Dear Ms. Christina Martinez;

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

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

**Partial Compliance with Conditions of Participation**
The following tags are identified as Condition of Participation Level Deficiencies:
- Tag # 1A32 and LS14 / 6L14 Individual Service Plan Implementation
- Tag # 1A09 Medication Delivery Routine Medication Administration

**Plan of Correction:**
The attached Report of Findings identifies the Standard Level and/or Condition of Participation deficiencies found during your agency’s 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.

**Submission of your Plan of Correction:**
Please submit your agency’s Plan of Correction in the space on the two right 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: Amanda Castaneda, Plan of Correction Coordinator**
   1170 North Solano Suite D Las Cruces, New Mexico 88001

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

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

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

**Billing Deficiencies:**
If you have deficiencies noted in this report of findings under the Service Domain: Medicaid Billing/Reimbursement, you must complete a Void/Adjust claims 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: Julie Ann Hill-Clapp
HSD/OIG
Program Integrity Unit
P.O. Box 2348
Santa Fe, New Mexico 87504-2348
```

Or if using UPS, FedEx, DHL (courier mail) send to physical address at:

```
Attention: Julie Ann Hill-Clapp
HSD/OIG
Program Integrity Unit
2025 S. Pacheco Street
Santa Fe, New Mexico 87505
```

Please be advised that there is a one-week lag period for applying payments received by check to Voided/Adjusted 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:

QMB Deputy Bureau Chief  
5301 Central Ave NE Suite #400  
Albuquerque, NM 87108  
Attention: IRF request

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 call the Plan of Correction Coordinator Amanda Castaneda at 575-373-5716 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,

*Florence G. Mulheron, BA*

Florence G. Mulheron, BA  
Team Lead/Healthcare Surveyor  
Division of Health Improvement  
Quality Management Bureau
Survey Process Employed:

Entrance Conference Date: August 3, 2015

Present: The Opportunity Center, Inc.
Clarence Wallace, Incident Coordinator/Quality Enhancement Coordinator

DOH/DHI/QMB
Florence G. Mulheron, BA, Team Lead/Healthcare Surveyor
Deb Russell, BS, Healthcare Surveyor

Exit Conference Date: August 6, 2015

Present: The Opportunity Center, Inc.
Christina Martinez, Director
Teresa Anschutz, Service Coordinator
Alexandra Elliott, Administrative Assistant
Doug Moots, Director of Nursing
Apryl Stickels, Finance Coordinator
Clarence Wallace, Incident Coordinator/Quality Enhancement Coordinator

DOH/DHI/QMB
Florence G. Mulheron, BA, Team Lead/Healthcare Surveyor
Deb Russell, BS, Healthcare Surveyor

DDSD - Southwest Regional Office
Angie Brooks, Generalist

Administrative Locations Visited Number: 1

Total Sample Size Number: 7
0 - Jackson Class Members
7 - Non-Jackson Class Members
6 - Supported Living
6 - Customized Community Supports
2 - Community Integrated Employment Services
1 - Customized In-Home Supports

Total Homes Visited Number: 4
 Supported Living Homes Visited Number: 4

Note: The following Individuals share a SL residence:
➢ #2, 3
➢ #5, 6

Persons Served Records Reviewed Number: 7

Persons Served Interviewed Number: 7
Direct Support Personnel Interviewed Number:  9
Direct Support Personnel Records Reviewed Number:  29
Service Coordinator Records Reviewed Number:  2

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
         MFEAD – NM Attorney General
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 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 575-373-5716 or email at AmandaE.Castaneda@state.nm.us. Requests for technical assistance must be requested through your Regional DDSD Office.

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

Instructions for Completing Agency POC:

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

If a deficiency has already been corrected, 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 Plan of Correction must address the six required Center for Medicare and Medicaid Services (CMS) core elements to address each deficiency cited in the Report of Findings:
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 individuals in similar situations.
3. What QA measures will be put into place or systemic changes made to ensure that the deficient practice will not recur.
4. Indicate how the agency plans to monitor its performance to make sure that solutions are sustained. The agency must develop a QA plan for ensuring that correction is achieved and sustained. This QA plan must be implemented, and the corrective action evaluated for its effectiveness. The plan of correction is integrated into the agency quality assurance system; and
5. Include dates when corrective action will be completed. The corrective action completion dates must be acceptable to the State.
6. The POC must be signed and dated by the agency director or other authorized official.

The following details should be considered when developing your Plan of Correction:
- Details about how and when Consumer, Personnel and Residential 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, as they are being used, and after they are completed;
- Your processes for ensuring that all staff are trained in Core Competencies, Abuse, Neglect and Exploitation Reporting, and Individual-Specific service requirements, etc.;
- How accuracy in Billing/Reimbursement documentation is assured;
- How health, safety is assured;
- For Case Management Providers, how Individual Specific Plans are reviewed to verify they meet requirements, how the timeliness of 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, Amanda Castaneda at 575-373-5716 or email at AmandaE.Castaneda@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 Amanda Castaneda, POC Coordinator in any of the following ways:
   a. Electronically at AmandaE.Castaneda@state.nm.us (preferred method)
   b. Fax to 575-528-5019, or
   c. Mail to POC Coordinator, 1170 North Solano Ste D, Las Cruces, New Mexico 88001
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.

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 the documents do not contain protected Health information (PHI) the preferred method is that you 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. In addition to this, we ask that you submit:
   • Evidence of an internal audit of billing/reimbursement conducted for a sample of individuals and timeframes of your choosing to verify POC implementation;
   • Copies of “void and adjust” forms submitted to Xerox State Healthcare, LLC to correct all unjustified units identified and submitted for payment during your internal audit.

Revisions, Modifications or Extensions to your Plan of Correction (post QMB approval) must be made in writing and submitted to the Plan of Correction Coordinator, prior to the due date and are approved on a case-by-case basis. No changes may be made to your POC or the timeframes for implementation without written approval of the POC Coordinator.
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 state and federal regulations. QMB has grouped the CMS assurances into five Service Domains: Level of Care; Plan of Care; 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 Management 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 in the QMB Report of Findings. 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.

Within the QMB Service Domains there are fundamental regulations, standards, or policies with which a provider must be in essential compliance in order to ensure the health and welfare of individuals served known as Conditions of Participation (CoPs).

The Determination of Compliance for each service type is based on a provider’s compliance with CoPs in three (3) Service Domains.

Case Management Services:
- Level of Care
- Plan of Care
- Qualified Providers

Community Inclusion Supports/ Living Supports:
- Qualified Provider
- Plan of Care
- Health, Welfare and Safety

**Conditions of Participation (CoPs)**

A CoP is an identified fundamental regulation, standard, or policy with which a provider must be in compliance in order to ensure the health and welfare of individuals served. CoPs are based on the Centers for Medicare and Medicaid Services, Home and Community-Based Waiver required assurances. A provider must be in compliance with CoPs to participate as a waiver provider.

QMB surveyors use professional judgment when reviewing the critical elements of each standard and regulation to determine when non-compliance with a standard level deficiency rises to the level of a CoP out of compliance. Only some deficiencies can rise to the level of a CoP (See the next section for a list of CoPs). The QMB survey team analyzes the relevant finding in terms of scope, actual harm or potential for harm, unique situations, patterns of performance, and other factors to determine if there is the potential for a negative outcome which would rise to the level of a CoP. A Standard level deficiency becomes a CoP out of compliance when the team’s analysis establishes that there is an identified potential for

QMB Report of Findings – The Opportunity Center, Inc. – Southwest Region – August 3 – 6, 2015
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significant harm or actual harm. It is then cited as a CoP out of compliance. If the deficiency does not rise to the level of a CoP out of compliance, it is cited as a Standard Level Deficiency.

The Division of Health Improvement (DHI) and the Developmental Disabilities Supports Division (DDSD) collaborated to revise the current Conditions of Participation (CoPs). There are seven Conditions of Participation in which providers must be in compliance.

CoPs and Service Domains for Case Management Supports are as follows:

**Service Domain: Level of Care**
Condition of Participation:
1. **Level of Care**: The Case Manager shall complete all required elements of the Long Term Care Assessment Abstract (LTCAA) to ensure ongoing eligibility for waiver services.

**Service Domain: Plan of Care**
Condition of Participation:
2. **Individual Service Plan (ISP) Creation and Development**: Each individual shall have an ISP. The ISP shall be developed in accordance with DDSD regulations and standards and is updated at least annually or when warranted by changes in the individual’s needs.

Condition of Participation:
3. **ISP Monitoring and Evaluation**: The Case Manager shall ensure the health and welfare of the individual through monitoring the implementation of ISP desired outcomes.

CoPs and Service Domain for ALL Service Providers is as follows:

**Service Domain: Qualified Providers**
Condition of Participation:
4. **Qualified Providers**: Agencies shall ensure support staff has completed criminal background screening and all mandated trainings as required by the DDSD.

CoPs and Service Domains for Living Supports and Inclusion Supports are as follows:

**Service Domain: Plan of Care**
Condition of Participation:
5. **ISP Implementation**: Services provided shall be consistent with the components of the ISP and implemented to achieve desired outcomes.

**Service Domain: Health, Welfare and Safety**
Condition of Participation:
6. **Individual Health, Safety and Welfare**: (Safety) Individuals have the right to live and work in a safe environment.

Condition of Participation:
7. **Individual Health, Safety and Welfare (Healthcare Oversight)**: The provider shall support individuals to access needed healthcare services in a timely manner. Nursing, healthcare services and healthcare oversight shall be available and provided as needed to address individuals’ health, safety and welfare.
QMB Determinations of Compliance

Compliance with Conditions of Participation
The QMB determination of *Compliance with Conditions of Participation* indicates that a provider is in compliance with all Conditions of Participation, (CoP). 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 with Conditions of Participation, the provider must be in compliance with all Conditions of Participation in all relevant Service Domains. The agency may also have Standard level deficiencies (deficiencies which are not at the condition level) out of compliance in any of the Service Domains.

Partial-Compliance with Conditions of Participation
The QMB determination of *Partial-Compliance with Conditions of Participation* indicates that a provider is out of compliance with Conditions of Participation in one (1) to two (2) Service Domains. The agency may have one or more Condition level tags within a Service Domain. 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. The agency may also have Standard level deficiencies (deficiencies which are not at the condition level) in any of the Service Domains.

Providers receiving a *repeat* determination of Partial-Compliance for repeat deficiencies at the level of a Condition in any Service Domain may be referred by the Quality Management Bureau to the Internal Review Committee (IRC) for consideration of remedies and possible actions or sanctions.

Non-Compliance with Conditions of Participation
The QMB determination of *Non-Compliance with Conditions of Participation* indicates a provider is significantly out of compliance with Conditions of Participation in multiple Service Domains. The agency may have one or more Condition level tags in each of 3 relevant Service Domains. 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. The agency may also have Standard level deficiencies (deficiencies which are not at the condition level) in any of the Service Domains.

Providers receiving a *repeat* determination of Non-Compliance will be referred by Quality Management Bureau to the Internal Review Committee (IRC) for consideration of remedies and possible actions or sanctions.
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 Deputy Bureau Chief within 10 business days of receipt of the final Report of Findings.
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: http://dhi.health.state.nm.us/qmb
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, Crystal Lopez-Beck at Crystal.Lopez-Beck@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.
## Standard of Care

<table>
<thead>
<tr>
<th>Service Domain: Service Plans: ISP Implementation</th>
<th>Deficiencies</th>
</tr>
</thead>
<tbody>
<tr>
<td>Services are delivered in accordance with the service plan, including type, scope, amount, duration and frequency specified in the service plan.</td>
<td></td>
</tr>
</tbody>
</table>

### Tag # 1A08

**Agency Case File**

**Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013**

**Chapter 5 (CIES) 3. Agency Requirements**

**H. Consumer Records Policy:** All Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Consumer Records Policy. Additional documentation that is required to be maintained at the administrative office includes:

1. Vocational Assessments that are of quality and contain content acceptable to DVR and DDSD;
2. Career Development Plans as incorporated in the ISP; and
3. Documentation of evidence that services provided under the DDW are not otherwise available under the Rehabilitation Act of 1973 (DVR).

**Chapter 6 (CCS) 3. Agency Requirements**

**G. Consumer Records Policy:** All Provider Agencies shall maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix.

<table>
<thead>
<tr>
<th>Standard Level Deficiency</th>
<th>Agency Plan of Correction, On-going QA/QI and Responsible Party</th>
</tr>
</thead>
<tbody>
<tr>
<td>Based on record review, the Agency did not maintain a complete and confidential case file at the administrative office for 5 of 7 individuals. Review of the Agency individual case files revealed the following items were not found, incomplete, and/or not current:</td>
<td></td>
</tr>
</tbody>
</table>
| • ISP budget forms MAD 046  
  ○ Not Found (#2, 7) |
| • Current Emergency and Personal Identification Information  
  ○ Did not contain the name and contact information of relatives and/or guardian. (#3) |
| • Behavior Crisis Intervention Plan (#2, 3, 5) |
| • Speech Therapy Plan (#6) |

**Provider:**

State your Plan of Correction for the deficiencies cited in this tag here: →

**Provider:**

Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →

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policy. Additional documentation that is required to be maintained at the administrative office includes:
  1. Vocational Assessments (if applicable) that are of quality and contain content acceptable to DVR and DDSD.

Chapter 7 (CIHS) 3. Agency Requirements:
E. Consumer Records Policy: All Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix policy.

Chapter 11 (FL) 3. Agency Requirements:
D. Consumer Records Policy: All Family Living Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix policy.

Chapter 12 (SL) 3. Agency Requirements:
D. Consumer Records Policy: All Living Supports - Supported Living Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix policy.

Chapter 13 (IMLS) 2. Service Requirements:
C. Documents to be maintained in the agency administrative office, include: (This is not an all-inclusive list refer to standard as it includes other items)
  • Emergency contact information;
  • Personal identification;
  • ISP budget forms and budget prior authorization;
  • ISP with signature page and all applicable assessments, including teaching and support strategies, Positive Behavior Support Plan
(PBSP), Behavior Crisis Intervention Plan (BCIP), or other relevant behavioral plans, Medical Emergency Response Plan (MERP), Healthcare Plan, Comprehensive Aspiration Risk Management Plan (CARMP), and Written Direct Support Instructions (WDSI);
• Dated and signed evidence that the individual has been informed of agency grievance/complaint procedure at least annually, or upon admission for a short term stay;
• Copy of Guardianship or Power of Attorney documents as applicable;
• Behavior Support Consultant, Occupational Therapist, Physical Therapist and Speech-Language Pathology progress reports as applicable, except for short term stays;
• Written consent by relevant health decision maker and primary care practitioner for self-administration of medication or assistance with medication from DSP as applicable;
• Progress notes written by DSP and nurses;
• Signed secondary freedom of choice form;
• Transition Plan as applicable for change of provider in past twelve (12) months.

DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION (DDSD): Director’s Release: Consumer Record Requirements eff. 1/1/2012

III. Requirement Amendments(s) or Clarifications:
A. All case management, living supports, customized in-home supports, community integrated employment and customized community supports providers must maintain records for individuals served through DD Waiver in accordance with the Individual Case File Matrix incorporated in this director’s release.

H. Readily accessible electronic records are accessible, including those stored through the Therap web-based system.

CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS: D. Provider Agency Case File for the Individual: All Provider Agencies shall maintain at the administrative office a confidential case file for each individual. Case records belong to the individual receiving services and copies shall be provided to the receiving agency whenever an individual changes providers. The record must also be made available for review when requested by DOH, HSD or federal government representatives for oversight purposes. The individual’s case file shall include the following requirements:

1. Emergency contact information, including the individual’s address, telephone number, names and telephone numbers of relatives, or guardian or conservator, physician’s name(s) and telephone number(s), pharmacy name, address and telephone number, and health plan if appropriate;
2. The individual’s complete and current ISP, with all supplemental plans specific to the individual, and the most current completed Health Assessment Tool (HAT);
3. Progress notes and other service delivery documentation;
4. Crisis Prevention/Intervention Plans, if there are any for the individual;
5. A medical history, which shall include at least demographic data, current and past medical diagnoses including the cause (if known) of the developmental disability, psychiatric diagnoses, allergies (food, environmental, medications), immunizations, and most recent physical exam;
6. When applicable, transition plans completed for individuals at the time of discharge from Fort Stanton Hospital or Los Lunas Hospital and Training School; and
7. Case records belong to the individual receiving services and copies shall be provided to the individual upon request.
8. The receiving Provider Agency shall be provided at a minimum the following records
whenever an individual changes provider agencies:
(a) Complete file for the past 12 months;
(b) ISP and quarterly reports from the current and prior ISP year;
(c) Intake information from original admission to services; and
(d) When applicable, the Individual Transition Plan at the time of discharge from Los Lunas Hospital and Training School or Ft. Stanton Hospital.

NMAC 8.302.1.17 RECORD KEEPING AND DOCUMENTATION REQUIREMENTS: A provider must maintain all the records necessary to fully disclose the nature, quality, amount and medical necessity of services furnished to an eligible recipient who is currently receiving or who has received services in the past.

B. Documentation of test results: Results of tests and services must be documented, which includes results of laboratory and radiology procedures or progress following therapy or treatment.
<table>
<thead>
<tr>
<th>Tag # 1A32 and LS14 / 6L14</th>
<th>Individual Service Plan Implementation</th>
<th>Condition of Participation Level Deficiency</th>
<th>Provider:</th>
</tr>
</thead>
<tbody>
<tr>
<td>NMAC 7.26.5.16.C and D Development of the ISP. Implementation of the ISP. The ISP shall be implemented according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan.</td>
<td></td>
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<tr>
<td><strong>C.</strong> The IDT shall review and discuss information and recommendations with the individual, with the goal of supporting the individual in attaining desired outcomes. The IDT develops an ISP based upon the individual's personal vision statement, strengths, needs, interests and preferences. The ISP is a dynamic document, revised periodically, as needed, and amended to reflect progress towards personal goals and achievements consistent with the individual's future vision. This regulation is consistent with standards established for individual plan development as set forth by the commission on the accreditation of rehabilitation facilities (CARF) and/or other program accreditation approved and adopted by the developmental disabilities division and the department of health. It is the policy of the developmental disabilities division (DDD), that to the extent permitted by funding, each individual receive supports and services that will assist and encourage independence and productivity in the community and attempt to prevent regression or loss of current capabilities. Services and supports include specialized and/or generic services, training, education and/or treatment as determined by the IDT and documented in the ISP.</td>
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<tr>
<td>After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur.</td>
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<tr>
<td>Based on record review, the Agency did not implement the ISP according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan for 5 of 7 individuals.</td>
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<tr>
<td>As indicated by Individuals ISP the following was found with regards to the implementation of ISP Outcomes:</td>
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<td></td>
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<tr>
<td><strong>Administrative Files Reviewed:</strong></td>
<td></td>
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<td></td>
</tr>
<tr>
<td><strong>Supported Living Data Collection/Data Tracking/Progress with regards to ISP Outcomes:</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Individual #3</td>
<td></td>
<td></td>
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<tr>
<td>• According to the Live Outcome; Action Step for &quot;With all needed supports … will do routine maintenance upgrades to his property” is to be completed 1 time per week, evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 4/2015 and 6/2015.</td>
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<td></td>
<td></td>
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<tr>
<td>Individual #4</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>• None found regarding: Live Outcome/Action Step: &quot;... will have support to research, purchase, and try new foods&quot; for 10/2014 - 11/2014. Action step is to be completed 1 time per week.</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Provider: State your Plan of Correction for the deficiencies cited in this tag here: →</td>
<td></td>
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<tr>
<td>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →</td>
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</tbody>
</table>
D. The intent is to provide choice and obtain opportunities for individuals to live, work and play with full participation in their communities. The following principles provide direction and purpose in planning for individuals with developmental disabilities.

[05/03/94; 01/15/97; Recompiled 10/31/01]

<p>| | |</p>
<table>
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<tr>
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</thead>
<tbody>
<tr>
<td></td>
<td>Agency’s Outcomes/Action Steps are as follows:</td>
</tr>
<tr>
<td></td>
<td>“… will have support to store, display, organize and make livable space in 2 rooms at his house.”</td>
</tr>
<tr>
<td></td>
<td>Annual ISP (10/2014 - 10/2015) Outcomes/Action Steps are as follows:</td>
</tr>
<tr>
<td></td>
<td>“… will have support to research, purchase and try new foods.”</td>
</tr>
<tr>
<td></td>
<td>Individual #5</td>
</tr>
<tr>
<td></td>
<td>None found regarding: Live Outcome/Action Step: “… will have support to purchase items, prepare dishes and deliver them to ARM” for 11/2014 - 5/2015. Action step is to be completed 1 times per week.</td>
</tr>
<tr>
<td></td>
<td>Individual #6</td>
</tr>
<tr>
<td></td>
<td>None found regarding: Live Outcome/Action Step: “… will follow appropriate conversation steps is to be completed 5 times per week” for 11/2015. Action step is to be completed 5 times per week.</td>
</tr>
<tr>
<td></td>
<td>According to the Live Outcome; Action Step for “… will follow appropriate conversation steps” is to be completed 5 x per week, evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 12/2014 - 7/2015.</td>
</tr>
</tbody>
</table>
Individual #7
- According to the Live Outcome; Action Step for “… will put her articles in her scrapbook” is to be completed 2 times per month, evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 3/2015 - 4/2015.

Customized Community Supports Data Collection/Data Tracking/Progress with regards to ISP Outcomes:

Individual #3
- No Outcomes or DDSD exemption/decision justification found for Customized Community Support Services. As indicated by NMAC 7.26.5.14 “Outcomes are required for any life area for which the individual receives services funded by the developmental disabilities Medicaid waiver.”

Individual #4
- None found regarding: Work/Education/Volunteer Outcome/Action Step: “… will select and consume a meal” for 10/2014 - 11/2014. Action step is to be completed 1 time per week.
- None found regarding: Work/Education/Volunteer Outcome/Action Step: “… will have assistance to prepare healthy delicious meals” for 10/2014 - 11/2014. Action step is to be completed 2 times per week.
- None found regarding: Work/Education/Volunteer Outcome/Action Step: “… will critique and submit his critique for publication” for 10/2014 - 11/2014.
Action step is to be completed 1 time per week.


**Agency’s Outcomes/Action Steps are as follows:**
- “…will have support to organize, invite and host card games.”

**Annual ISP (10/2014 - 10/2015) Outcomes/Action Steps are as follows:**
- “…will select and consume a meal.”
- “…will have assistance to prepare healthy delicious meals.”
- “…will critique and submit his critique for publication.”

- According to the Work/Education/Volunteer Outcome; Action Step for “Will select and consume a meal” is to be completed 1 time per week, evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 6/2015.

- According to the Work/Education/Volunteer Outcome; Action Step for “… will have assistance to prepare healthy delicious meals” is to be completed 2 time per week, evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 6/2015 – 7/2015.
- According to the Work/Education/Volunteer Outcome; Action Step for “Will critique and submit his critique for publication” is to be completed 1 time per week, evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 6/2015.

- None found regarding: Relationship/Have Fun Outcome/Action Step: “With all needed support … will research music of the 80s and find songs to download” for 10/2014 - 2/2015.

Individual #5

- None found regarding: Work/Education/Volunteer Outcome/Action Step: “… will have support to purchase needed items meet with local artists learn different styles and create different pieces of art work” for 11/2014 - 5/2015.

- None found regarding: Work/Education/Volunteer Outcome/Action Step: “… will have support to select a variety subjects to do editorials on and submit the editorials for publishing” for 5/2015 - 7/2015.

- None found regarding: Relationship/Have Fun Outcome/Action Step: “… will have support to purchase needed items, meet with local artists, learn different styles, and create different pieces of artwork” for 5/2015 - 7/2015.

Individual #6
• None found regarding: Relationship/Have Fun Outcome/Action Step: "… will select and participate" for 11/2015.

• According to the Relationship/Have Fun Outcome; Action Step for "… will select and participate," is to be completed "every other weekend" evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 1/2015 - 7/2015.

Individual #7
• None found regarding: Relationship/Have Fun Outcome/Action Step: "… will go out into the community to a place of her choosing" for 3/2015 - 6/2015.

Community Integrated Employment Services Data Collection/Data Tracking/Progress with regards to ISP Outcomes:

Individual #3
• According to the Work/Education/Volunteer Outcome; Action Step for "Staff will assist … in identifying tasks that he is interested in and capable of doing so that he proves his value at the workplace" is to be completed 1 time per week, evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 3/2014 - 6/2015.
### Standard Level Deficiency

Based on record review, the Agency did not complete written status reports as required for 1 of 6 individuals receiving Inclusion Services. Review of the Agency individual case files revealed the following items were not found, and/or incomplete:

**Customized Community Supports Semi-Annual Reports**

**Community Integrated Employment Services Semi-Annual Reports**

### Provider:
State your Plan of Correction for the deficiencies cited in this tag here: →

**Provider:**
Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →
meeting unless changes requiring team input need to be made (e.g., adding more hours to the Community Integrated Employment budget);

b. Written annual updates to the ISP work/learn action plan to DDSD;

2. VAP to the case manager if completed externally to the ISP;

3. Initial ISP reflecting the Vocational Assessment or the annual ISP with the updated VAP integrated or a copy of an external VAP if one was completed to DDSD;

4. Quarterly Community Integrated Employment Wage and Hour Reports for individuals employed and in job development to DDSD based on the DDSD fiscal year; and

a. Data related to the requirements of the Performance Contract to DDSD quarterly.

CHAPTER 6 (CCS) 3. Agency Requirements: H. Reporting Requirements: The Customized Community Supports Provider Agency shall submit the following:

1. Semi-annual progress reports one hundred ninety (190) days following the date of the annual ISP, and 14 days prior to the annual IDT meeting:

a. Identification of and implementation of a Meaningful Day definition for each person served;

b. Documentation for each date of service delivery summarizing the following:
   i. Choice based options offered throughout the day; and
ii. Progress toward outcomes using age appropriate strategies specified in each individual’s action steps in the ISP, and associated support plans/WDSI.

c. Record of personally meaningful community inclusion activities; and

d. Written updates, to the ISP Work/Learn Action Plan annually or as necessary due to change in work goals. These updates do not require an IDT meeting unless changes requiring team input need to be made.

e. Data related to the requirements of the Performance Contract to DDSD quarterly.


CHAPTER 5 IV. COMMUNITY INCLUSION SERVICES PROVIDER AGENCY REQUIREMENTS

E. Provider Agency Reporting Requirements:

All Community Inclusion Provider Agencies are required to submit written quarterly status reports to the individual’s Case Manager no later than fourteen (14) calendar days following the end of each quarter. In addition to reporting required by specific Community Access, Supported Employment, and Adult Habilitation Standards, the quarterly reports shall contain the following written documentation:

(1) Identification and implementation of a meaningful day definition for each person served;

(2) Documentation summarizing the following:

(a) Daily choice-based options; and
(b) Daily progress toward goals using age-appropriate strategies specified in each individual's action plan in the ISP.
(3) Significant changes in the individual's routine or staffing;
(4) Unusual or significant life events;
(5) Quarterly updates on health status, including changes in medication, assistive technology needs and durable medical equipment needs;
(6) Record of personally meaningful community inclusion;
(7) Success of supports as measured by whether or not the person makes progress toward his or her desired outcomes as identified in the ISP; and
(8) Any additional reporting required by DDSD.
<table>
<thead>
<tr>
<th>Tag # LS14 / 6L14</th>
<th>Standard Level Deficiency</th>
<th>Provider: State your Plan of Correction for the deficiencies cited in this tag here: →</th>
</tr>
</thead>
<tbody>
<tr>
<td>Residential Case File</td>
<td>Based on record review, the Agency did not maintain a complete and confidential case file in the residence for 5 of 6 Individuals receiving Supported Living Services. Review of the residential individual case files revealed the following items were not found, incomplete, and/or not current:</td>
<td></td>
</tr>
<tr>
<td></td>
<td><strong>ISP Teaching and Support Strategies</strong></td>
<td><strong>Provider:</strong> Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →</td>
</tr>
<tr>
<td></td>
<td>o Individual #3 - TSS not found for the following Action Steps:</td>
<td></td>
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<tr>
<td></td>
<td>o Live Outcome Statement</td>
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<tr>
<td></td>
<td>o “… will add at least 1 new plant each season.”</td>
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<td></td>
<td>o “… will care for and nurture plants and harvest any foods.”</td>
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<tr>
<td></td>
<td><strong>Positive Behavioral Plan (#2, 3)</strong></td>
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<td></td>
<td><strong>Behavior Crisis Intervention Plan (#2, 3, 5)</strong></td>
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<td></td>
<td><strong>Physical Therapy Plan (#3)</strong></td>
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<td></td>
<td><strong>Healthcare Passport (#2, 3, 5)</strong></td>
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<td></td>
<td><strong>Special Health Care Needs</strong></td>
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<tr>
<td></td>
<td>o Nutritional Plan (#2, 4, 7)</td>
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<td></td>
<td><strong>Progress Notes/Daily Contacts Logs:</strong></td>
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<tr>
<td></td>
<td>o Individual #3 - None found for 8/3/2015.</td>
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<td></td>
<td>o Individual #4 - None found for 8/1 – 2, 2015.</td>
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<tr>
<td><strong>CHAPTER 11 (FL) 3. Agency Requirements</strong></td>
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<tr>
<td><strong>C. Residence Case File:</strong> The Agency must maintain in the individual’s home a complete and current confidential case file for each individual. Residence case files are required to comply with the DDSD Individual Case File Matrix policy.</td>
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<tr>
<td><strong>CHAPTER 12 (SL) 3. Agency Requirements</strong></td>
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<tr>
<td><strong>C. Residence Case File:</strong> The Agency must maintain in the individual’s home a complete and current confidential case file for each individual. Residence case files are required to comply with the DDSD Individual Case File Matrix policy.</td>
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<tr>
<td><strong>CHAPTER 13 (IMLS) 2. Service Requirements</strong></td>
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<tr>
<td><strong>B.1. Documents To Be Maintained In The Home:</strong></td>
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<tr>
<td>a. Current Health Passport generated through the e-CHAT section of the Therap website and printed for use in the home in case of disruption in internet access;</td>
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<tr>
<td>b. Personal identification;</td>
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<tr>
<td>c. Current ISP with all applicable assessments, teaching and support strategies, and as applicable for the consumer, PBSP, BCIP, MERP, health care plans, CARMPs, Written Therapy Support Plans, and any other plans (e.g. PRN Psychotropic Medication Plans) as applicable;</td>
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<tr>
<td>d. Dated and signed consent to release information forms as applicable;</td>
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<tr>
<td>e. Current orders from health care practitioners;</td>
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<tr>
<td>f. Documentation and maintenance of accurate medical history in Therap website;</td>
<td></td>
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<tr>
<td>g. Medication Administration Records for the current month;</td>
<td></td>
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<tr>
<td>h. Record of medical and dental appointments for the current year, or during the period of stay for Based on record review, the Agency did not maintain a complete and confidential case file in the residence for 5 of 6 Individuals receiving Supported Living Services. Review of the residential individual case files revealed the following items were not found, incomplete, and/or not current: <strong>ISP Teaching and Support Strategies</strong> o Individual #3 - TSS not found for the following Action Steps: o Live Outcome Statement ➢ “… will add at least 1 new plant each season.” ➢ “… will care for and nurture plants and harvest any foods.” <strong>Positive Behavioral Plan (#2, 3)</strong> <strong>Behavior Crisis Intervention Plan (#2, 3, 5)</strong> <strong>Physical Therapy Plan (#3)</strong> <strong>Healthcare Passport (#2, 3, 5)</strong> <strong>Special Health Care Needs</strong> o Nutritional Plan (#2, 4, 7) <strong>Progress Notes/Daily Contacts Logs:</strong> o Individual #3 - None found for 8/3/2015. o Individual #4 - None found for 8/1 – 2, 2015.</td>
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</table>
short term stays, including any treatment provided;
i. Progress notes written by DSP and nurses;
j. Documentation and data collection related to ISP implementation;
k. Medicaid card;
l. Salud membership card or Medicare card as applicable; and
m. A Do Not Resuscitate (DNR) document and/or Advanced Directives as applicable.

DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION (DDSD): Director’s Release: Consumer Record Requirements eff. 11/1/2012
III. Requirement Amendments(s) or Clarifications:
A. All case management, living supports, customized in-home supports, community integrated employment and customized community supports providers must maintain records for individuals served through DD Waiver in accordance with the Individual Case File Matrix incorporated in this director’s release.

H. Readily accessible electronic records are accessible, including those stored through the Therap web-based system.

CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS
A. Residence Case File: For individuals receiving Supported Living or Family Living, the Agency shall maintain in the individual’s home a complete and current confidential case file for each individual. For individuals receiving Independent Living Services, rather than maintaining this file at the individual’s home, the complete and current confidential case file for each individual shall be maintained at the agency’s administrative site. Each file shall include the following:
(1) Complete and current ISP and all supplemental plans specific to the individual;
| (2) Complete and current Health Assessment Tool; |
| (3) Current emergency contact information, which includes the individual’s address, telephone number, names and telephone numbers of residential Community Living Support providers, relatives, or guardian or conservator, primary care physician's name(s) and telephone number(s), pharmacy name, address and telephone number and dentist name, address and telephone number, and health plan; |
| (4) Up-to-date progress notes, signed and dated by the person making the note for at least the past month (older notes may be transferred to the agency office); |
| (5) Data collected to document ISP Action Plan implementation |
| (6) Progress notes written by direct care staff and by nurses regarding individual health status and physical conditions including action taken in response to identified changes in condition for at least the past month; |
| (7) Physician’s or qualified health care providers written orders; |
| (8) Progress notes documenting implementation of a physician’s or qualified health care provider’s order(s); |
| (9) Medication Administration Record (MAR) for the past three (3) months which includes: |
| (a) The name of the individual; |
| (b) A transcription of the healthcare practitioners prescription including the brand and generic name of the medication; |
| (c) Diagnosis for which the medication is prescribed; |
| (d) Dosage, frequency and method/route of delivery; |
| (e) Times and dates of delivery; |
| (f) Initials of person administering or assisting with medication; and |
(g) An explanation of any medication irregularity, allergic reaction or adverse effect.

(h) For PRN medication an explanation for the use of the PRN must include:
   (i) Observable signs/symptoms or circumstances in which the medication is to be used, and
   (ii) Documentation of the effectiveness/result of the PRN delivered.

(i) A MAR is not required for individuals participating in Independent Living Services who self-administer their own medication. However, when medication administration is provided as part of the Independent Living Service a MAR must be maintained at the individual’s home and an updated copy must be placed in the agency file on a weekly basis.

(10) Record of visits to healthcare practitioners including any treatment provided at the visit and a record of all diagnostic testing for the current ISP year; and

(11) Medical History to include: demographic data, current and past medical diagnoses including the cause (if known) of the developmental disability and any psychiatric diagnosis, allergies (food, environmental, medications), status of routine adult health care screenings, immunizations, hospital discharge summaries for past twelve (12) months, past medical history including hospitalizations, surgeries, injuries, family history and current physical exam.
<table>
<thead>
<tr>
<th>Tag # LS17 / 6L17 Reporting Requirements (Community Living Reports)</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>7.26.5.17 DEVELOPMENT OF THE INDIVIDUAL SERVICE PLAN (ISP) - DISSEMINATION OF THE ISP, DOCUMENTATION AND COMPLIANCE: C. Objective quantifiable data reporting progress or lack of progress towards stated outcomes, and action plans shall be maintained in the individual’s records at each provider agency implementing the ISP. Provider agencies shall use this data to evaluate the effectiveness of services provided. Provider agencies shall submit to the case manager data reports and individual progress summaries quarterly, or more frequently, as decided by the IDT. These reports shall be included in the individual’s case management record, and used by the team to determine the ongoing effectiveness of the supports and services being provided. Determination of effectiveness shall result in timely modification of supports and services as needed.</td>
<td>Based on record review, the Agency did not complete written status reports for 1 of 6 individuals receiving Living Services. Review of the Agency individual case files revealed the following items were not found, and/or incomplete: <strong>Supported Living Semi-Annual Reports:</strong> • Individual #3 - None found for 1/2015 - 4/2015. <em>(ISP Meeting 4/28/2015)</em></td>
</tr>
<tr>
<td>Provider: State your Plan of Correction for the deficiencies cited in this tag here: → Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →</td>
<td></td>
</tr>
</tbody>
</table>
The semi-annual reports must contain the following written documentation:

a. Name of individual and date on each page;

b. Timely completion of relevant activities from ISP Action Plans;

c. Progress towards desired outcomes in the ISP accomplished during the past six month;

d. Significant changes in routine or staffing;

e. Unusual or significant life events, including significant change of health condition;

f. Data reports as determined by IDT members; and

g. Signature of the agency staff responsible for preparing the reports.

**CHAPTER 12 (SL) 3. Agency Requirements:**

**E. Living Supports- Supported Living Service Provider Agency Reporting Requirements:**

1. **Semi-Annual Reports:** Supported Living providers must submit written semi-annual status reports to the individual’s Case Manager and other IDT Members no later than one hundred ninety (190) calendar days after the ISP effective date. When reports are developed in any other language than English, it is the responsibility of the provider to translate the reports into English. The semi-annual reports must contain the following written documentation:

   a. Name of individual and date on each page;

   b. Timely completion of relevant activities from ISP Action Plans;
c. Progress towards desired outcomes in the ISP accomplished during the past six (6) months;

d. Significant changes in routine or staffing;

e. Unusual or significant life events, including significant change of health condition;

f. Data reports as determined by IDT members;

and

g. Signature of the agency staff responsible for preparing the reports.

CHAPTER 13 (IMLS) 3. Agency Requirements: F. Quality Assurance/Quality Improvement (QA/QI) Program:

4. Intensive Medical Living Services providers shall submit a written semi-annual (non-nursing) status report to the individual's case manager and other IDT members no later than the one hundred ninetieth (190th) day following ISP effective date. These semi-annual status reports shall contain at least the following information:

a. Status of completion of ISP Action Plans and associated support plans and/or WDSI;

b. Progress towards desired outcomes;

c. Significant changes in routine or staffing;

d. Unusual or significant life events; and

e. Data reports as determined by the IDT members;
<table>
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<tbody>
<tr>
<td><strong>CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS</strong></td>
</tr>
<tr>
<td>D. Community Living Service Provider Agency Reporting Requirements: <strong>All Community Living Support providers shall submit written quarterly status reports to the individual’s Case Manager and other IDT Members no later than fourteen (14) days following the end of each ISP quarter.</strong> The quarterly reports shall contain the following written documentation:</td>
</tr>
<tr>
<td>(1) Timely completion of relevant activities from ISP Action Plans</td>
</tr>
<tr>
<td>(2) Progress towards desired outcomes in the ISP accomplished during the quarter;</td>
</tr>
<tr>
<td>(3) Significant changes in routine or staffing;</td>
</tr>
<tr>
<td>(4) Unusual or significant life events;</td>
</tr>
<tr>
<td>(5) Updates on health status, including medication and durable medical equipment needs identified during the quarter; and</td>
</tr>
<tr>
<td>(6) Data reports as determined by IDT members.</td>
</tr>
</tbody>
</table>
**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.

<table>
<thead>
<tr>
<th>Tag # 1A20 Direct Support Personnel Training</th>
<th>Standard Level Deficiency</th>
<th>Agency Plan of Correction, On-going QA/QI and Responsible Party</th>
<th>Date Due</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy - Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007 - II. POLICY STATEMENTS:</strong></td>
<td>Based on record review, the Agency did not ensure Orientation and Training requirements were met for 25 of 29 Direct Support Personnel.</td>
<td><strong>Provider:</strong> State your Plan of Correction for the deficiencies cited in this tag here: →</td>
<td></td>
</tr>
<tr>
<td>A. Individuals shall receive services from competent and qualified staff.</td>
<td>Review of Direct Support Personnel training records found no evidence of the following required DOH/DDSD trainings and certification being completed:</td>
<td><strong>Provider:</strong> Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →</td>
<td></td>
</tr>
<tr>
<td>B. Staff shall complete individual-specific (formerly known as “Addendum B”) training requirements in accordance with the specifications described in the individual service plan (ISP) of each individual served.</td>
<td>• Pre- Service (DSP #220, 223, 224, 225, 226)</td>
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</tr>
<tr>
<td>C. Staff shall complete training on DOH-approved incident reporting procedures in accordance with 7 NMAC 1.13.</td>
<td>• Foundation for Health and Wellness (DSP #220, 223, 224, 225, 226)</td>
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</tr>
<tr>
<td>D. Staff providing direct services shall complete training in universal precautions on an annual basis. The training materials shall meet Occupational Safety and Health Administration (OSHA) requirements.</td>
<td>• Person-Centered Planning (1-Day) (DSP #220)</td>
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</tr>
<tr>
<td>E. Staff providing direct services shall maintain certification in first aid and CPR. The training materials shall meet OSHA requirements/guidelines.</td>
<td>• First Aid (DSP #202, 203, 205, 206, 208, 211, 213, 214, 215, 218, 219, 220, 227, 228, 229)</td>
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</tr>
<tr>
<td>F. Staff who may be exposed to hazardous chemicals shall complete relevant training in accordance with OSHA requirements.</td>
<td>• CPR (DSP #202, 203, 205, 206, 208, 211, 213, 214, 215, 218, 219, 220, 227, 228, 229)</td>
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</tr>
<tr>
<td>G. Staff shall be certified in a DDSD-approved behavioral intervention system (e.g., Mandt, CPI) before using physical restraint techniques. Staff</td>
<td>• Assisting With Medication Delivery (DSP #200, 206, 207, 208, 209, 210, 213, 215, 216, 220, 221, 228, 229)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Participatory Communication and Choice Making (DSP #207, 209)</td>
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</tbody>
</table>
members providing direct services shall maintain certification in a DDSD-approved behavioral intervention system if an individual they support has a behavioral crisis plan that includes the use of physical restraint techniques.

H. Staff shall complete and maintain certification in a DDSD-approved medication course in accordance with the DDSD Medication Delivery Policy M-001.

I. Staff providing direct services shall complete safety training within the first thirty (30) days of employment and before working alone with an individual receiving service.


<table>
<thead>
<tr>
<th>CHAPTER 5 (CIES) 3. Agency Requirements</th>
</tr>
</thead>
<tbody>
<tr>
<td>G. Training Requirements: 1. All Community Inclusion Providers must provide staff training in accordance with the DDSD policy T-003: Training Requirements for Direct Service Agency Staff Policy.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>CHAPTER 6 (CCS) 3. Agency Requirements</th>
</tr>
</thead>
<tbody>
<tr>
<td>F. Meet all training requirements as follows:</td>
</tr>
<tr>
<td>1. All Customized Community Supports Providers shall provide staff training in accordance with the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff Policy;</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>CHAPTER 7 (CIHS) 3. Agency Requirements</th>
</tr>
</thead>
<tbody>
<tr>
<td>C. Training Requirements: The Provider Agency must report required personnel training status to the DDSD Statewide Training Database as specified in the DDSD Policy T-001: Reporting and Documentation of DDSD Training Requirements Policy. The Provider Agency must ensure that the personnel support staff have completed training as specified in the</td>
</tr>
</tbody>
</table>

- Rights and Advocacy (DSP #207, 209)
- Supporting People with Challenging Behaviors (DSP #207, 209)
- Teaching and Support Strategies (DSP #207, 209)
<table>
<thead>
<tr>
<th>DDSD Policy T-003: Training Requirements for Direct Service Agency Staff Policy</th>
</tr>
</thead>
</table>

**CHAPTER 11 (FL) 3. Agency Requirements B. Living Supports- Family Living Services Provider Agency Staffing Requirements: 3. Training:**
A. All Family Living Provider agencies must ensure staff training in accordance with the Training Requirements for Direct Service Agency Staff policy. DSP’s or subcontractors delivering substitute care under Family Living must at a minimum comply with the section of the training policy that relates to Respite, Substitute Care, and personal support staff [Policy T-003: for Training Requirements for Direct Service Agency Staff; Sec. II-J, Items 1-4]. Pursuant to the Centers for Medicare and Medicaid Services (CMS) requirements, the services that a provider renders may only be claimed for federal match if the provider has completed all necessary training required by the state. All Family Living Provider agencies must report required personnel training status to the DDSD Statewide Training Database as specified in DDSD Policy T-001: Reporting and Documentation for DDSD Training Requirements.

**CHAPTER 12 (SL) 3. Agency Requirements B. Living Supports- Supported Living Services Provider Agency Staffing Requirements: 3. Training:**
A. All Living Supports- Supported Living Provider Agencies must ensure staff training in accordance with the DDSD Policy T-003: for Training Requirements for Direct Service Agency Staff. Pursuant to CMS requirements, the services that a provider renders may only be claimed for federal match if the provider has completed all necessary training required by the state.
state. All Supported Living provider agencies must report required personnel training status to the DDSD Statewide Training Database as specified in DDSD Policy T-001: Reporting and Documentation for DDSD Training Requirements.

CHAPTER 13 (IMLS) R. 2. Service Requirements. Staff Qualifications 2. DSP Qualifications. E. Complete training requirements as specified in the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff - effective March 1, 2007. Report required personnel training status to the DDSD Statewide Training Database as specified in the DDSD Policy T-001: Reporting and Documentation of DDSD Training Requirements Policy;
<table>
<thead>
<tr>
<th>Tag # 1A22</th>
<th>Agency Personnel Competency</th>
<th>Standard Level Deficiency</th>
<th>Provider</th>
</tr>
</thead>
<tbody>
<tr>
<td>Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy - Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007 - II. POLICY STATEMENTS:</td>
<td>Based on interview, the Agency did not ensure training competencies were met for 4 of 9 Direct Support Personnel.</td>
<td>State your Plan of Correction for the deficiencies cited in this tag here: →</td>
<td></td>
</tr>
<tr>
<td>A. Individuals shall receive services from competent and qualified staff.</td>
<td>When DSP were asked if they received training on the Individual’s Individual Service Plan and what the plan covered, the following was reported:</td>
<td></td>
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</tr>
<tr>
<td>B. Staff shall complete individual specific (formerly known as “Addendum B”) training requirements in accordance with the specifications described in the individual service plan (ISP) for each individual serviced.</td>
<td>• DSP #219 stated, “It’s the book.” (Individual #2)</td>
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<td></td>
</tr>
<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013 CHAPTER 5 (CIES) 3. Agency Requirements G. Training Requirements: 1. All Community Inclusion Providers must provide staff training in accordance with the DDSD policy T-003: Training Requirements for Direct Service Agency Staff Policy. 3. Ensure direct service personnel receives Individual Specific Training as outlined in each individual ISP, including aspects of support plans (healthcare and behavioral) or WDSI that pertain to the employment environment.</td>
<td>When DSP were asked if the Individual had a Positive Behavioral Supports Plan and if so, what the plan covered, the following was reported:</td>
<td></td>
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<tr>
<td></td>
<td>• DSP #230 stated, “Not that I’m aware of.” According to the Individual Specific Training Section of the ISP, the Individual requires a Positive Behavioral Supports Plan. (Individual #4)</td>
<td>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →</td>
<td></td>
</tr>
<tr>
<td></td>
<td>When DSP were asked if the individual had a Behavioral Crisis Intervention Plan and if so, what the plan covered, the following was reported:</td>
<td></td>
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<tr>
<td></td>
<td>• DSP #202 stated, “I’m pretty sure there would be a tab that said behavior crisis plan so since I’m not seeing one.” According to the Individual Specific Training Section of the ISP agency file, the individual has Behavioral Crisis Intervention Plan. (Individual #5)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>• DSP #219 stated, “I’d have to look it’s not in this book”. According to the Individual</td>
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<td></td>
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</tr>
</tbody>
</table>

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status to the DDSD Statewide Training Database as specified in the DDSD Policy T-001: Reporting and Documentation of DDSD Training Requirements Policy. The Provider Agency must ensure that the personnel support staff have completed training as specified in the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff Policy. 3. Staff shall complete individual specific training requirements in accordance with the specifications described in the ISP of each individual served; and 4. Staff that assists the individual with medication (e.g., setting up medication, or reminders) must have completed Assisting with Medication Delivery (AWMD) Training.

### CHAPTER 11 (FL) 3. Agency Requirements B. Living Supports - Family Living Services Provider Agency Staffing Requirements: 3. Training:

<table>
<thead>
<tr>
<th>Specific Training Section of the ISP agency file, the individual has Behavioral Crisis Intervention Plan. (Individual #2)</th>
</tr>
</thead>
<tbody>
<tr>
<td>• DSP #230 stated, “Not that I'm aware of.” According to the Individual Specific Training Section of the ISP agency file, the individual has Behavioral Crisis Intervention Plan. (Individual #4)</td>
</tr>
</tbody>
</table>

**When DSP were asked if the Individual had a Speech Therapy Plan and if so, what the plan covered, the following was reported:**

| • DSP #201 stated, “I haven't seen a speech therapist.” According to the Individual Specific Training Section of the ISP, the Individual requires a Speech Therapy Plan. (Individual #6) |

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DSP #230 stated, “Not that I'm aware of.” According to the Individual Specific Training Section of the ISP agency file, the individual has Behavioral Crisis Intervention Plan. (Individual #2)
and Documentation for DDSD Training Requirements.

B. Individual specific training must be arranged and conducted, including training on the Individual Service Plan outcomes, actions steps and strategies and associated support plans (e.g. health care plans, MERPs, PBSP and BCIP etc.), information about the individual's preferences with regard to privacy, communication style, and routines. Individual specific training for therapy related WDSI, Healthcare Plans, MERPs, CARMP, PBSP, and BCIP must occur at least annually and more often if plans change or if monitoring finds incorrect implementation. Family Living providers must notify the relevant support plan author whenever a new DSP is assigned to work with an individual, and therefore needs to receive training, or when an existing DSP requires a refresher. The individual should be present for and involved in individual specific training whenever possible.

CHAPTER 12 (SL) 3. Agency Requirements B. Living Supports - Supported Living Services Provider Agency Staffing Requirements: 3.

Training:
A. All Living Supports - Supported Living Provider Agencies must ensure staff training in accordance with the DDSD Policy T-003: for Training Requirements for Direct Service Agency Staff. Pursuant to CMS requirements, the services that a provider renders may only be claimed for federal match if the provider has completed all necessary training required by the state. All Supported Living provider agencies must report required personnel training status to the DDSD Statewide Training Database as specified in DDSD Policy T-001: Reporting and
Documentation for DDSD Training Requirements.

B Individual specific training must be arranged and conducted, including training on the ISP Outcomes, actions steps and strategies, associated support plans (e.g. health care plans, MERP, PBSP and BCIP, etc.), and information about the individual’s preferences with regard to privacy, communication style, and routines. Individual specific training for therapy related WDSI, Healthcare Plans, MERP, CARMP, PBSP, and BCIP must occur at least annually and more often if plans change or if monitoring finds incorrect implementation. Supported Living providers must notify the relevant support plan author whenever a new DSP is assigned to work with an individual, and therefore needs to receive training, or when an existing DSP requires a refresher. The individual should be present for and involved in individual specific. Training whenever possible.

CHAPTER 13 (IMLS) R. 2. Service Requirements. Staff Qualifications 2. DSP Qualifications. E. Complete training requirements as specified in the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff - effective March 1, 2007. Report required personnel training status to the DDSD Statewide Training Database as specified in the DDSD Policy T-001: Reporting and Documentation of DDSD Training Requirements Policy;
<table>
<thead>
<tr>
<th>Tag # 1A26</th>
<th>Consolidated On-line Registry</th>
<th>Employee Abuse Registry</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>NMAC 7.1.12.8 REGISTRY ESTABLISHED; PROVIDER INQUIRY REQUIRED: Upon the effective date of this rule, the department has established and maintains an accurate and complete electronic registry that contains the name, date of birth, address, social security number, and other appropriate identifying information of all persons who, while employed by a provider, have been determined by the department, as a result of an investigation of a complaint, to have engaged in a substantiated registry-referred incident of abuse, neglect or exploitation of a person receiving care or services from a provider. Additions and updates to the registry shall be posted no later than two (2) business days following receipt. Only department staff designated by the custodian may access, maintain and update the data in the registry.</td>
<td>Based on record review, the Agency did not maintain documentation in the employee’s personnel records that evidenced inquiry into the Employee Abuse Registry prior to employment for 1 of 31 Agency Personnel. The following Agency personnel records contained no evidence of the Employee Abuse Registry check being completed: Direct Support Personnel (DSP):</td>
<td>Provider: State your Plan of Correction for the deficiencies cited in this tag here: →</td>
<td></td>
</tr>
<tr>
<td>A. Provider requirement to inquire of registry. A provider, prior to employing or contracting with an employee, shall inquire of the registry whether the individual under consideration for employment or contracting is listed on the registry.</td>
<td></td>
<td>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →</td>
<td></td>
</tr>
<tr>
<td>B. Prohibited employment. A provider may not employ or contract with an individual to be an employee if the individual is listed on the registry as having a substantiated registry-referred incident of abuse, neglect or exploitation of a person receiving care or services from a provider.</td>
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<tr>
<td>D. Documentation of inquiry to registry. The provider shall maintain documentation in the employee’s personnel or employment records that evidences the fact that the provider made an</td>
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</tbody>
</table>
inquiry to the registry concerning that employee prior to employment. Such documentation must include evidence, based on the response to such inquiry received from the custodian by the provider, that the employee was not listed on the registry as having a substantiated registry-referred incident of abuse, neglect or exploitation.

E. **Documentation for other staff.** With respect to all employed or contracted individuals providing direct care who are licensed health care professionals or certified nurse aides, the provider shall maintain documentation reflecting the individual’s current licensure as a health care professional or current certification as a nurse aide.

F. **Consequences of noncompliance.** The department or other governmental agency having regulatory enforcement authority over a provider may sanction a provider in accordance with applicable law if the provider fails to make an appropriate and timely inquiry of the registry, or fails to maintain evidence of such inquiry, in connection with the hiring or contracting of an employee; or for employing or contracting any person to work as an employee who is listed on the registry. Such sanctions may include a directed plan of correction, civil monetary penalty not to exceed five thousand dollars ($5000) per instance, or termination or non-renewal of any contract with the department or other governmental agency.
<table>
<thead>
<tr>
<th>Tag # 1A28.1</th>
<th>Incident Mgt. System - Personnel Training</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Standard Level Deficiency</strong></td>
<td></td>
</tr>
<tr>
<td>Based on record review and interview, the Agency did not ensure Incident Management Training for 8 of 31 Agency Personnel.</td>
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</table>

**Direct Support Personnel (DSP):**
- Incident Management Training (Abuse, Neglect and Exploitation) (DSP# 204, 216, 218, 221, 222, 228)

**Service Coordination Personnel (SC):**
- Incident Management Training (Abuse, Neglect and Exploitation) (SC #231, 232)

**Provider:**
State your Plan of Correction for the deficiencies cited in this tag here: →

Provider:
Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →

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<table>
<thead>
<tr>
<th>C. Incident management system training curriculum requirements:</th>
</tr>
</thead>
<tbody>
<tr>
<td>(1) The community-based service provider shall conduct training or designate a knowledgeable representative to conduct training, in accordance with the written training curriculum provided electronically by the division that includes but is not limited to:</td>
</tr>
<tr>
<td>(a) an overview of the potential risk of abuse, neglect, or exploitation;</td>
</tr>
<tr>
<td>(b) informational procedures for properly filing the division's abuse, neglect, and exploitation or report of death form;</td>
</tr>
<tr>
<td>(c) specific instructions of the employees' legal responsibility to report an incident of abuse, neglect and exploitation, suspicious injury, and all deaths;</td>
</tr>
<tr>
<td>(d) specific instructions on how to respond to abuse, neglect, or exploitation;</td>
</tr>
<tr>
<td>(e) emergency action procedures to be followed in the event of an alleged incident or knowledge of abuse, neglect, exploitation, or suspicious injury.</td>
</tr>
<tr>
<td>(2) All current employees and volunteers shall receive training within 90 days of the effective date of this rule.</td>
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<tr>
<td>(3) All new employees and volunteers shall receive training prior to providing services to consumers.</td>
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</tbody>
</table>

D. Training documentation: All community-based service providers shall prepare training documentation for each employee and volunteer to include a signed statement indicating the date, time, and place they received their incident management reporting instruction. The community-based service provider shall maintain documentation of an employee or volunteer's training for a period of at least three years, or six months after termination of an employee's employment or the volunteer's work.
curricula shall be kept on the provider premises and made available upon request by the department. Training documentation shall be made available immediately upon a division representative's request. Failure to provide employee and volunteer training documentation shall subject the community-based service provider to the penalties provided for in this rule.

Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007

II. POLICY STATEMENTS:
A. Individuals shall receive services from competent and qualified staff.
C. Staff shall complete training on DOH-approved incident reporting procedures in accordance with 7 NMAC 1.13.
<table>
<thead>
<tr>
<th>Tag # 1A36</th>
<th>Service Coordination Requirements</th>
<th>Standard Level Deficiency</th>
<th>Provider</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tag # 1A36</td>
<td>Service Coordination Requirements</td>
<td>Standard Level Deficiency</td>
<td>Provider</td>
</tr>
<tr>
<td>Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy</td>
<td>Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007 - II. POLICY STATEMENTS:</td>
<td>Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007 - II. POLICY STATEMENTS:</td>
<td>State your Plan of Correction for the deficiencies cited in this tag here: →</td>
</tr>
<tr>
<td>K. In addition to the applicable requirements described in policy statements B – I (above), direct support staff, direct support supervisors, and internal service coordinators shall complete DDSD-approved core curriculum training. Attachments A and B to this policy identify the specific competency requirements for the following levels of core curriculum training:</td>
<td>Based on record review, the Agency did not ensure that Orientation and Training requirements were met for 1 of 2 Service Coordinators.</td>
<td>Based on record review, the Agency did not ensure that Orientation and Training requirements were met for 1 of 2 Service Coordinators.</td>
<td></td>
</tr>
<tr>
<td>1. Introductory Level – must be completed within thirty (30) days of assignment to his/her position with the agency.</td>
<td>Review of Service Coordinators training records found no evidence of the following required DOH/DDSD trainings being completed:</td>
<td>Review of Service Coordinators training records found no evidence of the following required DOH/DDSD trainings being completed:</td>
<td></td>
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<tr>
<td>2. Orientation – must be completed within ninety (90) days of assignment to his/her position with the agency.</td>
<td>• Pre-Service Part One (SC #232)</td>
<td>• Pre-Service Part One (SC #232)</td>
<td></td>
</tr>
<tr>
<td>3. Level I – must be completed within one (1) year of assignment to his/her position with the agency.</td>
<td>• Pre-Service Part Two (SC #232)</td>
<td>• Pre-Service Part Two (SC #232)</td>
<td></td>
</tr>
<tr>
<td>NMAC 7.26.5.7 “service coordinator”: the community provider staff member, sometimes called the program manager or the internal case manager, who supervises, implements and monitors the service plan within the community service provider agency</td>
<td>• Promoting Effective Teamwork (SC #232)</td>
<td>• Promoting Effective Teamwork (SC #232)</td>
<td></td>
</tr>
<tr>
<td>NMAC 7.26.5.11 (b) service coordinator: the service coordinators of the community provider agencies shall assure that appropriate staff develop strategies specific to their responsibilities in the ISP; the service coordinators shall assure the action plans and strategies are implemented consistent with the</td>
<td>• ISP Critique (SC #232)</td>
<td>• ISP Critique (SC #232)</td>
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<td></td>
<td>• Sexuality for People with Developmental Disabilities (SC #232)</td>
<td>• Sexuality for People with Developmental Disabilities (SC #232)</td>
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<td></td>
<td>• Level 1 Health (SC #232)</td>
<td>• Level 1 Health (SC #232)</td>
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</tbody>
</table>

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provisions of the ISP, and shall report to the case manager on ISP implementation and the individual's progress on action plans within their agencies; for persons funded solely by state general funds, the service coordinator shall assume all the duties of the independent case manager described within these regulations; if there are two or more “key” community service provider agencies with two or more service coordinator staff, the IDT shall designate which service coordinator shall assume the duties of the case manager; the criteria to guide the IDTs selection are set forth as follows:

(i) the designated service coordinator shall have the skills necessary to carry out the duties and responsibilities of the case manager as defined in these regulations;
(ii) the designated service coordinator shall have the time and interest to fulfill the functions of the case manager as defined in these regulations;
(iii) the designated service coordinator shall be familiar with and understand community service delivery and supports;
(iv) the designated service coordinator shall know the individual or be willing to become familiar and develop a relationship with the individual being served;
<table>
<thead>
<tr>
<th>Tag # 1A37</th>
<th>Individual Specific Training</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy - Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007 - II. POLICY STATEMENTS:</td>
<td>Based on record review, the Agency did not ensure that Individual Specific Training requirements were met for 5 of 31 Agency Personnel.</td>
<td></td>
</tr>
<tr>
<td>A. Individuals shall receive services from competent and qualified staff.</td>
<td>Review of personnel records found no evidence of the following:</td>
<td></td>
</tr>
<tr>
<td>B. Staff shall complete individual specific (formerly known as “Addendum B”) training requirements in accordance with the specifications described in the individual service plan (ISP) for each individual serviced.</td>
<td><strong>Direct Support Personnel (DSP):</strong></td>
<td></td>
</tr>
<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013</td>
<td>• Individual Specific Training (DSP #209, 223, 224, 225, 226)</td>
<td></td>
</tr>
<tr>
<td><strong>CHAPTER 5 (CIES) 3. Agency Requirements</strong></td>
<td><strong>Provider:</strong></td>
<td></td>
</tr>
<tr>
<td>G. Training Requirements: 1. All Community Inclusion Providers must provide staff training in accordance with the DDSD policy T-003: Training Requirements for Direct Service Agency Staff Policy. 3. Ensure direct service personnel receives Individual Specific Training as outlined in each individual ISP, including aspects of support plans (healthcare and behavioral) or WDSI that pertain to the employment environment.</td>
<td>State your Plan of Correction for the deficiencies cited in this tag here: →</td>
<td></td>
</tr>
<tr>
<td><strong>CHAPTER 6 (CCS) 3. Agency Requirements</strong></td>
<td><strong>Provider:</strong></td>
<td></td>
</tr>
<tr>
<td>F. Meet all training requirements as follows: 1. All Customized Community Supports Providers shall provide staff training in accordance with the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff Policy;</td>
<td>Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →</td>
<td></td>
</tr>
<tr>
<td><strong>CHAPTER 7 (CIHS) 3. Agency Requirements</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>C. Training Requirements: The Provider Agency must report required personnel training</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Provider: State your Plan of Correction for the deficiencies cited in this tag here: →

Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →
status to the DDSD Statewide Training Database as specified in the DDSD Policy T-001: Reporting and Documentation of DDSD Training Requirements Policy. The Provider Agency must ensure that the personnel support staff have completed training as specified in the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff Policy. 3. Staff shall complete individual specific training requirements in accordance with the specifications described in the ISP of each individual served; and 4. Staff that assists the individual with medication (e.g., setting up medication, or reminders) must have completed Assisting with Medication Delivery (AWMD) Training.

CHAPTER 11 (FL) 3. Agency Requirements B. Living Supports- Family Living Services Provider Agency Staffing Requirements: 3. Training:
A. All Family Living Provider agencies must ensure staff training in accordance with the Training Requirements for Direct Service Agency Staff policy. DSP’s or subcontractors delivering substitute care under Family Living must at a minimum comply with the section of the training policy that relates to Respite, Substitute Care, and personal support staff [Policy T-003: for Training Requirements for Direct Service Agency Staff; Sec. II-J, Items 1-4]. Pursuant to the Centers for Medicare and Medicaid Services (CMS) requirements, the services that a provider renders may only be claimed for federal match if the provider has completed all necessary training required by the state. All Family Living Provider agencies must report required personnel training status to the DDSD Statewide Training Database as specified in DDSD Policy T-001: Reporting
and Documentation for DDSD Training Requirements.

B. Individual specific training must be arranged and conducted, including training on the Individual Service Plan outcomes, actions steps and strategies and associated support plans (e.g. health care plans, MERP, PBSP and BCIP etc.), information about the individual's preferences with regard to privacy, communication style, and routines. Individual specific training for therapy related WDSI, Healthcare Plans, MERPs, CARMP, PBSP, and BCIP must occur at least annually and more often if plans change or if monitoring finds incorrect implementation. Family Living providers must notify the relevant support plan author whenever a new DSP is assigned to work with an individual, and therefore needs to receive training, or when an existing DSP requires a refresher. The individual should be present for and involved in individual specific training whenever possible.

CHAPTER 12 (SL) 3. Agency Requirements B. Living Supports- Supported Living Services Provider Agency Staffing Requirements: 3. Training:

A. All Living Supports- Supported Living Provider Agencies must ensure staff training in accordance with the DDSD Policy T-003: for Training Requirements for Direct Service Agency Staff. Pursuant to CMS requirements, the services that a provider renders may only be claimed for federal match if the provider has completed all necessary training required by the state. All Supported Living provider agencies must report required personnel training status to the DDSD Statewide Training Database as specified in DDSD Policy T-001: Reporting and
Documentation for DDSD Training Requirements.

B. Individual specific training must be arranged and conducted, including training on the ISP Outcomes, actions steps and strategies, associated support plans (e.g. health care plans, MERP, PBSP and BCIP, etc.), and information about the individual’s preferences with regard to privacy, communication style, and routines. Individual specific training for therapy related WDSI, Healthcare Plans, MERP, CARMP, PBSP, and BCIP must occur at least annually and more often if plans change or if monitoring finds incorrect implementation. Supported Living providers must notify the relevant support plan author whenever a new DSP is assigned to work with an individual, and therefore needs to receive training, or when an existing DSP requires a refresher. The individual should be present for and involved in individual specific. Training whenever possible.

CHAPTER 13 (IMLS) R. 2. Service Requirements. Staff Qualifications 2. DSP Qualifications. E. Complete training requirements as specified in the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff - effective March 1, 2007. Report required personnel training status to the DDSD Statewide Training Database as specified in the DDSD Policy T-001: Reporting and Documentation of DDSD Training Requirements Policy;
<table>
<thead>
<tr>
<th>Tag # 1A43 General Events Reporting</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Department of Health (DOH) Developmental Disabilities Supports Division (DDSD)</strong> Policy: General Events Reporting Effective 1/1/2012</td>
<td></td>
</tr>
</tbody>
</table>
| **1. Purpose**  
To report, track and analyze significant events experiences by adult participants of the DD Waiver program, which do not meet criteria for abuse, neglect or exploitation, or other “reportable incident” as defined by the Incident Management Bureau of the Division of Health Improvement, Department of Health, but which pose a risk to individuals served. Analysis of reported significant events is intended to identify emerging patterns so that preventative actions can be identified at the individual, provider agency, regional and statewide levels. |
| **II. Policy Statements** |
| **A.** Designated employees of each agency will enter specified information into the General Events Reporting section of the secure website operated under contract by Therap Services within 2 business days of the occurrence or knowledge by the reporting agency of any of the following defined events in which DDSD requires reporting: Chocking, Missing Person, Suicide Attempt or Threat, Restraint related to Behavior, Serious Injury including Skin Breakdown, Fall (with or without injury), Out of Home Placement and Infections...Providers shall utilize the “Significant Events Reporting System Guide” to assure that events are reported correctly for DDSD tracking purposes. At providers’ discretion additional events may be tracked. Based on record review, observation and/or interview the Agency did not follow the General Events Reporting requirements as indicated by the policy. |
| **Review of the agency 2015 Quality improvement plan indicated the following:** |
| - "Therap will be used as the information site to add reports. The Director Nursing will enter all medical reports. The Service Coordinator, Quality Enhancement Coordinator or Executive Director will enter all non-medical reports.” |
| **Review of GER on Therap revealed the following:** |
| - Internal reports that are medical and non-medical are not being entered into Therap within 2 business days as required per Standard. |
| - There was no reporting on certain medication errors, falls, injury, self-harm etc. as required per Standard. |
| - There were 8 medical incident reports entered into General Events Reporting System in Therap for the reporting period July 2014 – July 2015. However, the reports were not approved in the system by the agency. |
| - Non-Medical Reports that are required to be reported through the General Events Reporting System were not being entered into therap as required per Standard. |

Provider:  
State your Plan of Correction for the deficiencies cited in this tag here: →

Provider:  
Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →
within the Therap General Events Reporting which are not required by DDSD such as medication errors.

B. General Events Reporting does not replace agency obligations to report abuse, neglect, exploitation and other reportable incidents in compliance with policies and procedures issued by the Department's Incident Management Bureau of the Division of Health Improvement.

When #233 was asked by Surveyors to explain and show how they utilize the General Events Reporting System the follow was reported and observed:

- The Opportunity Center maintains their own access database, this reporting system does not allow the user to review, track and tread internal reports. In addition, it prints the entire history of internal reports. This system does not allow the user to run a report based on time period.
**Standard of Care**  | **Deficiencies**  | **Agency Plan of Correction, On-going QA/QI and Responsible Party**  | **Date Due**
---|---|---|---

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

<table>
<thead>
<tr>
<th>Tag #1A08.2 Healthcare Requirements</th>
<th>Standard Level Deficiency</th>
<th>Provider: State your Plan of Correction for the deficiencies cited in this tag here: →</th>
</tr>
</thead>
</table>
| NMAC 8.302.1.17 RECORD KEEPING AND DOCUMENTATION REQUIREMENTS: A provider must maintain all the records necessary to fully disclose the nature, quality, amount and medical necessity of services furnished to an eligible recipient who is currently receiving or who has received services in the past. | Based on record review, the Agency did not provide documentation of annual physical examinations and/or other examinations as specified by a licensed physician for 7 of 7 individuals receiving Community Inclusion, Living Services and Other Services. Review of the administrative individual case files revealed the following items were not found, incomplete, and/or not current: **Community Inclusion Services / Other Services Healthcare Requirements (Individuals Receiving Inclusion / Other Services Only):**  
- **Follow up to Annual Physical**  
  - Individual #1 - As indicated by collateral documentation reviewed, the exam was completed on 10/23/2014. Physician’s progress notes stated “Would like to see every 6 months”. No evidence of follow up were found.  
  
**Community Living Services / Community Inclusion Services (Individuals Receiving Multiple Services):**  
- **Annual Physical (#4)**  
- **Dental Exam** | Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: → |

**DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION (DDSD): Director’s Release: Consumer Record Requirements eff. 11/1/2012 III. Requirement Amendments(s) or Clarifications:**  
A. All case management, living supports, customized in-home supports, community integrated employment and customized community supports providers must maintain records for individuals served through DD Waiver in accordance with the Individual Case File Matrix incorporated in this director’s release.  
H. Readily accessible electronic records are accessible, including those stored through the Therap web-based system.
| Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013 | **Chapter 5 (CIES) 3. Agency Requirements**  
**H. Consumer Records Policy:** All Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Consumer Records Policy. |
|---|---|
| **Chapter 6 (CCS) 3. Agency Requirements:**  
**G. Consumer Records Policy:** All Provider Agencies shall maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix policy. |  
| **Chapter 7 (CIHS) 3. Agency Requirements:**  
**E. Consumer Records Policy:** All Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix policy. |
| **Chapter 11 (FL) 3. Agency Requirements:**  
**D. Consumer Records Policy:** All Family Living Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix policy. |  
| **Chapter 12 (SL) 3. Agency Requirements:**  
**D. Consumer Records Policy:** All Living Supports - Supported Living Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix policy. |
| **Vision Exam**  
| |  
| Individual #2 - As indicated by the DDSD file matrix, Vision Exams are to be conducted every other year. No evidence of exam was found. |
| Individual #3 - As indicated by collateral documentation reviewed, exam was completed on 10/7/2013. Follow-up was to be completed in 1 year. No evidence of follow-up found. |
| Individual #4 - As indicated by the DDSD file matrix Dental Exams are to be conducted annually. No evidence of exam was found. |
| Individual #5 - As indicated by the DDSD file matrix Dental Exams are to be conducted annually. No evidence of exam was found. |
| Individual #6 - As indicated by collateral documentation reviewed, exam was conducted annually. No evidence of exam results were found. |  
| Individual #7 - As indicated by collateral documentation reviewed, the exam was completed on 7/21/2015. No evidence of exam results were found. |
comply with the DDSD Individual Case File Matrix policy.

Chapter 13 (IMLS) 2. Service Requirements:
C. Documents to be maintained in the agency administrative office, include: (This is not an all-inclusive list refer to standard as it includes other items)…


CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS: D. Provider Agency Case File for the Individual: All Provider Agencies shall maintain at the administrative office a confidential case file for each individual. Case records belong to the individual receiving services and copies shall be provided to the receiving agency whenever an individual changes providers. The record must also be made available for review when requested by DOH, HSD or federal government representatives for oversight purposes. The individual’s case file shall include the following requirements:
(5) A medical history, which shall include at least demographic data, current and past medical diagnoses including the cause (if known) of the developmental disability, psychiatric diagnoses, allergies (food, environmental, medications), immunizations, and most recent physical exam;

CHAPTER 6. VI. GENERAL REQUIREMENTS FOR COMMUNITY LIVING
G. Health Care Requirements for Community Living Services.
(1) The Community Living Service providers shall ensure completion of a HAT for each individual receiving this service. The HAT shall completed on 5/29/2015. Follow-up was to be completed in 1 year. No evidence of follow-up found.

- Nutritional Evaluation
  ▪ Individual #6 - According to Nutritional evaluation the individual is required to have a 6 month follow-up. No evidence of follow-up found.
be completed 2 weeks prior to the annual ISP meeting and submitted to the Case Manager and all other IDT Members. A revised HAT is required to also be submitted whenever the individual’s health status changes significantly. For individuals who are newly allocated to the DD Waiver program, the HAT may be completed within 2 weeks following the initial ISP meeting and submitted with any strategies and support plans indicated in the ISP, or within 72 hours following admission into direct services, whichever comes first.

(2) Each individual will have a Health Care Coordinator, designated by the IDT. When the individual’s HAT score is 4, 5 or 6 the Health Care Coordinator shall be an IDT member, other than the individual. The Health Care Coordinator shall oversee and monitor health care services for the individual in accordance with these standards. In circumstances where no IDT member voluntarily accepts designation as the health care coordinator, the community living provider shall assign a staff member to this role.

(3) For each individual receiving Community Living Services, the provider agency shall ensure and document the following:

(a) Provision of health care oversight consistent with these Standards as detailed in Chapter One section III E: Healthcare Documentation by Nurses For Community Living Services, Community Inclusion Services and Private Duty Nursing Services.

b) That each individual with a score of 4, 5, or 6 on the HAT, has a Health Care Plan developed by a licensed nurse.

c) That an individual with chronic condition(s) with the potential to exacerbate into a life threatening
condition, has Crisis Prevention/Intervention Plan(s) developed by a licensed nurse or other appropriate professional for each such condition.

(4) That an average of 3 hours of documented nutritional counseling is available annually, if recommended by the IDT.

(5) That the physical property and grounds are free of hazards to the individual's health and safety.

(6) In addition, for each individual receiving Supported Living or Family Living Services, the provider shall verify and document the following:
   (a) The individual has a primary licensed physician;
   (b) The individual receives an annual physical examination and other examinations as specified by a licensed physician;
   (c) The individual receives annual dental check-ups and other check-ups as specified by a licensed dentist;
   (d) The individual receives eye examinations as specified by a licensed optometrist or ophthalmologist; and
   (e) Agency activities that occur as follow-up to medical appointments (e.g. treatment, visits to specialists, changes in medication or daily routine).
<table>
<thead>
<tr>
<th>Tag #</th>
<th>1A03</th>
<th>CQI System</th>
<th>Standard Level Deficiency</th>
<th>Provider</th>
</tr>
</thead>
<tbody>
<tr>
<td>STATE OF NEW MEXICO DEPARTMENT OF HEALTH DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION PROVIDER AGREEMENT: ARTICLE 17. PROGRAM EVALUATIONS</td>
<td>Based on record review and interview, the Agency had not fully implemented their Continuous Quality Management System as required by standard.</td>
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<td>d. PROVIDER shall have a Quality Management and Improvement Plan in accordance with the current MF Waiver Standards and/or the DD Waiver Standards specified by the DEPARTMENT. The Quality Management and Improvement Plan for DD Waiver Providers must describe how the PROVIDER will determine that each waiver assurance and requirement is met. The applicable assurances and requirements are: (1) level of care determination; (2) service plan; (3) qualified providers; (4) health and welfare; (5) administrative authority; and, (6) financial accountability. For each waiver assurance, this description must include:</td>
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<td>i. Activities or processes related to discovery, i.e., monitoring and recording the findings. Descriptions of monitoring/oversight activities that occur at the individual and provider level of service delivery. These monitoring activities provide a foundation for Quality Management by generating information that can be aggregated and analyzed to measure the overall system performance;</td>
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<td>ii. The entities or individuals responsible for conducting the discovery/monitoring processes;</td>
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<td>iii. The types of information used to measure performance; and,</td>
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<td>iv. The frequency with which performance is measured.</td>
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<tr>
<td>CHAPTER 5 (CIES) 3. Agency Requirements: J. Quality Assurance/Quality Improvement (QA/QI) Program: Agencies must develop and maintain an</td>
<td>Provider:</td>
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<td>State your Plan of Correction for the deficiencies cited in this tag here: →</td>
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</table>
active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QA/QI activities.

1. **Development of a QA/QI plan:** The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working.

2. **Implementing a QA/QI Committee:** The QA/QI committee must convene on at least a quarterly basis and as needed to review service reports, to identify any deficiencies, trends, patterns or concerns as well as opportunities for quality improvement. The QA/QI meeting must be documented. The QA/QI review should address at least the following:
   a. **Implementation of ISPs:** extent to which services are delivered in accordance with ISPs and associated support plans with WDSI including the type, scope, amount, duration and frequency specified in the ISP as well as effectiveness of such implementation as indicated by achievement of outcomes;

3. The Provider Agency must complete a QA/QI report annually by February 15th of each calendar year or as otherwise requested by DOH. The report must be kept on file at the agency, made
available for review by DOH and upon request from
DDSD; the report must be submitted to the relevant
DDSD Regional Offices. The report will
summarize:

a. Analysis of General Events Reports data in
Therap;
b. Compliance with Caregivers Criminal History
Screening requirements;
c. Compliance with Employee Abuse Registry
requirements;
d. Compliance with DDSD training requirements;
e. Patterns of reportable incidents;
f. Results of improvement actions taken in
previous quarters;
g. Sufficiency of staff coverage;
h. Effectiveness and timeliness of implementation
of ISPs, and associated support including trends
in achievement of individual desired outcomes;
i. Results of General Events Reporting data
analysis;
j. Action taken regarding individual grievances;
k. Presence and completeness of required
documentation;
l. A description of how data collected as part of the
agency’s QA/QI Plan was used; what quality
improvement initiatives were undertaken and
what were the results of those efforts, including
discovery and remediation of any service
delivery deficiencies discovered through the
QA/QI process; and
m. Significant program changes.

CHAPTER 6 (CCS) 3. Agency Requirements: I.
Quality Assurance/Quality Improvement (QA/QI)
Program: Agencies must develop and maintain an
active QA/QI program in order to assure the
provision of quality services. This includes the
development of a QA/QI plan, data gathering and
analysis, and routine meetings to analyze the
results of QI activities.
1. Development of a QI plan: The quality
management plan is used by an agency to
continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working.

2. Implementing a QI Committee: The QA/QI committee shall convene at least quarterly and as needed to review service reports, to identify any deficiencies, trends, patterns or concerns as well as opportunities for quality improvement. The QA/QI meeting shall be documented. The QA/QI review should address at least the following:
   a. The extent to which services are delivered in accordance with ISPs, associated support plans and WDSI including the type, scope, amount, duration and frequency specified in the ISP as well as effectiveness of such implementation as indicated by achievement of outcomes;
   b. Analysis of General Events Reports data;
   c. Compliance with Caregivers Criminal History Screening requirements;
   d. Compliance with Employee Abuse Registry requirements;
   e. Compliance with DDSD training requirements;
   f. Patterns of reportable incidents; and
   g. Results of improvement actions taken in previous quarters.

3. The Provider Agencies must complete a QA/QI report annually by February 15th of each year, or as otherwise requested by DOH. The report must be kept on file at the agency, made available for
review by DOH and upon request from DDSD the report must be submitted to the relevant DDSD Regional Offices. The report will summarize:

a. Sufficiency of staff coverage;
b. Effectiveness and timeliness of implementation of ISPs, associated support plans, and WDSI, including trends in achievement of individual desired outcomes;
c. Results of General Events Reporting data analysis;
d. Action taken regarding individual grievances;
e. Presence and completeness of required documentation;
f. A description of how data collected as part of the agency’s QI plan was used; what quality improvement initiatives were undertaken and what were the results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QI process; and

g. Significant program changes.

CHAPTER 7 (CIHS) 3. Agency Requirements: G. Quality Assurance/Quality Improvement (QA/QI) Program: Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QA/QI activities.

1. Development of a QA/QI plan: The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should
describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working.

2. **Implementing a QA/QI Committee:** The QA/QI committee shall convene on at least a quarterly basis and as needed to review monthly service reports, to identify any deficiencies, trends, patterns or concerns as well as opportunities for quality improvement. The QA/QI meeting must be documented. The QA/QI review should address at least the following:

   a. **Implementation of ISPs:** The extent to which services are delivered in accordance with ISPs and associated support plans and/or WDSI including the type, scope, amount, duration and frequency specified in the ISP as well as effectiveness of such implementation as indicated by achievement of outcomes;

   b. Analysis of General Events Reports data;

   c. Compliance with Caregivers Criminal History Screening requirements;

   d. Compliance with Employee Abuse Registry requirements;

   e. Compliance with DDSD training requirements;

   f. Patterns of reportable incidents; and

   g. Results of improvement actions taken in previous quarters.

3. The Provider Agency must complete a QA/QI report annually by February 15th of each calendar year, or as otherwise request by DOH. The report must be kept on file at the agency, made available for review by DOH and, upon request from DDSD
the report must be submitted to the relevant DDSD Regional Offices. The report will summarize:

a. Sufficiency of staff coverage;

b. Effectiveness and timeliness of implementation of ISPs and associated support plans and/or WDSI, including trends in achievement of individual desired outcomes;

c. Results of General Events Reporting data analysis;

d. Action taken regarding individual grievances;

e. Presence and completeness of required documentation;

f. A description of how data collected as part of the agency’s QA/QI plan was used; what quality improvement initiatives were undertaken and what were the results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QI process; and

g. Significant program changes.

CHAPTER 11 (FL) 3. Agency Requirements: H. Quality Improvement/Quality Assurance (QA/QI) Program: Family Living Provider Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QA/QI activities.

1. Development of a QA/QI plan: The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in
each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working.

2. Implementing a QA/QI Committee: The QA/QI committee must convene on at least a quarterly basis and as needed to review monthly service reports, to identify any deficiencies, trends, patterns or concerns as well as opportunities for quality improvement. The QA/QI meeting must be documented. The QA/QI review should address at least the following:

a. The extent to which services are delivered in accordance with the ISP including the type, scope, amount, duration and frequency specified in the ISP as well as effectiveness of such implementation as indicated by achievement of outcomes;

b. Analysis of General Events Reports data;

c. Compliance with Caregivers Criminal History Screening requirements;

d. Compliance with Employee Abuse Registry requirements;

e. Compliance with DDSD training requirements;

f. Patterns in reportable incidents; and

g. Results of improvement actions taken in previous quarters.

3. The Provider Agency must complete a QA/QI report annually by February 15th of each year, or as otherwise requested by DOH. The report must be kept on file at the agency, made available for review by DOH and upon request from DDSD; the report must be submitted to the relevant DDSD Regional Offices. The report will summarize:

a. Sufficiency of staff coverage;
b. Effectiveness and timeliness of implementation of ISPs, including trends in achievement of individual desired outcomes;
c. Results of General Events Reporting data analysis, Trends in category II significant events;
d. Patterns in medication errors;
e. Action taken regarding individual grievances;
f. Presence and completeness of required documentation;
g. A description of how data collected as part of the agency's QI plan was used;
h. What quality improvement initiatives were undertaken and what were the results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QI process; and
i. Significant program changes.

CHAPTER 12 (SL) 3. Agency Requirements: B. Quality Assurance/Quality Improvement (QA/QI) Program: Supported Living Provider Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QA/QI activities.

1. Development of a QA/QI plan: The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to
evaluate whether implementation of improvements are working.

2. **Implementing a QA/QI Committee**: The QA/QI committee must convene on at least a quarterly basis and as needed to review monthly service reports, to identify any deficiencies, trends, patterns, or concerns as well as opportunities for quality improvement. The QA/QI meeting must be documented. The QA/QI review should address at least the following:
   
a. Implementation of the ISP and the extent to which services are delivered in accordance with the ISP including the type, scope, amount, duration, and frequency specified in the ISP as well as effectiveness of such implementation as indicated by achievement of outcomes;
   b. Analysis of General Events Reports data;
   c. Compliance with Caregivers Criminal History Screening requirements;
   d. Compliance with Employee Abuse Registry requirements;
   e. Compliance with DDSD training requirements;
   f. Patterns in reportable incidents; and
   g. Results of improvement actions taken in previous quarters.

2. The Provider Agency must complete a QA/QI report annually by February 15th of each calendar year, or as otherwise requested by DOH. The report must be kept on file at the agency, made available for review by DOH, and upon request from DDSD the report must be submitted to the relevant DDSD Regional Offices. The report will summarize:
   
a. Sufficiency of staff coverage;
   b. Effectiveness and timeliness of implementation of ISPs, including trends in achievement of individual desired outcomes;
   c. Results of General Events Reporting data analysis, Trends in Category II significant events;
d. Patterns in medication errors;
e. Action taken regarding individual grievances;
f. Presence and completeness of required documentation;
g. A description of how data collected as part of the agency’s QA/QI plan was used, what quality improvement initiatives were undertaken, and the results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QI process; and
h. Significant program changes.

CHAPTER 13 (IMLS) 3. Service Requirements:
F. Quality Assurance/Quality Improvement (QA/QI) Program: Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QI activities.

1. Development of a QI plan: The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working.

2. Implementing a QA/QI Committee: The QA/QI committee shall convene on at least a quarterly basis and as needed to review service reports, to identify any deficiencies, trends, patterns or
concerns, as well as opportunities for quality improvement. For Intensive Medical Living providers, at least one nurse shall be a member of this committee. The QA meeting shall be documented. The QA review should address at least the following:

a. Implementation of the ISPs, including the extent to which services are delivered in accordance with the ISPs and associated support plans and/or WDSI including the type, scope, amount, duration, and frequency specified in the ISPs as well as effectiveness of such implementation as indicated by achievement of outcomes;

b. Trends in General Events as defined by DDSD;

c. Compliance with Caregivers Criminal History Screening Requirements;

d. Compliance with DDSD training requirements;

e. Trends in reportable incidents; and

f. Results of improvement actions taken in previous quarters.

3. The Provider Agency must complete a QA/QI report annually by February 15th of each calendar year, or as otherwise requested by DOH. The report must be kept on file at the agency, made available for review by DOH and upon request from DDSD; the report must be submitted to the relevant DDSD Regional Offices. The report will summarize:

a. Sufficiency of staff coverage;

b. Effectiveness and timeliness of implementation of ISPs and associated Support plans and/or WDSI including trends in achievement of individual desired outcomes;

c. Trends in reportable incidents;

d. Trends in medication errors;

e. Action taken regarding individual grievances;

f. Presence and completeness of required documentation;

g. How data collected as part of the agency's QA/QI was used, what quality improvement initiatives were undertaken, and what were the
results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QI process; and h. Significant program changes.

CHAPTER 14 (ANS) 3. Service Requirements:
N. Quality Assurance/Quality Improvement (QA/QI) Program: Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QI activities.

1. Development of a QI plan: The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working.

2. Implementing a QA/QI Committee: The QA/QI committee shall convene on at least on a quarterly basis and as needed to review service reports, to identify any deficiencies, trends, patterns or concerns, as well as opportunities for quality improvement. For Intensive Medical Living providers, at least one nurse shall be a member of this committee. The QA meeting shall be documented. The QA review should address at least the following:
   a. Trends in General Events as defined by DDSD;
b. Compliance with Caregivers Criminal History Screening Requirements;
c. Compliance with DDSD training requirements;
d. Trends in reportable incidents; and
e. Results of improvement actions taken in previous quarters.

3. The Provider Agency must complete a QA/QI report annually by February 15th of each calendar year, or as otherwise requested by DOH. The report must be kept on file at the agency, made available for review by DOH and upon request from DDSD; the report must be submitted to the relevant DDSD Regional Offices. The report will summarizes:
   a. Sufficiency of staff coverage;
b. Trends in reportable incidents;
c. Trends in medication errors;
d. Action taken regarding individual grievances;
e. Presence and completeness of required documentation;
f. How data collected as part of the agency’s QA/QI was used, what quality improvement initiatives were undertaken, and what were the results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QI process; and

g. Significant program changes

NMAC 7.1.14.8 INCIDENT MANAGEMENT SYSTEM REPORTING REQUIREMENTS FOR COMMUNITY-BASED SERVICE PROVIDERS:
F. Quality assurance/quality improvement program for community-based service providers:
The community-based service provider shall establish and implement a quality improvement program for reviewing alleged complaints and incidents of abuse, neglect, or exploitation against them as a provider after the division’s investigation is complete. The incident management program shall include written documentation of corrective actions
taken. The community-based service provider shall take all reasonable steps to prevent further incidents. The community-based service provider shall provide the following internal monitoring and facilitating quality improvement program:

(1) community-based service providers shall have current abuse, neglect, and exploitation management policy and procedures in place that comply with the department's requirements;

(2) community-based service providers providing intellectual and developmental disabilities services must have a designated incident management coordinator in place; and

(3) community-based service providers providing intellectual and developmental disabilities services must have an incident management committee to identify any deficiencies, trends, patterns, or concerns as well as opportunities for quality improvement, address internal and external incident reports for the purpose of examining internal root causes, and to take action on identified issues.
Tag # 1A09
Medication Delivery
Routine Medication Administration

<table>
<thead>
<tr>
<th>MINIMUM STANDARDS:</th>
<th>Condition of Participation Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>A. MINIMUM STANDARDS FOR THE DISTRIBUTION, STORAGE, HANDLING AND RECORD KEEPING OF DRUGS:</td>
<td>After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur.</td>
</tr>
<tr>
<td>(d) The facility shall have a Medication Administration Record (MAR) documenting medication administered to residents, including over-the-counter medications.</td>
<td>Medication Administration Records (MAR) were reviewed for the months of July and August 2015.</td>
</tr>
<tr>
<td>This documentation shall include:</td>
<td>Based on record review, 4 of 6 individuals had Medication Administration Records (MAR), which contained missing medications entries and/or other errors:</td>
</tr>
<tr>
<td>(i) Name of resident;</td>
<td>Individual #2</td>
</tr>
<tr>
<td>(ii) Date given;</td>
<td>July 2015</td>
</tr>
<tr>
<td>(iii) Drug product name;</td>
<td>Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries:</td>
</tr>
<tr>
<td>(iv) Dosage and form;</td>
<td>• Lactulose 30 ml (2 times daily) – Blank 7/25 (8 PM)</td>
</tr>
<tr>
<td>(v) Strength of drug;</td>
<td>Medication Administration Record document did not contain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose for the following medications:</td>
</tr>
<tr>
<td>(vi) Route of administration;</td>
<td>• Ability 20 mg (1 time daily)</td>
</tr>
<tr>
<td>(vii) How often medication is to be taken;</td>
<td>• Abilify 5 mg (1 time daily)</td>
</tr>
<tr>
<td>(viii) Time taken and staff initials;</td>
<td>• Anafranil 100 mg (1 time daily)</td>
</tr>
<tr>
<td>(ix) Dates when the medication is discontinued or changed;</td>
<td>• Ativan 1 mg (4 times daily)</td>
</tr>
<tr>
<td>(x) The name and initials of all staff administering medications.</td>
<td>• Benztropine 1 mg (2 times daily)</td>
</tr>
</tbody>
</table>

Model Custodial Procedure Manual

D. Administration of Drugs

Unless otherwise stated by practitioner, patients will not be allowed to administer their own medications. Document the practitioner’s order authorizing the self-administration of medications.

All PRN (As needed) medications shall have complete detail instructions regarding the administering of the medication. This shall include:

- symptoms that indicate the use of the medication,
exact dosage to be used, and
the exact amount to be used in a 24 hour period.


CHAPTER 5 (CIES) 1. Scope of Service B. Self Employment 8. Providing assistance with medication delivery as outlined in the ISP; C. Individual Community Integrated Employment 3. Providing assistance with medication delivery as outlined in the ISP; D. Group Community Integrated Employment 4. Providing assistance with medication delivery as outlined in the ISP; and

B. Community Integrated Employment Agency Staffing Requirements: o. Comply with DDSD Medication Assessment and Delivery Policy and Procedures;

CHAPTER 6 (CCS) 1. Scope of Services A. Individualized Customized Community Supports 19. Providing assistance or supports with medications in accordance with DDSD Medication Assessment and Delivery policy. C. Small Group Customized Community Supports 19. Providing assistance or supports with medications in accordance with DDSD Medication Assessment and Delivery policy. D. Group Customized Community Supports 19. Providing assistance or supports with medications in accordance with DDSD Medication Assessment and Delivery policy.

CHAPTER 11 (FL) 1 SCOPE OF SERVICES A. Living Supports- Family Living Services: The scope of Family Living Services includes, but is not limited to the following as identified by the Interdisciplinary Team (IDT):

- Flonase 1 spray (1 time daily)
- Lactulose 30 ml (2 times daily)
- Luvox 150 mg (2 times daily)
- Oxcarbazepine 150 mg (1 time daily)
- Oxcarbazepine 300 mg (1 time daily)
- Synthroid 100 mcg (1 time daily)

August 2015
Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries:
- Benadryl 50 mg (1 time daily) – Blank 8/1 - 3 (AM)
- Luvox (2 times daily)

Individual #3
July 2015
Medication Administration Record document did not contain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose for the following medications:
- Atenolol 25 mg (1 time daily)
- Ativan 1 mg (3 times daily)
- Baclofen 5 mg (2 times daily)
19. Assisting in medication delivery, and related monitoring, in accordance with the DDSD’s Medication Assessment and Delivery Policy, New Mexico Nurse Practice Act, and Board of Pharmacy regulations including skill development activities leading to the ability for individuals to self-administer medication as appropriate; and

I. Healthcare Requirements for Family Living.

3. B. Adult Nursing Services for medication oversight are required for all surrogate Living Supports- Family Living direct support personnel if the individual has regularly scheduled medication. Adult Nursing services for medication oversight are required for all surrogate Family Living Direct Support Personnel (including substitute care), if the individual has regularly scheduled medication.

6. Support Living - Family Living Provider Agencies must have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in accordance with DDSD Medication Assessment and Delivery Policy and Procedures, the New Mexico Nurse Practice Act and Board of Pharmacy standards and regulations.

   a. All twenty-four (24) hour residential home sites serving two (2) or more unrelated individuals must be licensed by the Board of Pharmacy, per current regulations;
   b. When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) must be maintained and include:

      i. The name of the individual, a transcription of the physician’s or licensed health care provider’s prescription including the brand and generic name of the medication, and

<table>
<thead>
<tr>
<th>Medication</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cogentin 5 mg</td>
<td>2 times daily</td>
</tr>
<tr>
<td>Flomax 4 mg</td>
<td>1 time daily</td>
</tr>
<tr>
<td>Geodon 80 mg</td>
<td>1 time daily</td>
</tr>
<tr>
<td>Geodon 20 mg</td>
<td>1 time daily</td>
</tr>
<tr>
<td>Glucophage 500 mg</td>
<td>1 time daily</td>
</tr>
<tr>
<td>Lovastatin 20 mg</td>
<td>1 time daily</td>
</tr>
<tr>
<td>Proscar 5 mg</td>
<td>1 time daily</td>
</tr>
<tr>
<td>Sertraline 50 mg</td>
<td>1 time daily</td>
</tr>
<tr>
<td>Zaditor 1 gttsp</td>
<td>2 times daily</td>
</tr>
</tbody>
</table>

August 2015
Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries:

- Ativan 1 mg (3 times daily) – Blank 8/4 (8AM and 2PM)

Individual #5
July 2015
Medication Administration Record document did not contain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose for the following medications:

- Cymbalta 60 mg (1 time daily)
- Depakote 500 mg (2 times daily)
- Dolxazsoin 4 mg (2 times daily)
diagnosis for which the medication is prescribed;
ii. Prescribed dosage, frequency and method/route of administration, times and dates of administration;
iii. Initials of the individual administering or assisting with the medication delivery;
iv. Explanation of any medication error;
v. Documentation of any allergic reaction or adverse medication effect; and
vi. For PRN medication, instructions for the use of the PRN medication must include observable signs/symptoms or circumstances in which the medication is to be used, and documentation of effectiveness of PRN medication administered.

c. The Family Living Provider Agency must also maintain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose; and
d. Information from the prescribing pharmacy regarding medications must be kept in the home and community inclusion service locations and must include the expected desired outcomes of administering the medication, signs and symptoms of adverse events and interactions with other medications.
e. Medication Oversight is optional if the individual resides with their biological family (by affinity or consanguinity). If Medication Oversight is not selected as an Ongoing Nursing Service, all elements of medication administration and oversight are the sole responsibility of the individual and their biological family. Therefore, a monthly medication administration record (MAR) is not required unless the family requests it and

<table>
<thead>
<tr>
<th>Medicine</th>
<th>Dosage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Enalapril 20 mg</td>
<td>1 times daily</td>
</tr>
<tr>
<td>Ensure 1 can</td>
<td>2 times daily</td>
</tr>
<tr>
<td>Ferrous Sulfate 325 mg</td>
<td>3 times daily</td>
</tr>
<tr>
<td>Multivitamin 1 unit</td>
<td>1 time daily</td>
</tr>
<tr>
<td>Natural Tears 1 gtts</td>
<td>4 times daily</td>
</tr>
<tr>
<td>Pravastatin 40 mg</td>
<td>1 time daily</td>
</tr>
<tr>
<td>Prilosec 20 mg</td>
<td>2 times daily</td>
</tr>
<tr>
<td>Proscar 5 mg</td>
<td>1 time daily</td>
</tr>
<tr>
<td>Vitamin C 500 mg</td>
<td>3 times daily</td>
</tr>
<tr>
<td>Zaditor 1 gtts</td>
<td>2 times daily</td>
</tr>
<tr>
<td>Zonegran 100 mg</td>
<td>2 times daily</td>
</tr>
</tbody>
</table>

Medication Administration Records contained missing entries. Documentation found indicated reason for missing entries was medication was not available:
- Multivitamin 1 unit (1 time daily) – Blank 7/28 (8 AM)
- Pravastatin 40 mg (1 time daily) – Blank 7/28 (8 PM)
- Proscar 5 mg (1 time daily) – Blank 7/28 (8 AM)
- Zonegran 100 mg (2 time daily) Blank 7/28 (8 AM)
continually communicates all medication changes to the provider agency in a timely manner to insure accuracy of the MAR.

i. The family must communicate at least annually and as needed for significant change of condition with the agency nurse regarding the current medications and the individual’s response to medications for purpose of accurately completing required nursing assessments.

ii. As per the DDSD Medication Assessment and Delivery Policy and Procedure, paid DSP who are not related by affinity or consanguinity to the individual may not deliver medications to the individual unless they have completed Assisting with Medication Delivery (AWMD) training. DSP may also be under a delegation relationship with a DDW agency nurse or be a Certified Medication Aide (CMA). Where CMAs are used, the agency is responsible for maintaining compliance with New Mexico Board of Nursing requirements.

iii. If the substitute care provider is a surrogate (not related by affinity or consanguinity) Medication Oversight must be selected and provided.

**CHAPTER 12 (SL) 2. Service Requirements L. Training and Requirements: 3. Medication Delivery:** Supported Living Provider Agencies must have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in accordance with DDSD Medication Assessment and Delivery Policy and Procedures, New Mexico Nurse Practice Act, and Board of Pharmacy standards and regulations.

<table>
<thead>
<tr>
<th>Individual #6</th>
<th>July 2015</th>
</tr>
</thead>
<tbody>
<tr>
<td>Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries:</td>
<td></td>
</tr>
<tr>
<td>• Acycovir Ointment (2 times daily) – Blank 7/1 (8 PM)</td>
<td></td>
</tr>
<tr>
<td>• Psyllium 2-4 unit (1 time daily) – Blank 7/5 (8 AM)</td>
<td></td>
</tr>
</tbody>
</table>

Medication Administration Record document did not contain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose for the following medications:

• Acycovir Ointment (2 times daily)
• Baclofen 20 mg (2 times daily)
• Multivitamin 1 unit (1 time daily)
• Nitrofurantoin 100 mg (1 time daily)
• Oxybutynin 10 mg (1 time daily)
• Polyethylene glycol ½ cap (1 time daily)
• Psyllium 2-4 unit (1 time daily)
• Vitamin D3 2000 units (1 time daily)
• Zink 50 mg (1 time daily)
• Zoloft 100 mg (1 time daily)

Medication Administration Records contained missing entries. Documentation found
h. All twenty-four (24) hour residential home sites serving two (2) or more unrelated individuals must be licensed by the Board of Pharmacy, per current regulations;

i. When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) must be maintained and include:

   i. The name of the individual, a transcription of the physician’s or licensed health care provider’s prescription including the brand and generic name of the medication, and diagnosis for which the medication is prescribed;

   ii. Prescribed dosage, frequency and method/route of administration, times and dates of administration;

   iii. Initials of the individual administering or assisting with the medication delivery;

   iv. Explanation of any medication error;

   v. Documentation of any allergic reaction or adverse medication effect; and

   vi. For PRN medication, instructions for the use of the PRN medication must include observable signs/symptoms or circumstances in which the medication is to be used, and documentation of effectiveness of PRN medication administered.

j. The Supported Living Provider Agency must also maintain a signature page that designates the full name that corresponds to the indicated reason for missing entries was medication was not available:

   • Zink 50 mg (1 time daily) – Blank 7/6-8 (8 AM)
each initial used to document administered or assisted delivery of each dose; and

k. Information from the prescribing pharmacy regarding medications must be kept in the home and community inclusion service locations and must include the expected desired outcomes of administrating the medication, signs, and symptoms of adverse events and interactions with other medications.

CHAPTER 13 (IMLS) 2. Service Requirements.
B. There must be compliance with all policy requirements for Intensive Medical Living Service Providers, including written policy and procedures regarding medication delivery and tracking and reporting of medication errors consistent with the DDSD Medication Delivery Policy and Procedures, relevant Board of Nursing Rules, and Pharmacy Board standards and regulations.


CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS:
E. Medication Delivery: Provider Agencies that provide Community Living, Community Inclusion or Private Duty Nursing services shall have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in accordance with DDSD Medication Assessment and Delivery Policy and Procedures, the Board of Nursing Rules and Board of Pharmacy standards and regulations.

(2) When required by the DDSD Medication Assessment and Delivery Policy, Medication
Administration Records (MAR) shall be maintained and include:
(a) The name of the individual, a transcription of the physician's written or licensed health care provider's prescription including the brand and generic name of the medication, diagnosis for which the medication is prescribed;
(b) Prescribed dosage, frequency and method/route of administration, times and dates of administration;
(c) Initials of the individual administering or assisting with the medication;
(d) Explanation of any medication irregularity;
(e) Documentation of any allergic reaction or adverse medication effect; and
(f) For PRN medication, an explanation for the use of the PRN medication shall include observable signs/symptoms or circumstances in which the medication is to be used, and documentation of effectiveness of PRN medication administered.

(3) The Provider Agency shall also maintain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose;
(4) MARs are not required for individuals participating in Independent Living who self-administer their own medications;
(5) Information from the prescribing pharmacy regarding medications shall be kept in the home and community inclusion service locations and shall include the expected desired outcomes of administering the medication, signs and symptoms of adverse events and interactions with other medications;
<table>
<thead>
<tr>
<th>Tag # 1A09.1</th>
<th>Standard Level Deficiency</th>
<th>Provider:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Medication Delivery</td>
<td>Medication Administration Records (MAR) were reviewed for the months of July and August, 2015.</td>
<td>State your Plan of Correction for the deficiencies cited in this tag here: →</td>
</tr>
<tr>
<td>PRN Medication Administration</td>
<td>Based on record review, 3 of 6 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard:</td>
<td></td>
</tr>
<tr>
<td>NMAC 16.19.11.8 MINIMUM STANDARDS:</td>
<td>Individual #2</td>
<td>Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →</td>
</tr>
<tr>
<td>A. MINIMUM STANDARDS FOR THE DISTRIBUTION, STORAGE, HANDLING AND RECORD KEEPING OF DRUGS:</td>
<td>July 2015</td>
<td></td>
</tr>
<tr>
<td>(d) The facility shall have a Medication Administration Record (MAR) documenting medication administered to residents, including over-the-counter medications. This documentation shall include:</td>
<td>No Effectiveness was noted on the Medication Administration Record for the following PRN medication:</td>
<td></td>
</tr>
<tr>
<td></td>
<td>(i) Name of resident;</td>
<td>• Ibuprofen 800 mg – PRN – 7/17/2015 (given 1 time).</td>
</tr>
<tr>
<td></td>
<td>(ii) Date given;</td>
<td></td>
</tr>
<tr>
<td></td>
<td>(iii) Drug product name;</td>
<td>Individual #4</td>
</tr>
<tr>
<td></td>
<td>(iv) Dosage and form;</td>
<td>July 2015</td>
</tr>
<tr>
<td></td>
<td>(v) Strength of drug;</td>
<td>No Effectiveness was noted on the Medication Administration Record for the following PRN medication:</td>
</tr>
<tr>
<td></td>
<td>(vi) Route of administration;</td>
<td>• Tylenol 325 mg – PRN – 7/18/2015 (given 1 time).</td>
</tr>
<tr>
<td></td>
<td>(vii) How often medication is to be taken;</td>
<td>Individual #5</td>
</tr>
<tr>
<td></td>
<td>(viii) Time taken and staff initials;</td>
<td>July 2015</td>
</tr>
<tr>
<td></td>
<td>(ix) Dates when the medication is discontinued or changed;</td>
<td>No evidence of documented Signs/Symptoms were found for the following PRN medication:</td>
</tr>
<tr>
<td></td>
<td>(x) The name and initials of all staff administering medications.</td>
<td>• Caloseptine (Lotion) – PRN – 7/10 -13, 21 – 23, 27 – 29, 2015 (given 1 - 2 times daily).</td>
</tr>
<tr>
<td>Model Custodial Procedure Manual</td>
<td></td>
<td></td>
</tr>
<tr>
<td>D. Administration of Drugs</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Unless otherwise stated by practitioner, patients will not be allowed to administer their own medications. Document the practitioner’s order authorizing the self-administration of medications.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>All PRN (As needed) medications shall have complete detail instructions regarding the administering of the medication. This shall include:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>➢ symptoms that indicate the use of the medication,</td>
<td></td>
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</table>
exact dosage to be used, and
the exact amount to be used in a 24 hour
period.

**Department of Health Developmental Disabilities Supports Division (DDSD) Medication Assessment and Delivery Policy - Eff. November 1, 2006**

**F. PRN Medication**

3. Prior to self-administration, self-administration with physical assist or assisting with delivery of PRN medications, the direct support staff must contact the agency nurse to describe observed symptoms and thus assure that the PRN medication is being used according to instructions given by the ordering PCP. In cases of fever, respiratory distress (including coughing), severe pain, vomiting, diarrhea, change in responsiveness/level of consciousness, the nurse must strongly consider the need to conduct a face-to-face assessment to assure that the PRN does not mask a condition better treated by seeking medical attention. This does not apply to home based/family living settings where the provider is related by affinity or by consanguinity to the individual.

4. The agency nurse shall review the utilization of PRN medications routinely. Frequent or escalating use of PRN medications must be reported to the PCP and discussed by the Interdisciplinary for changes to the overall support plan (see Section H of this policy).

**H. Agency Nurse Monitoring**

1. Regardless of the level of assistance with medication delivery that is required by the individual or the route through which the medication is delivered, the agency nurses
must monitor the individual’s response to the effects of their routine and PRN medications. The frequency and type of monitoring must be based on the nurse’s assessment of the individual and consideration of the individual’s diagnoses, health status, stability, utilization of PRN medications and level of support required by the individual’s condition and the skill level and needs of the direct care staff. Nursing monitoring should be based on prudent nursing practice and should support the safety and independence of the individual in the community setting. The health care plan shall reflect the planned monitoring of the individual’s response to medication.

Department of Health Developmental Disabilities Supports Division (DDSD) - Procedure Title: Medication Assessment and Delivery

Procedure Eff Date: November 1, 2006

C. 3. Prior to delivery of the PRN, direct support staff must contact the agency nurse to describe observed symptoms and thus assure that the PRN is being used according to instructions given by the ordering PCP. In cases of fever, respiratory distress (including coughing), severe pain, vomiting, diarrhea, change in responsiveness/level of consciousness, the nurse must strongly consider the need to conduct a face-to-face assessment to assure that the PRN does not mask a condition better treated by seeking medical attention.

(References: Psychotropic Medication Use Policy, Section D, page 5 Use of PRN Psychotropic Medications; and, Human Rights Committee Requirements Policy, Section B, page 4 Interventions Requiring Review and Approval – Use of PRN Medications).
a. Document conversation with nurse including all reported signs and symptoms, advice given and action taken by staff.

4. Document on the MAR each time a PRN medication is used and describe its effect on the individual (e.g., temperature down, vomiting lessened, anxiety increased, the condition is the same, improved, or worsened, etc.).


CHAPTER 11 (FL) 1 SCOPE OF SERVICES
A. Living Supports- Family Living Services:
The scope of Family Living Services includes, but is not limited to the following as identified by the Interdisciplinary Team (IDT):

19. Assisting in medication delivery, and related monitoring, in accordance with the DDSD’s Medication Assessment and Delivery Policy, New Mexico Nurse Practice Act, and Board of Pharmacy regulations including skill development activities leading to the ability for individuals to self-administer medication as appropriate; and

I. Healthcare Requirements for Family Living.
3. B. Adult Nursing Services for medication oversight are required for all surrogate Family Living direct support personnel if the individual has regularly scheduled medication. Adult Nursing services for medication oversight are required for all surrogate Family Living Direct Support Personnel (including substitute care), if the individual has regularly scheduled medication.

6. Support Living- Family Living Provider Agencies must have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in
f. All twenty-four (24) hour residential home sites serving two (2) or more unrelated individuals must be licensed by the Board of Pharmacy, per current regulations;
g. When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) must be maintained and include:

i. The name of the individual, a transcription of the physician’s or licensed health care provider’s prescription including the brand and generic name of the medication, and diagnosis for which the medication is prescribed;
ii. Prescribed dosage, frequency and method/route of administration, times and dates of administration;
iii. Initials of the individual administering or assisting with the medication delivery;
iv. Explanation of any medication error;
v. Documentation of any allergic reaction or adverse medication effect; and
vi. For PRN medication, instructions for the use of the PRN medication must include observable signs/symptoms or circumstances in which the medication is to be used, and documentation of effectiveness of PRN medication administered.

h. The Family Living Provider Agency must also maintain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose; and
<p>| | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>i.</td>
<td>Information from the prescribing pharmacy regarding medications must be kept in the home and community inclusion service locations and must include the expected desired outcomes of administering the medication, signs and symptoms of adverse events and interactions with other medications.</td>
</tr>
<tr>
<td>j.</td>
<td>Medication Oversight is optional if the individual resides with their biological family (by affinity or consanguinity). If Medication Oversight is not selected as an Ongoing Nursing Service, all elements of medication administration and oversight are the sole responsibility of the individual and their biological family. Therefore, a monthly medication administration record (MAR) is not required unless the family requests it and continually communicates all medication changes to the provider agency in a timely manner to insure accuracy of the MAR.</td>
</tr>
<tr>
<td>iv.</td>
<td>The family must communicate at least annually and as needed for significant change of condition with the agency nurse regarding the current medications and the individual’s response to medications for purpose of accurately completing required nursing assessments.</td>
</tr>
</tbody>
</table>
v. | As per the DDSD Medication Assessment and Delivery Policy and Procedure, paid DSP who are not related by affinity or consanguinity to the individual may not deliver medications to the individual unless they have completed Assisting with Medication Delivery (AWMD) training. DSP may also be under a delegation relationship with a DDW agency nurse or be a Certified Medication Aide (CMA). Where CMAs are used, the agency is responsible for |
maintaining compliance with New Mexico Board of Nursing requirements.

vi. If the substitute care provider is a surrogate (not related by affinity or consanguinity) Medication Oversight must be selected and provided.

CHAPTER 12 (SL) 2. Service Requirements L. Training and Requirements: 3. Medication Delivery: Supported Living Provider Agencies must have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in accordance with DDSD Medication Assessment and Delivery Policy and Procedures, New Mexico Nurse Practice Act, and Board of Pharmacy standards and regulations.

l. All twenty-four (24) hour residential home sites serving two (2) or more unrelated individuals must be licensed by the Board of Pharmacy, per current regulations;

h. When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) must be maintained and include:

i. The name of the individual, a transcription of the physician’s or licensed health care provider’s prescription including the brand and generic name of the medication, and diagnosis for which the medication is prescribed;

ii. Prescribed dosage, frequency and method/route of administration, times and dates of administration;
iii. Initials of the individual administering or assisting with the medication delivery;

iv. Explanation of any medication error;

v. Documentation of any allergic reaction or adverse medication effect; and

vi. For PRN medication, instructions for the use of the PRN medication must include observable signs/symptoms or circumstances in which the medication is to be used, and documentation of effectiveness of PRN medication administered.

n. The Supported Living Provider Agency must also maintain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose; and

o. Information from the prescribing pharmacy regarding medications must be kept in the home and community inclusion service locations and must include the expected desired outcomes of administrating the medication, signs, and symptoms of adverse events and interactions with other medications.

CHAPTER 13 (IMLS) 2. Service Requirements. B. There must be compliance with all policy requirements for Intensive Medical Living Service Providers, including written policy and procedures regarding medication delivery and tracking and reporting of medication errors consistent with the DDSD Medication Delivery Policy and Procedures,
relevant Board of Nursing Rules, and Pharmacy Board standards and regulations.


CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS: The objective of these standards is to establish Provider Agency policy, procedure and reporting requirements for DD Medicaid Waiver program. These requirements apply to all such Provider Agency staff, whether directly employed or subcontracting with the Provider Agency. Additional Provider Agency requirements and personnel qualifications may be applicable for specific service standards.

E. Medication Delivery: Provider Agencies that provide Community Living, Community Inclusion or Private Duty Nursing services shall have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in accordance with DDSD Medication Assessment and Delivery Policy and Procedures, the Board of Nursing Rules and Board of Pharmacy standards and regulations.

(2) When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) shall be maintained and include:
   (a) The name of the individual, a transcription of the physician's written or licensed health care provider's prescription including the brand and generic name of the medication, diagnosis for which the medication is prescribed;
(b) Prescribed dosage, frequency and method/route of administration, times and dates of administration;
(c) Initials of the individual administering or assisting with the medication;
(d) Explanation of any medication irregularity;
(e) Documentation of any allergic reaction or adverse medication effect; and
(f) For PRN medication, an explanation for the use of the PRN medication shall include observable signs/symptoms or circumstances in which the medication is to be used, and documentation of effectiveness of PRN medication administered.

(3) The Provider Agency shall also maintain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose;

(4) MARs are not required for individuals participating in Independent Living who self-administer their own medications;

(5) Information from the prescribing pharmacy regarding medications shall be kept in the home and community inclusion service locations and shall include the expected desired outcomes of administering the medication, signs and symptoms of adverse events and interactions with other medications;
<table>
<thead>
<tr>
<th>Tag # 1A11</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Transportation Policy and Procedure</strong></td>
</tr>
<tr>
<td><strong>Standard Level Deficiency</strong></td>
</tr>
<tr>
<td><strong>STATE OF NEW MEXICO DEPARTMENT OF HEALTH DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION PROVIDER AGREEMENT:</strong></td>
</tr>
<tr>
<td><strong>ARTICLE 2. APPLICABLE LAWS:</strong> This Provider Agreement shall be governed by the laws of the State of New Mexico.</td>
</tr>
<tr>
<td><strong>Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy:</strong></td>
</tr>
<tr>
<td><strong>Training Requirements for Direct Service Agency Staff Policy Eff Date:</strong> March 1, 2007</td>
</tr>
<tr>
<td><strong>II. POLICY STATEMENTS:</strong></td>
</tr>
<tr>
<td>I. Staff providing direct services shall complete safety training within the first thirty (30) days of employment and before working alone with an individual receiving services. The training shall address at least the following:</td>
</tr>
<tr>
<td>1. Operating a fire extinguisher</td>
</tr>
<tr>
<td>2. Proper lifting procedures</td>
</tr>
<tr>
<td>3. General vehicle safety precautions (e.g., pre-trip inspection, removing keys from the ignition when not in the driver’s seat)</td>
</tr>
<tr>
<td>4. Assisting passengers with cognitive and/or physical impairments (e.g., general guidelines for supporting individuals who may be unaware of safety issues involving traffic or those who require physical assistance to enter/exit a vehicle)</td>
</tr>
<tr>
<td>5. Operating wheelchair lifts (if applicable to the staff’s role)</td>
</tr>
<tr>
<td>6. Wheelchair tie-down procedures (if applicable to the staff’s role)</td>
</tr>
<tr>
<td>7. Emergency and evacuation procedures (e.g., roadside emergency, fire emergency)</td>
</tr>
<tr>
<td><strong>Based on record review, the Agency did not have written policies and procedures regarding the safe transportation of individuals in the community, which comply with New Mexico regulations governing the operation of motor vehicles to transport individuals.</strong></td>
</tr>
<tr>
<td><strong>Review of Agency’s policies and procedures indicated the following elements were not found:</strong></td>
</tr>
<tr>
<td>(1) Operating a fire extinguisher</td>
</tr>
<tr>
<td>(2) Individual safety, including safe locations for boarding and disembarking passengers, appropriate responses to hazardous weather and other adverse driving conditions,</td>
</tr>
<tr>
<td>(4) Staff training regarding the safe operation of the vehicle, assisting passengers and safe lifting procedures,</td>
</tr>
<tr>
<td>(5) Emergency Plans, including vehicle evacuation techniques,</td>
</tr>
<tr>
<td><strong>Provider:</strong></td>
</tr>
<tr>
<td><strong>State your Plan of Correction for the deficiencies cited in this tag here:</strong></td>
</tr>
<tr>
<td><strong>Provider:</strong></td>
</tr>
<tr>
<td><strong>Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here:</strong></td>
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</tbody>
</table>

QMB Report of Findings – The Opportunity Center, Inc. – Southwest Region – August 3 – 6, 2015

Survey Report #: Q.16.1.DDW.D1556.3.RTN.01.15.261
transportation during Job Development activities; and B. Self Employment: 7. Arranging or providing transportation during Job Development activities; and C. Integrated Employment Services: 2. Arranging or providing transportation or supporting public transportation during Individual Community Integrated Employment Services; Integrated Employment Services: D. 3. Arranging or providing transportation or supporting public transportation during Group Community Integrated Employment Services;

CHAPTER 6 (CCS) I. Scope of Service A. Individualized Customized Community Supports 17. Providing transportation or assisting with transportation arrangements for participating in Customized Community Supports; C. Small Group Customized Community Supports 17. Providing or assisting with transportation during provision of Customized Community Supports; D. Group Customized Community Supports 17. Providing or assisting with transportation during provision of Customized Community Supports;

CHAPTER 11 (FL) 2. Service Requirements: I. Healthcare Requirements for Family Living: 10. Family Living provider agencies must have a written policy and procedures regarding the safe transportation of individuals in the community, and comply with New Mexico regulations governing the operation of motor vehicles to transport individuals, and which are consistent with DDSD guidelines issued July 1, 1999 titled “Client Transportation Safety”. The policy and procedures must address at least the following topics:

a. Drivers’ requirements;
b. Individual safety, including safe locations for boarding and disembarking passengers, appropriate responses to hazardous weather and other adverse driving conditions;
c. Vehicle maintenance and safety inspections;
d. DSP training regarding the safe operation of the vehicle, assisting passengers and safe lifting procedures;

e. Emergency Plans, including vehicle evacuation techniques;

f. Accident Procedures; and

g. Written documentation of vehicle maintenance, safety inspections, and staffing training.

CHAPTER 12 (SL) 2. Service Requirements: L. Training and Requirements 7. Transportation:
Supported Living provider agencies must have a written policy and procedures regarding the safe transportation of individuals in the community, and comply with New Mexico regulations governing the operation of motor vehicles to transport individuals, and which are consistent with DDSD guidelines issued July 1, 1999 titled “Client Transportation Safety.” The policy and procedures must address at least the following topics:

a. Drivers’ requirements;

b. Individual safety, including safe locations for boarding and disembarking passengers, appropriate responses to hazardous weather and other adverse driving conditions;

c. Vehicle maintenance and safety inspections;

d. DSP training regarding the safe operation of the vehicle, assisting passengers and safe lifting procedures;

e. Emergency Plans, including vehicle evacuation techniques;

f. Accident Procedures; and

g. Written documentation of vehicle maintenance, safety inspections, and staffing training.

CHAPTER 13 (IMLS) 2. Service Requirements: N. Services provider agencies must develop and implement policies and procedures regarding the safe transportation of individuals in the community which comply with New Mexico regulations governing operation of motor vehicles to transport individuals and which are consistent with DDSD
guidelines issued July 1, 1999 titled “Client Transportation Safety”. The policy and procedures must address at least the following:

1. Documented evidence of driver requirements;
2. Individual safety including locations for boarding and disembarking passengers, and appropriate response to hazardous weather and other adverse driving conditions, including securing all equipment and supplies needed to assure health and safety during transport;
3. Vehicle maintenance and safety inspections;
4. Documented evidence of driver training regarding safe operation of the vehicle, assisting passengers, and safe lifting procedures;
5. Emergency plans including vehicle evacuation techniques; and
6. Accident procedures.
### Tag # 1A27.2
Duty to Report IRs Filed During On-Site and/or IRs Not Reported by Provider

<table>
<thead>
<tr>
<th>Standard Level Deficiency</th>
<th>Provider: State your Plan of Correction for the deficiencies cited in this tag here: →</th>
</tr>
</thead>
<tbody>
<tr>
<td>NMAC 7.1.14 ABUSE, NEGLECT, EXPLOITATION, AND DEATH REPORTING, TRAINING AND RELATED REQUIREMENTS FOR COMMUNITY PROVIDERS</td>
<td>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →</td>
</tr>
<tr>
<td>NMAC 7.1.14.8 INCIDENT MANAGEMENT SYSTEM REPORTING REQUIREMENTS FOR COMMUNITY-BASED SERVICE PROVIDERS:</td>
<td></td>
</tr>
<tr>
<td>A. Duty to report:</td>
<td></td>
</tr>
<tr>
<td>(1) All community-based providers shall immediately report alleged crimes to law enforcement or call for emergency medical services as appropriate to ensure the safety of consumers.</td>
<td></td>
</tr>
<tr>
<td>(2) All community-based service providers, their employees and volunteers shall immediately call the department of health improvement (DHI) hotline at 1-800-445-6242 to report abuse, neglect, exploitation, suspicious injuries or any death and also to report an environmentally hazardous condition which creates an immediate threat to health or safety.</td>
<td></td>
</tr>
<tr>
<td>B. Reporter requirement. All community-based service providers shall ensure that the employee or volunteer with knowledge of the alleged abuse, neglect, exploitation, suspicious injury, or death calls the division’s hotline to report the incident.</td>
<td></td>
</tr>
<tr>
<td>C. Initial reports, form of report, immediate action and safety planning, evidence preservation, required initial notifications:</td>
<td></td>
</tr>
<tr>
<td>(1) Abuse, neglect, and exploitation, suspicious injury or death reporting: Any person may report an allegation of abuse, neglect, or exploitation, suspicious injury or a</td>
<td></td>
</tr>
</tbody>
</table>

Based on record review, the Agency did not report suspected abuse, neglect, or exploitation, unexpected and natural/expected deaths; or other reportable incidents to the Division of Health Improvement for 2 of 7 Individuals.

During the on-site survey 8/3 - 6, 2015, surveyors found evidence of 2 internal agency incident reports, which had not been reported to DHI, as required by regulation.

The following internal incidents were reported as a result of the on-site survey:

**Individual #5**
- Incident date 5/5/2015 (7 am). Type of incident identified was abuse. Incident was brought to the attention of the Agency by Surveyor. Incident report was filed on 8/27/2015 by DHI/QMB.

**Individual #6**
- Incident date 5/5/2015 (7 am). Type of incident identified was abuse. Incident was brought to the attention of the Agency by Surveyor. Incident report was filed on 8/27/2015 by DHI/QMB.
death by calling the division’s toll-free hotline number 1-800-445-6242. Any consumer, family member, or legal guardian may call the division’s hotline to report an allegation of abuse, neglect, or exploitation, suspicious injury or death directly, or may report through the community-based service provider who, in addition to calling the hotline, must also utilize the division’s abuse, neglect, and exploitation or report of death form. The abuse, neglect, and exploitation or report of death form and instructions for its completion and filing are available at the division's website, http://dhi.health.state.nm.us, or may be obtained from the department by calling the division’s toll free hotline number, 1-800-445-6242.

(2) **Use of abuse, neglect, and exploitation or report of death form and notification by community-based service providers:** In addition to calling the division’s hotline as required in Paragraph (2) of Subsection A of 7.1.14.8 NMAC, the community-based service provider shall also report the incident of abuse, neglect, exploitation, suspicious injury, or death utilizing the division’s abuse, neglect, and exploitation or report of death form consistent with the requirements of the division’s abuse, neglect, and exploitation reporting guide. The community-based service provider shall ensure all abuse, neglect, exploitation or death reports describing the alleged incident are completed on the division’s abuse, neglect, and exploitation or report of death form and received by the division within 24 hours of the verbal report. If the provider has internet access, the report form shall be submitted via the division’s website at http://dhi.health.state.nm.us; otherwise it may be submitted via fax to 1-800-584-6057. The
community-based service provider shall ensure that the reporter with the most direct knowledge of the incident participates in the preparation of the report form.

(3) **Limited provider investigation:** No investigation beyond that necessary in order to be able to report the abuse, neglect, or exploitation and ensure the safety of consumers is permitted until the division has completed its investigation.

(4) **Immediate action and safety planning:** Upon discovery of any alleged incident of abuse, neglect, or exploitation, the community-based service provider shall:
   (a) develop and implement an immediate action and safety plan for any potentially endangered consumers, if applicable;
   (b) be immediately prepared to report that immediate action and safety plan verbally, and revise the plan according to the division’s direction, if necessary; and
   (c) provide the accepted immediate action and safety plan in writing on the immediate action and safety plan form within 24 hours of the verbal report. If the provider has internet access, the report form shall be submitted via the division’s website at http://dhi.health.state.nm.us; otherwise it may be submitted by faxing it to the division at 1-800-584-6057.

(5) **Evidence preservation:** The community-based service provider shall preserve evidence related to an alleged incident of abuse, neglect, or exploitation, including records, and do nothing to disturb the evidence. If physical evidence must be removed or affected, the provider shall take photographs or do whatever is reasonable to document the location and type of evidence found which appears related to the incident.
(6) **Legal guardian or parental notification:** The responsible community-based service provider shall ensure that the consumer’s legal guardian or parent is notified of the alleged incident of abuse, neglect and exploitation within 24 hours of notice of the alleged incident unless the parent or legal guardian is suspected of committing the alleged abuse, neglect, or exploitation, in which case the community-based service provider shall leave notification to the division’s investigative representative.

(7) **Case manager or consultant notification by community-based service providers:** The responsible community-based service provider shall notify the consumer’s case manager or consultant within 24 hours that an alleged incident involving abuse, neglect, or exploitation has been reported to the division. Names of other consumers and employees may be redacted before any documentation is forwarded to a case manager or consultant.

(8) **Non-responsible reporter:** Providers who are reporting an incident in which they are not the responsible community-based service provider shall notify the responsible community-based service provider within 24 hours of an incident or allegation of an incident of abuse, neglect, and exploitation.
<table>
<thead>
<tr>
<th>Tag # 1A28.2</th>
<th>Incident Mgt. System - Parent/Guardian Training</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>7.1.14.9 INCIDENT MANAGEMENT SYSTEM REQUIREMENTS:</td>
<td>Based on record review, the Agency did not provide documentation indicating consumer, family members, or legal guardians had received an orientation packet including incident management system policies and procedural information concerning the reporting of Abuse, Neglect and Exploitation, for 1 of 7 individuals.</td>
<td></td>
</tr>
<tr>
<td><strong>A. General:</strong> All community-based service providers shall establish and maintain an incident management system, which emphasizes the principles of prevention and staff involvement. The community-based service provider shall ensure that the incident management system policies and procedures require all employees and volunteers to be competently trained to respond to, report, and preserve evidence related to incidents in a timely and accurate manner.</td>
<td>Provider: State your Plan of Correction for the deficiencies cited in this tag here: →</td>
<td></td>
</tr>
<tr>
<td><strong>E. Consumer and guardian orientation packet:</strong> Consumers, family members, and legal guardians shall be made aware of and have available immediate access to the community-based service provider incident reporting processes. The community-based service provider shall provide consumers, family members, or legal guardians an orientation packet to include incident management systems policies and procedural information concerning the reporting of abuse, neglect, exploitation, suspicious injury, or death. The community-based service provider shall include a signed statement indicating the date, time, and place they received their orientation packet to be contained in the consumer’s file. The appropriate consumer, family member, or legal guardian shall sign this at the time of orientation.</td>
<td>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →</td>
<td></td>
</tr>
</tbody>
</table>

- Parent/Guardian Incident Management Training (Abuse, Neglect and Exploitation) (#2)
<table>
<thead>
<tr>
<th>Tag #1A39</th>
<th>Standard Level Deficiency</th>
<th>Provider: State your Plan of Correction for the deficiencies cited in this tag here: →</th>
</tr>
</thead>
</table>
| **Assistive Technology and Adaptive Equipment** | **CHAPTER 1. III. PROVIDER AGENCY DOCUMENTATION OF SERVICE DELIVERY AND LOCATION**  
**F. Sanitation:**  
(1) Equipment and utensils shall be kept clean and in good repair; and | ![Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →](#) |
| **7.26.5.13 DEVELOPMENT OF THE INDIVIDUAL SERVICE PLAN (ISP) - ASSESSMENTS:** | **7.26.5.14 DEVELOPMENT OF THE INDIVIDUAL SERVICE PLAN (ISP) - CONTENT OF INDIVIDUAL SERVICE PLANS:** Each ISP shall contain: | ![](#) |
| **F. Assistive technology:** Necessary support mechanisms and devices, including the rationale for the use of assistive technology or adaptive equipment when a need has been identified shall be documented in the ISP. The rationale shall include the environments and situations in which assistive technology is used. Selection of assistive technology shall support the individual's independence and functional capabilities in as non-intrusive a fashion as possible. | **Based on record review, observation and interview the Agency did not ensure the necessary support mechanisms and devices, including the rationale for the use of assistive technology or adaptive equipment as in place for 2 of 6 Individuals.**  
**During interview DSP were asked if the Individual had any assistive device or adaptive equipment and was it in functioning order.**  
- DSP #208 reported the following, “No his iPad and electric wheel chair are broken and have been for about 7 months or so.” (Individual #4)  
Review of documents indicated an iPad for communication, and an electric wheelchair were required to be used by the Individual.  
**During interview DSP were asked if the Individual had any assistive device or adaptive equipment and was it in functioning order.**  
- DSP # 202 reported the following, “he refuses to use it.” (Individual #5)  
According to the Health and section of the ISP a C Pap is needed for sleep apnea and is to be maintained by the Occupational Center Nurse. During home visit on 08/03/2015 Surveyors observed the Individual’s C Pap machine in a box and written on the box was the following | ![](#) |
| **CHAPTER 5 VI. SCOPE OF SUPPORTED EMPLOYMENT SERVICES**  
(7) Facilitating job accommodations and use of assistive technology, including the use of communication devices; | | |
| **CHAPTER 5 VII. SUPPORTED EMPLOYMENT SERVICES REQUIREMENTS**  
**D. Provider Agency Requirements**  
(6) Qualification and Competencies for Supported Employment Staff (includes intensive): Qualifications and competencies for staff providing | | |
| | | |
job coaching/consultation services shall, at a minimum, are able to:

CHAPTER 5 XI. COMMUNITY ACCESS SERVICES REQUIREMENTS
F. Community Access Services Provider Agency Staff Qualifications and Competencies
(1) Qualifications and Competencies for Community Access Coaches. The Community Access Coach shall, at a minimum, demonstrate the ability to:

(q) Communicate effectively with the individual including communication through the use of adaptive equipment and use of a communication dictionary when the individual uses these modes of communication;

(j) Communicate effectively with the individual including communication through the use of adaptive equipment as well as the individual’s Communication Dictionary, if applicable, at the work site;

CHAPTER 6. II. SCOPE OF COMMUNITY LIVING SERVICES.
A. The scope of Community Living Services includes, but is not limited the following as identified by the IDT:

(8) Implementation of the ISP, Therapy, Meal-time, Positive Behavioral Supports, Health Care, and Crisis Prevention/Interventions Plans, if applicable;

(9) Assistance in developing health maintenance supports, as well as monitoring the effectiveness of such supports;

(12) Assist the individual as needed, in coordination with the designated healthcare
coordinator and others on the IDT, with access to medical, dental, therapy, nutritional, behavioral and nursing practitioners and in the timely implementation of healthcare orders, monitoring and recording of therapeutic plans or activities as prescribed, to include: health care and crisis prevention/intervention plans;

CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS

H. Community Living Services Provider Agency Staffing Requirements

(1) Community Living Service Staff Qualifications and Competencies: Individuals working as direct support staff and supervisors for Community Living Service Provider Agencies shall demonstrate the following:

(b) The ability to assist the individual to meet his or her physical (e.g., health, grooming, toileting, eating) and personal management needs, by teaching skills, providing supports, and building on individual strengths and capabilities;

L. Residence Requirements for Family Living Services and Supported Living Services

(1) Supported Living Services and Family Living Services providers shall assure that each individual’s residence has:

(5) Kitchen area shall:

(b) Arrangements will be made, in consultation with the IDT for environmental accommodations and assistive technology devices specific to the needs of the individual(s); and
Tag # LS25 / 6L25  
Residential Health and Safety (SL/FL)  

<table>
<thead>
<tr>
<th>Standard Level Deficiency</th>
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</thead>
<tbody>
<tr>
<td>Based on observation, the Agency did not ensure that each individual’s residence met all requirements within the standard for 3 of 4 Supported Living.</td>
</tr>
<tr>
<td>Review of the residential records and observation of the residence revealed the following items were not found, not functioning or incomplete:</td>
</tr>
<tr>
<td><strong>Supported Living Requirements:</strong></td>
</tr>
<tr>
<td>- Battery operated or electric smoke detectors, heat sensors, or a sprinkler system installed in the residence (#2, 3)</td>
</tr>
<tr>
<td>- Water temperature in home does not exceed safe temperature (110°F)</td>
</tr>
<tr>
<td>- Water temperature in home measured 128.1°F (#2, 3)</td>
</tr>
<tr>
<td>- Water temperature in home measured 139°F (#4)</td>
</tr>
<tr>
<td>- Accessible written procedures for the safe storage of all medications with dispensing instructions for each individual that are consistent with the Assisting with Medication Administration training or each individual’s ISP (#2, 3, 5, 6)</td>
</tr>
<tr>
<td>- Accessible written procedures for emergency placement and relocation of individuals in the event of an emergency evacuation that makes the residence unsuitable for occupancy. The emergency evacuation procedures shall address, but are not limited</td>
</tr>
</tbody>
</table>

Provider:  
State your Plan of Correction for the deficiencies cited in this tag here: →

Provider:  
Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here: →

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Survey Report #: Q.16.1.DDW.D1556.3.RTN.01.15.261

Page 107 of 115
q. Have accessible written procedures for emergency placement and relocation of individuals in the event of an emergency evacuation that makes the residence unsuitable for occupancy. The emergency evacuation procedures must address, but are not limited to, fire, chemical and/or hazardous waste spills, and flooding.

**CHAPTER 12 (SL) Living Supports – Supported Living Agency Requirements G. Residence Requirements for Living Supports-Supported Living Services:** 1. Supported Living Provider Agencies must assure that each individual’s residence is maintained to be clean, safe, and comfortable and accommodates the individual’s daily living, social, and leisure activities. In addition the residence must:

   f. Maintain basic utilities, i.e., gas, power, water, and telephone;
   
g. Provide environmental accommodations and assistive technology devices in the residence including modifications to the bathroom (i.e., shower chairs, grab bars, walk in shower, raised toilets, etc.) based on the unique needs of the individual in consultation with the IDT;
   
h. Ensure water temperature in home does not exceed safe temperature (110°F);
   
i. Have a battery operated or electric smoke detectors and carbon monoxide detectors, fire extinguisher, or a sprinkler system;
   
j. Have a general-purpose First Aid kit;

   to, fire, chemical and/or hazardous waste spills, and flooding (#7)

**Note:** The following Individuals share a residence:

- #2, 3
- #5, 6
k. Allow at a maximum of two (2) individuals to share, with mutual consent, a bedroom and each individual has the right to have his or her own bed;

l. Have accessible written documentation of actual evacuation drills occurring at least three (3) times a year. For Supported Living evacuation drills must occur at least once a year during each shift;

m. Have accessible written procedures for the safe storage of all medications with dispensing instructions for each individual that are consistent with the Assisting with Medication Delivery training or each individual’s ISP; and

n. Have accessible written procedures for emergency placement and relocation of individuals in the event of an emergency evacuation that makes the residence unsuitable for occupancy. The emergency evacuation procedures must address, but are not limited to, fire, chemical and/or hazardous waste spills, and flooding.

CHAPTER 13 (IMLS) 2. Service Requirements R. Staff Qualifications: 3. Supervisor Qualifications And Requirements:
S Each residence shall include operable safety equipment, including but not limited to, an operable smoke detector or sprinkler system, a carbon monoxide detector if any natural gas appliance or heating is used, fire extinguisher, general purpose first aid kit, written procedures for emergency evacuation due to fire or other emergency and documentation of evacuation drills occurring at least annually during each shift, phone number for poison control within line of site of the telephone, basic utilities, general household appliances, kitchen and dining utensils, adequate food and drink for three
meals per day, proper food storage, and cleaning supplies.

T Each residence shall have a blood borne pathogens kit as applicable to the residents’ health status, personal protection equipment, and any ordered or required medical supplies shall also be available in the home.

U If not medically contraindicated, and with mutual consent, up to two (2) individuals may share a single bedroom. Each individual shall have their own bed. All bedrooms shall have doors that may be closed for privacy. Individuals have the right to decorate their bedroom in a style of their choosing consistent with safe and sanitary living conditions.

V For residences with more than two (2) residents, there shall be at least two (2) bathrooms. Toilets, tubs/showers used by the individuals shall provide for privacy and be designed or adapted for the safe provision of personal care. Water temperature shall be maintained at a safe level to prevent injury and ensure comfort and shall not exceed one hundred ten (110) degrees.


CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS
L. Residence Requirements for Family Living Services and Supported Living Services
Service Domain: Medicaid Billing/Reimbursement – State financial oversight exists to assure that claims are coded and paid for in accordance with the reimbursement methodology specified in the approved waiver.

TAG #1A12
All Services Reimbursement (No Deficiencies Found)


CHAPTER 5 (CIES) 6. REIMBURSEMENT All Provider Agencies must maintain all records necessary to fully disclose the type, quality, quantity and clinical necessity of services furnished to individuals who are currently receiving services. The Provider Agency records must be sufficiently detailed to substantiate the date, time, individual name, servicing provider, nature of services, and length of a session of service billed.

1. The documentation of the billable time spent with an individual must be kept on the written or electronic record that is prepared prior to a request for reimbursement from the HSD. For each unit billed, the record must contain the following:
   a. Date, start, and end time of each service encounter or other billable service interval;
   b. A description of what occurred during the encounter or service interval; and
   c. The signature or authenticated name of staff providing the service.

CHAPTER 6 (CCS) 4. REIMBURSEMENT A. Required Records: All Provider Agencies must maintain all records necessary to fully disclose the type, quality, quantity and clinical necessity of services furnished to individuals who are currently receiving services. The Provider Agency records must be sufficiently detailed to substantiate the date, time, individual name, servicing Provider Agency, nature of services, and length of a session of service billed.

1. The documentation of the billable time spent with an individual shall be kept on the written or electronic record that is prepared prior to a request for reimbursement from the Human Services Department (HSD). For each unit billed, the record shall contain the following:
   a. Date, start and end time of each service encounter or other billable service interval;
   b. A description of what occurred during the encounter or service interval; and
   c. The signature or authenticated name of staff providing the service.

CHAPTER 7 (CIHS) 4. REIMBURSEMENT. A. All Provider Agencies must maintain all records necessary to fully disclose the service, quality, quantity and clinical necessity furnished to individuals who are currently receiving services. The Provider Agency records shall be sufficiently detailed to substantiate the individual’s name, date, time, Provider Agency name, nature of services and length of a session of service billed.

1. The documentation of the billable time spent with an individual shall be kept on the written or electronic record that is prepared prior to a request for reimbursement from the Human Services Department (HSD). For each unit billed, the record shall contain the following:
   a. Date, start and end time of each service encounter or other billable service interval;
   b. A description of what occurred during the encounter or service interval; and
   c. The signature or authenticated name of staff providing the service.
CHAPTER 11 (FL) 4. REIMBURSEMENT A. Family Living Services Provider Agencies must maintain all records necessary to fully disclose the type, quality, quantity and clinical necessity of services furnished to individuals who are currently receiving services. The Family Living Services Provider Agency records must be sufficiently detailed to substantiate the date, time, individual name, servicing provider, nature of services, and length of a session of service billed.

1. The documentation of the billable time spent with an individual must be kept on the written or electronic record that is prepared prior to a request for reimbursement from the Human Services Department (HSD). For each unit billed, the record must contain the following:
   a. Date, start and end time of each service encounter or other billable service interval;
   b. A description of what occurred during the encounter or service interval; and
   c. The signature or authenticated name of staff providing the service.

CHAPTER 12 (SL) 2. REIMBURSEMENT
A. Supported Living Provider Agencies must maintain all records necessary to fully disclose the type, quality, quantity, and clinical necessity of services furnished to individuals who are currently receiving services. The Supported Living Services Provider Agency records must be sufficiently detailed to substantiate the date, time, individual name, servicing provider, nature of services, and length of a session of service billed.

1. The documentation of the billable time spent with an individual must be kept on the written or electronic record that is prepared prior to a request for reimbursement from the Human Services Department (HSD). For each unit billed, the record must contain the following:
   a. Date, start and end time of each service encounter or other billable service interval;
   b. A description of what occurred during the encounter or service interval;
   c. The signature or authenticated name of staff providing the service;

CHAPTER 13 (IMLS) 1. REIMBURSEMENT
A. All Living Supports- Intensive Medical Living Services Provider Agencies must maintain all records necessary to fully disclose the type, quality, quantity, and clinical necessity of services furnished to individuals who are currently receiving services. The Intensive Medical Living Services Provider Agency records shall be sufficiently detailed to substantiate the date, time, individual’s name, servicing provider, nature of services, and length of a session of service billed.

1. The documentation of the billable time spent with an individual shall be kept on the written or electronic record that is prepared prior to a request for reimbursement from the Human Services Department (HSD). For each unity billed, the record shall contain the following:
   a. Date, start and end time of each service encounter or other billable service interval;
   b. A description of what occurred during the encounter or service interval; and
   c. The signature or authenticated name of staff providing the service

Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 Chapter 1. III. PROVIDER AGENCY DOCUMENTATION OF SERVICE DELIVERY AND LOCATION B. Billable Units: The documentation of the billable time spent with an individual shall be kept on the written or electronic record that is prepared prior to a request for reimbursement from the HSD. For each unit billed, the record shall contain the following:

(1) Date, start and end time of each service encounter or other billable service interval;
(2) A description of what occurred during the encounter or service interval; and
(3) The signature or authenticated name of staff providing the service.
Billing for **2012**: Living Supports (Supported Living); Inclusion Supports (Customized Community Supports, Community Integrated Employment Services) and Other (Customized In-Home Supports) services was reviewed for 7 of 7 individuals. *Progress notes and billing records supported billing activities for the months of April, May and June 2015.*
Date: January 4, 2016

To: Christina Martinez, Executive Director
Provider: The Opportunity Center, Inc.
Address: 905 Tenth Street
State/Zip: Alamogordo, New Mexico 88310

E-mail Address: christina_oppcenter@hotmail.com

CC: Philip Gutierrez, President of the Board
Address: 1300 N. White Sands
State/Zip: Alamogordo, New Mexico 88310

Vice President: Dr. Norman Lindley
E-Mail Address: elmedico23@hotmail.com

Region: Southwest
Survey Date: August 3 - 6, 2015
Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: 2012: Living Supports (Supported Living); Inclusion Supports (Customized Community Supports, Community Integrated Employment Services) and Other (Customized In-Home Supports)

Survey Type: Routine

Dear Ms. Christina Martinez;

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,

Amanda Castañeda

Amanda Castañeda
Plan of Correction Coordinator
Quality Management Bureau/DHI

Q.16.1.DDW.D1556.3.RTN.09.16.4