regarding the level of assistance with medication delivery (AWMD) to the IDT. A copy of the MAAT will be sent to all the team members two weeks before the annual ISP meeting and the original MAAT will be retained in the Provider Agency records.

3. Decisions about medication delivery are made by the IDT to promote a person's maximum independence and community integration. The IDT will reach consensus regarding which criteria the person meets, as indicated by the results of the MAAT and the nursing recommendations, and the decision is documented this in the ISP.

**13.2.9 Healthcare Plans (HCP):**

<p>1. At the nurse’s discretion, based on prudent nursing practice, interim HCPs may be developed to address issues that must be implemented immediately after admission, readmission or change of medical condition to provide safe services prior to completion of the e-CHAT and formal care planning process. This includes interim ARM plans for those persons newly identified at moderate or high risk for aspiration. All interim plans must be removed if the plan is no longer needed or when final HCP including CARMPs are in place to avoid duplication of plans.</p> <p>2. In collaboration with the IDT, the agency nurse is required to create HCPs that address all the areas identified as required in the most current e-CHAT summary report which is indicated by “R” in the HCP column. At the nurse’s sole discretion, based on prudent nursing practice, HCPs may be combined where clinically appropriate. The nurse should use nursing judgment to determine whether to also include HCPs for any of the areas indicated by “C” on the e-CHAT summary report. The nurse may also create other HCPs plans that the nurse determines are warranted.</p> <p><b>13.2.10 Medical Emergency Response Plan (MERP):</b></p> <p>1. The agency nurse is required to develop a Medical Emergency Response Plan (MERP) for all conditions marked with an "R" in the e-CHAT summary report. The agency nurse should use her/his clinical judgment and input from the Interdisciplinary Team (IDT) to determine whether shown as "C" in the e-CHAT summary report or other conditions also warrant a MERP.</p> <p>2. MERPs are required for persons who have one or more conditions or illnesses that present a likely potential to become a life-threatening situation.</p>			
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**Chapter 20: Provider Documentation and Client Records: 20.5.3 Health Passport and Physician Consultation Form:** All Primary and Secondary Provider Agencies must use the Health Passport and Physician Consultation form from the Therap system. This standardized document contains individual, physician and emergency contact information, a complete list of current medical diagnoses, health and safety risk factors, allergies, and information regarding insurance, guardianship, and advance directives. The Health Passport also includes a standardized form to use at medical appointments called the Physician Consultation form.

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Completion Date
<b>Service Domain: Medicaid Billing/Reimbursement</b> – State financial oversight exists to assure that claims are coded and paid for in accordance with the reimbursement methodology specified in the approved waiver.			
<b>Tag # IS30 Customized Community Supports Reimbursement</b>	<b>Standard Level Deficiency</b>		
<p>Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019</p> <p><b>Chapter 21: Billing Requirements: 21.4 Recording Keeping and Documentation Requirements:</b> 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:</p> <ol style="list-style-type: none"> <li>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.</li> <li>2. Comprehensive documentation of direct service delivery must include, at a minimum: <ol style="list-style-type: none"> <li>a. the agency name;</li> <li>b. the name of the recipient of the service;</li> <li>c. the location of the service;</li> <li>d. the date of the service;</li> <li>e. the type of service;</li> <li>f. the start and end times of the service;</li> <li>g. the signature and title of each staff member who documents their time; and</li> <li>h. the nature of services.</li> </ol> </li> <li>3. 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.</li> <li>4. A Provider Agency that receives payment for treatment, services or goods must retain all medical and business records relating to any</li> </ol>	<p>Based on record review, the Agency did not provide written or electronic documentation as evidence for each unit billed for Customized Community Supports for 6 of 11 individuals.</p> <p>Individual #2 December 2021</p> <ul style="list-style-type: none"> <li>• The Agency billed 153 units of Customized Community Supports (Individual) (H2021 HB U1) from 12/16/2021 through 12/29/2021. Documentation received accounted for 152 units.</li> </ul> <p>Individual #4 December 2021</p> <ul style="list-style-type: none"> <li>• The Agency billed 4 units of Customized Community Supports (Individual) (H2021 HB U1). No documentation was found for 12/16/2021 through 12/29/2021 to justify the 4 units billed.</li> </ul> <p>January 2022</p> <ul style="list-style-type: none"> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 1/6/2022 through 1/12/2022. Documentation received accounted for 18 units.</li> </ul> <p>Individual #5 January 2022</p> <ul style="list-style-type: none"> <li>• The Agency billed 4 units of Customized Community Supports (Group) (T2021 HB U7) from 1/13/2022 through 1/19/2022. Documentation received accounted for 0 units.</li> </ul>	<p><b>Provider:</b> 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?): →</p> <p><b>Provider:</b> 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?): →</p>	

<p>of the following for a period of at least six years from the payment date:</p> <ol style="list-style-type: none"> <li>treatment or care of any eligible recipient;</li> <li>services or goods provided to any eligible recipient;</li> <li>amounts paid by MAD on behalf of any eligible recipient; and</li> <li>any records required by MAD for the administration of Medicaid.</li> </ol> <p><b>21.9 Billable Units:</b> 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.</p> <p><b>21.9.1 Requirements for Daily Units:</b> For services billed in daily units, Provider Agencies must adhere to the following:</p> <ol style="list-style-type: none"> <li>A day is considered 24 hours from midnight to midnight.</li> <li>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.</li> <li>The maximum allowable billable units cannot exceed 340 calendar days per ISP year or 170 calendar days per six months.</li> <li>When a person transitions from one Provider Agency to another during the ISP year, a standard formula to calculate the units billed by each Provider Agency must be applied as follows: <ol style="list-style-type: none"> <li>The discharging Provider Agency bills the number of calendar days that services were provided multiplied by .93 (93%).</li> <li>The receiving Provider Agency bills the remaining days up to 340 for the ISP</li> </ol> </li> </ol>	<p>February 2022</p> <ul style="list-style-type: none"> <li>The Agency billed 4 units of Customized Community Supports (Group) (T2021 HB U7) from 2/17/2022 through 2/23/2022. Documentation received accounted for 0 units.</li> <li>The Agency billed 5 units of Customized Community Supports (Group) (T2021 HB U7) from 2/24/2022 through 3/1/2022. Documentation received accounted for 0 units.</li> </ul> <p>Individual #9 January 2022</p> <ul style="list-style-type: none"> <li>The Agency billed 47 units of Customized Community Supports (Group) (T2021 HB U8) from 1/6/2022 through 1/12/2022. No Documentation was found for 1/6/2022 through 1/12/2022 to justify the 47 units billed.</li> </ul> <p>Individual #11 December 2021</p> <ul style="list-style-type: none"> <li>The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 12/2/2021 through 12/8/2021. Documentation received accounted for 100 units.</li> <li>The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 12/9/2021 through 12/15/2021. Documentation received accounted for 112 units.</li> <li>The Agency billed 240 units of Customized Community Supports (Individual) (H2021 HB U1) from 12/16/2021 through 12/29/2021.</li> </ul>		
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<p>year.</p> <p><b>21.9.2 Requirements for Monthly Units:</b> For services billed in monthly units, a Provider Agency must adhere to the following:</p> <ol style="list-style-type: none"> <li>1. A month is considered a period of 30 calendar days.</li> <li>2. At least one hour of face-to-face billable services shall be provided during a calendar month where any portion of a monthly unit is billed.</li> <li>3. Monthly units can be prorated by a half unit.</li> <li>4. Agency transfers not occurring at the beginning of the 30-day interval are required to be coordinated in the middle of the 30-day interval so that the discharging and receiving agency receive a half unit.</li> </ol> <p><b>21.9.3 Requirements for 15-minute and hourly units:</b> For services billed in 15-minute or hourly intervals, Provider Agencies must adhere to the following:</p> <ol style="list-style-type: none"> <li>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.</li> <li>2. Services that last in their entirety less than eight minutes cannot be billed.</li> </ol>	<p>Documentation received accounted for 188 units.</p> <ul style="list-style-type: none"> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 12/30/2021 through 1/5/2022. Documentation received accounted for 88 units.</li> </ul> <p>January 2022</p> <ul style="list-style-type: none"> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 1/6/2022 through 1/12/2022. Documentation received accounted for 108 units.</li> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 1/13/2022 through 1/19/2022. Documentation received accounted for 104 units.</li> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 1/20/2022 through 1/26/2022. Documentation received accounted for 88 units.</li> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 1/27/2022 through 2/2/2022. Documentation received accounted for 80 units.</li> </ul> <p>February 2022</p> <ul style="list-style-type: none"> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 2/3/2022</li> </ul>		
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	<p>through 2/9/2022. Documentation received accounted for 100 units.</p> <ul style="list-style-type: none"> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 2/10/2022 through 2/16/2022. Documentation received accounted for 104 units.</li> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 2/17/2022 through 2/23/2022. Documentation received accounted for 92 units.</li> </ul> <p>Individual #12 December 2021</p> <ul style="list-style-type: none"> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 12/2/2021 through 12/8/2021. Documentation received accounted for 78 units.</li> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 12/9/2021 through 12/15/2021. Documentation received accounted for 92 units.</li> <li>• The Agency billed 216 units of Customized Community Supports (Individual) (H2021 HB U1) from 12/16/2021 through 12/29/2021. Documentation received accounted for 160 units.</li> <li>• The Agency billed 24 units of Customized Community Supports (Group) (T2021 HB</li> </ul>		
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	<p>U8) from 12/16/2022 through 12/29/2021. Documentation received accounted for 16 units.</p> <ul style="list-style-type: none"> <li>• The Agency billed 24 units of Customized Community Supports (Group) (T2021 HB U8) from 12/16/2022 through 12/29/2021. Documentation received accounted for 16 units.</li> <li>• The Agency billed 96 units of Customized Community Supports (Individual) (H2021 HB U1) from 12/30/2021 through 1/5/2022. Documentation received accounted for 60 units.</li> </ul> <p>January 2022</p> <ul style="list-style-type: none"> <li>• The Agency billed 120 units of Customized Community Supports (Individual) (H2021 HB U1) from 1/13/2022 through 1/19/2022. No documentation was found for 1/13/2022 through 1/19/2022.</li> <li>• The Agency billed 120 units of Customized Community Supports (Group) (T2021 HB U8) from 1/20/2022 through 1/26/2022. Documentation received accounted for 92 units.</li> <li>• The Agency billed 120 units of Customized Community Supports (Group) (T2021 HB U8) from 1/27/2022 through 2/2/2022. Documentation received accounted for 64 units.</li> </ul> <p>February 2022</p> <ul style="list-style-type: none"> <li>• The Agency billed 120 units of Customized Community Supports (Group) (T2021 HB U8) from 2/3/2022 through 2/9/2022. Documentation received accounted for 116 units.</li> </ul>		
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	<ul style="list-style-type: none"><li>• The Agency billed 120 units of Customized Community Supports (Group) (T2021 HB U8) from 2/10/2022 through 2/16/2022. Documentation received accounted for 108 units.</li><li>• The Agency billed 120 units of Customized Community Supports (Group) (T2021 HB U8) from 2/17/2022 through 2/23/2022. Documentation received accounted for 108 units.</li></ul>		
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MICHELLE LUJAN GRISHAM  
Governor

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

Date: July 28, 2022

To: Melissa McCue, Executive Director

Provider: Mandy's Special Farm dba Mandy's  
Address: 3501 Campus Blvd. NE  
State/Zip: Albuquerque, New Mexico 87106

E-mail Address: [melissa@mandysfarm.org](mailto:melissa@mandysfarm.org)

Board Chair: Ruthie Robbins, Board President  
E-Mail Address: [drmrobb@att.net](mailto:drmrobb@att.net)

Region: Metro  
Survey Date: April 4 - 15, 2022

Program Surveyed: Developmental Disabilities Waiver

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

Survey Type: Routine

Dear Ms. McCue:

The Division of Health Improvement/Quality Management Bureau has received, reviewed and approved the supporting documents you submitted for your Plan of Correction. The documents you provided verified that all previously cited survey Deficiencies have been corrected.

**The Plan of Correction process is now complete.**

**Furthermore, your agency is now determined to be in Compliance with all Conditions of Participation.**

To maintain ongoing compliance with standards and regulations, continue to use the Quality Assurance (self-auditing) processes you described in your Plan of Correction.

Consistent use of these Quality Assurance processes will enable you to identify and promptly respond to problems, enhance your service delivery, and result in fewer deficiencies cited in future QMB surveys.

Thank you for your cooperation with the Plan of Correction process, for striving to come into compliance with standards and regulations, and for helping to provide the health, safety and personal growth of the people you serve.



Sincerely,

*Monica Valdez, BS*

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

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