Dear Mr. Johnson and Mr. Stack;

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:

**Compliance with all Conditions of Participation.**
This determination is based on your agency’s compliance with CMS waiver assurances at the Condition of Participation level. The attached QMB Report of Findings indicates Standard Level deficiencies identified and requires implementation of a Plan of Correction.

**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.

During the exit interview of your on-site survey Attachment A on the Plan of Correction Process was provided to you. Please refer to Attachment A for specific instruction on completing your Plan of Correction. At a minimum your Plan of Correction should address the following for each Tag cited:

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

**On-going Quality Assurance/Quality Improvement Processes:**
- What is going to be done? (i.e. file reviews, periodic check with checklist, etc.)
- How many individuals is this going to effect? (i.e. percentage of individuals reviewed, number of files reviewed, etc.)
- How often will this be completed? (i.e. weekly, monthly, quarterly, etc.)
- Who is responsible? (responsible position)
- What steps will be taken if issues are found? (i.e. retraining, requesting documents, filing RORI, etc.)

**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: Lisa Medina-Lujan  
HSD/OIG  
Program Integrity Unit  
2025 S. Pacheco Street  
Santa Fe, New Mexico 87505  

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

Attention: Lisa Medina-Lujan  
HSD/OIG  
Program Integrity Unit  
1474 Rodeo Road  
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,

Barbara Kane, BAS  
Barbara Kane, BAS  
Team Lead/Healthcare Surveyor  
Division of Health Improvement  
Quality Management Bureau
Survey Process Employed:

Entrance Conference Date: September 19, 2016

Present: Community Options, Inc.
- Hector Johnson, State Director
- Jessica Adamchak, RN
- Marsha Ford, Program Coordinator/Service Coordinator/Direct Support Personnel

DOH/DHI/QMB
- Barbara Kane, BAS, Team Lead/Healthcare Surveyor
- Chris Melon, MPA, Healthcare Surveyor
- Corrina Strain, BSN, RN, Healthcare Surveyor

Exit Conference Date: September 21, 2016

Present: Community Options, Inc.
- Hector Johnson, State Director
- Marsha Ford, Program Coordinator/Service Coordinator/Direct Support Personnel
- Ashley Hatfield, Incident Management Coordinator/Quality Assurance/Trainer

DOH/DHI/QMB
- Barbara Kane, BSA, Team Lead/Healthcare Surveyor
- Chris Melon, MPA, Healthcare Surveyor
- Corrina Strain, BSN, RN, Healthcare Surveyor

DDSD - SW Regional Office
- Scott Doan, Bureau Chief (via phone)

Administrative Locations Visited Number: 1

Total Sample Size Number: 7
- 1 - Jackson Class Member
- 6 - Non-Jackson Class Members
- 1 - Supported Living
- 1 - Adult Habilitation
- 1 - Supported Employment
- 5 - Customized Community Supports
- 3 - Community Integrated Employment Services

Total Homes Visited Number: 1
- Supported Living Homes Visited Number: 1

Persons Served Records Reviewed Number: 7

Persons Served Interviewed Number: 6

Persons Served Not Seen and/or Not Available Number: 1 (One Individual was not available during the on-site survey)
Direct Support Personnel Interviewed Number: 6
Direct Support Personnel Records Reviewed Number: 9 (Program Coordinator/Service Coordinator also performs duties as Direct Support Personnel)
Service Coordinator Records Reviewed Number: 1
Administrative Interviews Number: 2

Administrative Processes and Records Reviewed:

- Medicaid Billing/Reimbursement Records for all Services Provided
- Accreditation Records
- Individual Medical and Program Case Files, including, but not limited to:
  o Individual Service Plans
  o Progress on Identified Outcomes
  o Healthcare Plans
  o Medication Administration Records
  o Medical Emergency Response Plans
  o Therapy Evaluations and Plans
  o Healthcare Documentation Regarding Appointments and Required Follow-Up
  o 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.Casaneda@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.

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.

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

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

The Division of Health Improvement, Quality Management Bureau (QMB) surveys compliance of the Developmental Disabilities Waiver (DDW) standards and 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 the following Service Domains.

Case Management Services (Four Service Domains):
- Plan of Care: ISP Development & Monitoring
- Level of Care
- Qualified Providers
- Health, Safety and Welfare

Community Living Supports / Inclusion Supports (Three Service Domains):
- Service Plans: ISP Implementation
- Qualified Provider
- Health, Safety and Welfare

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
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: Plan of Care ISP Development & Monitoring**

Condition of Participation:

1. **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:

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

**Service Domain: Level of Care**

Condition of Participation:

3. **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.

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: Service Plan: ISP Implementation**

Condition of Participation:

5. **ISP Implementation**: Services provided shall be consistent with the components of the ISP and implemented to achieve desired outcomes / action step.

**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 and/or 6 or more total Condition level tags in the Report of Findings. 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.
### Agency:
Community Options, Inc. – Metro Region

### Program:
Developmental Disabilities Waiver

### Service:
- **2012:** Inclusion Supports (Customized Community Supports, Community Integrated Employment Services)
- **2007:** Community Living (Supported Living) and Community Inclusion (Adult Habilitation, Supported Employment)

### Monitoring Type:
Routine, Survey

### Survey Date:
September 16 – 21, 2016

### Standard of Care

#### Deficiencies

<table>
<thead>
<tr>
<th>Tag #</th>
<th>Agency Case File</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A08</td>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015 Chapter 5 (CIES) 3. Agency Requirements J. Consumer Records Policy: Community Integrated Employment 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 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. Additional documentation that is required to be maintained at the administrative office includes:</td>
</tr>
<tr>
<td></td>
<td>Based on record review, the Agency did not maintain a complete and confidential case file at the administrative office for 1 of 7 individuals. Review of the Agency individual case files revealed the following items were not found, incomplete, and/or not current:</td>
</tr>
<tr>
<td></td>
<td>- Annual ISP</td>
</tr>
<tr>
<td></td>
<td>° Not Current (#8) (ISP does not reflect a change of service. Community Integrated Employment Services were discontinued on 7/06/16.) (ISP effective date 7/30/2016 – 7/29/2017)</td>
</tr>
</tbody>
</table>

#### Agency Plan of Correction, On-going QA/QI and Responsible Party

Provider:
State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?):

Provider:
Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?):

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Survey Report #: Q.17.1.DDW.D3124.5.RTN.01.16.294

Page 13 of 56
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. 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.

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.
| Tag # 1A08.1 | Agency Case File - Progress Notes | Standard Level Deficiency | Provider: State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?): →

Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): → |

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**Chapter 5 (CIES) 3. Agency Requirements: 6. Reimbursement A. 1.** Provider Agencies must maintain all records necessary to fully disclose the service, quality...The documentation of the billable time spent with an individual shall be kept on the written or electronic record...

**Chapter 6 (CCS) 3. Agency Requirements: 4. Reimbursement A. Record Requirements 1.** Provider Agencies must maintain all records necessary to fully disclose the service, quality...The documentation of the billable time spent with an individual shall be kept on the written or electronic record...

**Chapter 7 (CIHS) 3. Agency Requirements: 4. Reimbursement A. 1.** Provider Agencies must maintain all records necessary to fully disclose the service, quality...The documentation of the billable time spent with an individual shall be kept on the written or electronic record...

**Chapter 11 (FL) 3. Agency Requirements: 4. Reimbursement A. 1.** Provider Agencies must maintain all records necessary to fully disclose the service, quality...The documentation of the billable time spent with an individual shall be kept on the written or electronic record...

**Chapter 12 (SL) 3. Agency Requirements: 2. Reimbursement A. 1.** Provider Agencies must maintain all records necessary to fully disclose the service, quality...The documentation of the billable time spent with an individual shall be kept on the written or electronic record...

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Based on record review, the Agency did not maintain progress notes and other service delivery documentation for 1 of 7 Individuals.

Review of the Agency individual case files revealed the following items were not found:

**Supported Employment Progress Notes/Daily Contact Logs**
- Individual #7 - None found for 6/01/2016; 7/09/2016.

Based on record review, the Agency did not maintain progress notes and other service delivery documentation for 1 of 7 Individuals.
Chapter 13 (IMLS) 3. Agency Requirements: 4. Reimbursement A. 1. …Provider Agencies must maintain all records necessary to fully disclose the service, quality…The documentation of the billable time spent with an individual shall be kept on the written or electronic record…

Chapter 15 (ANS) 4. Reimbursement A. 1. …Provider Agencies must maintain all records necessary to fully disclose the service, quality…The documentation of the billable time spent with an individual shall be kept on the written or electronic record…


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:

(3) Progress notes and other service delivery documentation;
**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 # 1A11.1</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>Transportation Training</strong></td>
<td>Based on record review, the Agency did not provide and/or have documentation for staff training regarding the safe operation of the vehicle, assisting passengers and safe lifting procedures for 1 of 9 Direct Support Personnel. No documented evidence was found of the following required training:</td>
<td><strong>Provider:</strong> State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?): →</td>
<td></td>
</tr>
<tr>
<td><strong>Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy</strong></td>
<td><strong>Provider:</strong> Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): →</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Training Requirements for Direct Service Agency Staff Policy Eff. Date:</strong> March 1, 2007 <strong>II. POLICY STATEMENTS:</strong></td>
<td>1. Operating a fire extinguisher 2. Proper lifting procedures 3. General vehicle safety precautions (e.g., pre-trip inspection, removing keys from the ignition when not in the driver’s seat) 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) 5. Operating wheelchair lifts (if applicable to the staff’s role) 6. Wheelchair tie-down procedures (if applicable to the staff’s role) 7. Emergency and evacuation procedures (e.g., roadside emergency, fire emergency) <strong>NMAC 7.9.2 F. TRANSPORTATION:</strong> (1) Any employee or agent of a regulated facility or agency who is responsible for assisting a resident in boarding or alighting from a motor vehicle must provide and/or have documentation regarding the safe operation of the vehicle, assisting passengers and safe lifting procedures.</td>
<td></td>
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</tr>
</tbody>
</table>


Survey Report #: Q.17.1.DDW.D3124.5.RTN.01.16.294
vehicle must complete a state-approved training program in passenger transportation assistance before assisting any resident. The passenger transportation assistance program shall be comprised of but not limited to the following elements: resident assessment, emergency procedures, supervised practice in the safe operation of equipment, familiarity with state regulations governing the transportation of persons with disabilities, and a method for determining and documenting successful completion of the course. The course requirements above are examples and may be modified as needed.

(2) Any employee or agent of a regulated facility or agency who drives a motor vehicle provided by the facility or agency for use in the transportation of clients must complete:

(a) A state approved training program in passenger assistance and

(b) A state approved training program in the operation of a motor vehicle to transport clients of a regulated facility or agency. The motor vehicle transportation assistance program shall be comprised of but not limited to the following elements: resident assessment, emergency procedures, supervised practice in the safe operation of motor vehicles, familiarity with state regulations governing the transportation of persons with disabilities, maintenance and safety record keeping, training on hazardous driving conditions and a method for determining and documenting successful completion of the course. The course requirements above are examples and may be modified as needed.

(c) A valid New Mexico driver’s license for the type of vehicle being operated consistent with State of New Mexico requirements.

(3) Each regulated facility and agency shall establish and enforce written policies (including training) and procedures for employees who provide assistance to clients with boarding or
alighting from motor vehicles.

(4) Each regulated facility and agency shall establish and enforce written polices (including training and procedures for employees who operate motor vehicles to transport clients.


**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.

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

**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;

**CHAPTER 7 (CIHS) 3. Agency Requirements**

**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 DDSD Policy T-003: Training Requirements for Direct Service Agency Staff Policy

**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. 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 #</th>
<th>A20</th>
<th>Direct Support Personnel Training</th>
<th>Standard Level Deficiency</th>
<th>Provider:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Based on record review, the Agency did not ensure Orientation and Training requirements were met for 1 of 9 Direct Support Personnel. Review of Direct Support Personnel training records found no evidence of the following required DOH/DDSD trainings and certification being completed:</td>
<td>State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?): →</td>
</tr>
</tbody>
</table>

- Assisting with Medication Delivery (DSP #206)

Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): →

<table>
<thead>
<tr>
<th>Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy - Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007</th>
<th>II. POLICY STATEMENTS:</th>
</tr>
</thead>
<tbody>
<tr>
<td>A. Individuals shall receive services from competent and qualified staff.</td>
<td>A. Individuals shall receive services from competent and qualified staff.</td>
</tr>
<tr>
<td>B. Staff shall complete individual-specific (formerly known as &quot;Addendum B&quot;) training requirements in accordance with the specifications described in the individual service plan (ISP) of each individual served.</td>
<td>B. Staff shall complete individual-specific (formerly known as &quot;Addendum B&quot;) training requirements in accordance with the specifications described in the individual service plan (ISP) of each individual served.</td>
</tr>
<tr>
<td>C. Staff shall complete training on DOH-approved incident reporting procedures in accordance with 7 NMAC 1.13.</td>
<td>C. Staff shall complete training on DOH-approved incident reporting procedures in accordance with 7 NMAC 1.13.</td>
</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>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>
</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>E. Staff providing direct services shall maintain certification in first aid and CPR. The training materials shall meet OSHA requirements/guidelines.</td>
</tr>
<tr>
<td>F. Staff who may be exposed to hazardous chemicals shall complete relevant training in accordance with OSHA requirements.</td>
<td>F. Staff who may be exposed to hazardous chemicals shall complete relevant training in accordance with OSHA requirements.</td>
</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 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.</td>
<td>G. Staff shall be certified in a DDSD-approved behavioral intervention system (e.g., Mandt, CPI) before using physical restraint techniques. Staff 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.</td>
</tr>
<tr>
<td>H. Staff shall complete and maintain certification in a DDSD-approved medication course in accordance with the DDSD Medication Delivery Policy M-001.</td>
<td>H. Staff shall complete and maintain certification in a DDSD-approved medication course in accordance with the DDSD Medication Delivery Policy M-001.</td>
</tr>
<tr>
<td>I. Staff providing direct services shall complete</td>
<td>I. Staff providing direct services shall complete</td>
</tr>
</tbody>
</table>
safety training within the first thirty (30) days of employment and before working alone with an individual receiving service.


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.

CHAPTER 6 (CCS) 3. Agency Requirements
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;

CHAPTER 7 (CIHS) 3. Agency Requirements
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 DDSD Policy T-003: Training Requirements for Direct Service Agency Staff Policy

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. 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: A. Individuals shall receive services from competent and qualified staff. B. Staff shall complete individual specific (formerly known as &quot;Addendum B&quot;) training requirements in accordance with the specifications described in the individual service plan (ISP) for each individual serviced. Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015 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>Based on interview, the Agency did not ensure training competencies were met for 2 of 6 Direct Support Personnel. When DSP were asked if the Individual had a CARMP and if so, what the plan covered, the following was reported: • DSP #207 stated, &quot;No.&quot; According to the Individual Specific Training Section of the ISP, the Individual requires a CARMP. (Individual #4)</td>
<td>State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?): →</td>
<td></td>
</tr>
<tr>
<td>CHAPTER 6 (CCS) 3. Agency Requirements 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>When DSP were asked if the Individual had Health Care Plans and if so, what the plan(s) covered, the following was reported: • DSP #204 stated, &quot;No.&quot; According to the Electronic Comprehensive Health Assessment Tool, the Individual requires a Health Care Plan for Seizures. (Individual #2)</td>
<td>→</td>
<td></td>
</tr>
<tr>
<td>CHAPTER 7 (CIHS) 3. Agency Requirements C. Training Requirements: The Provider Agency must report required personnel training status to the DDSD Statewide Training</td>
<td>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): →</td>
<td></td>
<td></td>
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</tbody>
</table>
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.

<table>
<thead>
<tr>
<th>CHAPTER 12 (SL) 3. Agency Requirements</th>
</tr>
</thead>
<tbody>
<tr>
<td>B. Living Supports- Supported Living</td>
</tr>
<tr>
<td>Services Provider Agency Staffing</td>
</tr>
<tr>
<td>Requirements: 3. Training:</td>
</tr>
<tr>
<td>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.</td>
</tr>
<tr>
<td>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,</td>
</tr>
</tbody>
</table>
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;
Service Domain: Health and Welfare – The state, on an ongoing basis, identifies, addresses and seeks to prevent occurrences of abuse, neglect and exploitation. Individuals shall be afforded their basic human rights. The provider supports individuals to access needed healthcare services in a timely manner.

Tag #1A08.2 Healthcare Requirements

<table>
<thead>
<tr>
<th>Standard of Care</th>
<th>Deficiencies</th>
<th>Agency Plan of Correction, On-going QA/QI and Responsible Party</th>
<th>Date Due</th>
</tr>
</thead>
<tbody>
<tr>
<td>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.</td>
<td>Based on record review, the Agency did not provide documentation of annual physical examinations and/or other examinations as specified by a licensed physician for 2 of 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:</td>
<td>Provider: State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?): →</td>
<td></td>
</tr>
<tr>
<td>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.</td>
<td><strong>Community Inclusion Services / Other Services Healthcare Requirements (Individuals Receiving Inclusion / Other Services Only):</strong></td>
<td><strong>Provider:</strong> Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): →</td>
<td></td>
</tr>
<tr>
<td>DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION (DDSD): Director’s Release: Consumer Record Requirements eff. 11/1/2012 III. Requirement Amendments(s) or Clarifications:</td>
<td><strong>Auditory Exam</strong>  ◦ Individual #3 - As indicated by collateral documentation reviewed, exam was completed on 5/8/2014. Next exam was to be completed in 5/2016. No evidence of</td>
<td><strong>Provider:</strong> Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): →</td>
<td></td>
</tr>
<tr>
<td>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.</td>
<td>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; 6/15/2015</td>
<td></td>
<td></td>
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</tbody>
</table>
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.

Chapter 13 (IMLS) 2. Service Requirements:

- Blood Levels
  - Individual #8 - As indicated by collateral documentation reviewed, lab work was ordered on 12/17/2015. No evidence of lab results was found.

- Nutritional Evaluation
  - Individual #8 - As indicated by collateral documentation reviewed, Evaluation was completed on 10/14/2015. Follow-up was to be completed in 6 months. No evidence of follow-up found.
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 provider. 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 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
(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).
**Tag # 1A09**  
**Medication Delivery**  
**Routine Medication Administration**

**NMAC 16.19.11.8 MINIMUM STANDARDS:**  
A. MINIMUM STANDARDS FOR THE DISTRIBUTION, STORAGE, HANDLING AND RECORD KEEPING OF DRUGS:  
(d) The facility shall have a Medication Administration Record (MAR) documenting medication administered to residents, **including over-the-counter medications.**

This documentation shall include:  
(i) Name of resident;  
(ii) Date given;  
(iii) Drug product name;  
(iv) Dosage and form;  
(v) Strength of drug;  
(vi) Route of administration;  
(vii) How often medication is to be taken;  
(viii) Time taken and staff initials;  
(ix) Dates when the medication is discontinued or changed;  
(x) The name and initials of all staff administering medications.

**Model Custodial Procedure Manual**  
**D. Administration of Drugs**

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

All PRN (As needed) medications shall have complete detail instructions regarding the administering of the medication. This shall include:  
- symptoms that indicate the use of the medication,  
- exact dosage to be used, and  
- the exact amount to be used in a 24-hour period.

| Standard Level Deficiency | Provider:  
State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?): → |
|---------------------------|-----------------------------------------------------------|
| Medication Administration Records (MAR) were reviewed for the month of September 2016.  
Based on record review, 1 of 2 individuals had Medication Administration Records (MAR), which contained missing medications entries and/or other errors:  
Individual #7  
September 2016  
Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries:  
- Melatonin 5 mg 2 tablets (1 time daily) – Blank 9/18 (8:00 PM)  
- Oxybutynin Chloride 5 mg 1 tablet (3 times daily) – Blank 9/16 (3:00 PM)  
- Vitamin C 250 mg 1 tablet (2 times daily) – Blank 9/16 (4:00 PM)  
- Vitamin D3 400 units 2 tablets (1 time daily) – Blank 9/2 (8:00 AM) | Provider:  
Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): → |

**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:**

- 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):

- 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

<table>
<thead>
<tr>
<th>I. Healthcare Requirements for Family Living.</th>
</tr>
</thead>
<tbody>
<tr>
<td>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.</td>
</tr>
<tr>
<td>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.</td>
</tr>
<tr>
<td>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;</td>
</tr>
<tr>
<td>b. When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) must be maintained and include:</td>
</tr>
<tr>
<td>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;</td>
</tr>
<tr>
<td>ii. Prescribed dosage, frequency and method/route of administration, times and dates of administration;</td>
</tr>
<tr>
<td>iii. Initials of the individual administering or assisting with the medication delivery;</td>
</tr>
</tbody>
</table>
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 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.

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 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 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

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.

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 # 1A31</th>
<th>Standard Level Deficiency</th>
<th>Provider:</th>
</tr>
</thead>
</table>
| Client Rights/Human Rights | 7.26.3.11 RESTRICTIONS OR LIMITATION OF CLIENT’S RIGHTS:  
A. A service provider shall not restrict or limit a client's rights except:  
(1) where the restriction or limitation is allowed in an emergency and is necessary to prevent imminent risk of physical harm to the client or another person; or  
(2) where the interdisciplinary team has determined that the client's limited capacity to exercise the right threatens his or her physical safety; or  
(3) as provided for in Section 10.1.14 [now Subsection N of 7.26.3.10 NMAC].  
B. Any emergency intervention to prevent physical harm shall be reasonable to prevent harm, shall be the least restrictive intervention necessary to meet the emergency, shall be allowed no longer than necessary and shall be subject to interdisciplinary team (IDT) review. The IDT upon completion of its review may refer its findings to the office of quality assurance. The emergency intervention may be subject to review by the service provider’s behavioral support committee or human rights committee in accordance with the behavioral support policies or other department regulation or policy.  
C. The service provider may adopt reasonable program policies of general applicability to clients served by that service provider that do not violate client rights. [09/12/94; 01/15/97; Recompiled 10/31/01]  
Long Term Services Division  
Policy Title: Human Rights Committee  
Requirements Eff Date: March 1, 2003  
IV. POLICY STATEMENT - Human Rights | State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?): →  
Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): → |
Committees are required for residential service provider agencies. The purpose of these committees with respect to the provision of Behavior Supports is to review and monitor the implementation of certain Behavior Support Plans.

Human Rights Committees may not approve any of the interventions specifically prohibited in the following policies:
- Aversive Intervention Prohibitions
- Psychotropic Medications Use
- Behavioral Support Service Provision.

A Human Rights Committee may also serve other agency functions as appropriate, such as the review of internal policies on sexuality and incident management follow-up.

A. HUMAN RIGHTS COMMITTEE ROLE IN BEHAVIOR SUPPORTS
Only those Behavior Support Plans with an aversive intervention included as part of the plan or associated Crisis Intervention Plan need to be reviewed prior to implementation. Plans not containing aversive interventions do not require Human Rights Committee review or approval.

2. The Human Rights Committee will determine and adopt a written policy stating the frequency and purpose of meetings. Behavior Support Plans approved by the Human Rights Committee will be reviewed at least quarterly.

3. Records, including minutes of all meetings will be retained at the agency with primary responsibility for implementation for at least five years from the completion of each individual’s Individual Service Plan.
| Disabilities Supports Division (DDSD) - Procedure Title: Medication Assessment and Delivery  
  | Procedure Eff Date: November 1, 2006  
  | B. 1. e. If the PRN medication is to be used in response to psychiatric and/or behavioral symptoms in addition to the above requirements, obtain current written consent from the individual, guardian or surrogate health decision maker and submit for review by the agency's Human Rights Committee (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). |
### 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.

<table>
<thead>
<tr>
<th>Tag # IS25 / 5I25 Community Integrated Employment Services / Supported Employment Reimbursement</th>
<th>Standard of Care</th>
<th>Deficiencies</th>
<th>Agency Plan of Correction, On-going QA/QI and Responsible Party</th>
<th>Date Due</th>
</tr>
</thead>
<tbody>
<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015</td>
<td>CHAPTER 5 (CIES) 6. REIMBURSEMENT: A. 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:</td>
<td>Based on record review, the Agency did not provide written or electronic documentation as evidence for each unit billed for Supported Employment Services for 1 of 4 individuals</td>
<td>Provider: State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?):</td>
<td></td>
</tr>
<tr>
<td>Individual #7</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>June 2016</td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>• The Agency billed 1 unit of Supported Employment (T2013) on 6/1/2016. No documentation was found for 6/1/2016 to justify the 1 unit billed.</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>July 2016</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>• The Agency billed 1 unit of Supported Employment (T2013) on 7/9/2016. No documentation was found for 7/9/2016 to justify the 1 unit billed.</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Provider:**

State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?):

**Provider:**

Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?):

**Provider:**


provided to the Medicaid recipient. Services that have been billed to Medicaid, but are not substantiated in a treatment plan and/or patient records for the recipient are subject to recoupment.
<table>
<thead>
<tr>
<th>Tag # 5l44</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Adult Habilitation Reimbursement</strong></td>
<td>Based on record review, the Agency did not provide written or electronic documentation as evidence for each unit billed for Adult Habilitation Services for 1 of 1 individual.</td>
</tr>
<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</td>
<td>Individual #7</td>
</tr>
<tr>
<td><strong>CHAPTER 1 III. PROVIDER AGENCY DOCUMENTATION OF SERVICE DELIVERY AND LOCATION</strong></td>
<td>July 2016</td>
</tr>
<tr>
<td>A. <strong>General:</strong> All Provider Agencies shall 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 date, time, individual name, servicing Provider Agency, level of services, and length of a session of service billed.</td>
<td>• The Agency billed 103 units of Adult Habilitation (T2021 U1) from 7/18/2016 through 7/22/2016. Documentation received accounted for 99 units.</td>
</tr>
<tr>
<td><strong>B. Billable Units:</strong> 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:</td>
<td><strong>Provider:</strong></td>
</tr>
<tr>
<td>(1) Date, start and end time of each service encounter or other billable service interval;</td>
<td>State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?): →</td>
</tr>
<tr>
<td>(2) A description of what occurred during the encounter or service interval; and</td>
<td><strong>Provider:</strong></td>
</tr>
<tr>
<td>(3) The signature or authenticated name of staff providing the service.</td>
<td>Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): →</td>
</tr>
</tbody>
</table>

**MAD-MR: 03-59 Eff 1/1/2004**

**8.314.1 BI RECORD KEEPING AND DOCUMENTATION REQUIREMENTS:**

Providers must maintain all records necessary to fully disclose the extent of the services provided to the Medicaid recipient. Services that have been billed to Medicaid, but are not substantiated in a treatment plan and/or patient records for the recipient are subject to recoupment.
### Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007

#### CHAPTER 5 XVI. REIMBURSEMENT

**A. Billable Unit.** A billable unit for Adult Habilitation Services is in 15-minute increments hour. The rate is based on the individual's level of care.

**B. Billable Activities**

1. The Community Inclusion Provider Agency can bill for those activities listed and described on the ISP and within the Scope of Service. Partial units are allowable. Billable units are face-to-face, except that Adult Habilitation services may be non-face-to-face under the following conditions: (a) Time that is non-face-to-face is documented separately and clearly identified as to the nature of the activity; and (b) Non face-to-face hours do not exceed 5% of the monthly billable hours.

2. Adult Habilitation Services can be provided with any other services, insofar as the services are not reported for the same hours on the same day, except that Therapy Services and Case Management may be provided and billed for the same hours.
| Tag # IS30 | Customized Community Supports Reimbursement |
|-----------------------------------------------|
| **Standard Level Deficiency** |
| Based on record review, the Agency did not provide written or electronic documentation as evidence for each unit billed for Customized Community Supports for 2 of 5 individuals. |
| **Provider:** |
| State your Plan of Correction for the deficiencies cited in this tag here (How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?): |
| Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): |

<p>| | | |</p>
<table>
<thead>
<tr>
<th></th>
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</tr>
</thead>
<tbody>
<tr>
<td><strong>Individual #2</strong></td>
<td><strong>July 2016</strong></td>
<td></td>
</tr>
<tr>
<td>The Agency billed 41 units of Customized Community Supports (Individual) (H2021 HB U1) from 7/5/2016 through 7/8/2016. Documentation received accounted for 14 units.</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Individual #8</strong></td>
<td><strong>August 2016</strong></td>
<td></td>
</tr>
<tr>
<td>The Agency billed 117 units of Customized Community Supports (Group) (T2021 HB U7) from 8/15/2016 through 8/19/2016. Documentation received accounted for 108 units.</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>The Agency billed 111 units of Customized Community Supports (Group) (T2021 HB U7) from 8/22/2016 through 8/26/2016. Documentation received accounted for 109 units.</td>
<td></td>
</tr>
</tbody>
</table>

**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.

2. **Billable Unit:**
   1. The billable unit for Individual Customized Community Supports is a fifteen (15) minute unit.

   2. The billable unit for Community Inclusion Aide is a fifteen (15) minute unit.
3. The billable unit for Group Customized Community Supports is a fifteen (15) minute unit, with the rate category based on the NM DDW group.

4. The time at home is intermittent or brief; e.g. one-hour time period for lunch and/or change of clothes. The Provider Agency may bill for providing this support under Customized Community Supports without prior approval from DDSD.

5. The billable unit for Intensive Behavioral Customized Community Supports is a fifteen (15) minute unit. (There is a separate rate established for individuals who require one-to-one (1:1) support either in the community or in a group day setting due to behavioral challenges (NM DDW group G).

6. The billable unit for Fiscal Management for Adult Education is dollars charged for each class including a 10% administrative processing fee.

C. **Billable Activities:**

1. All DSP activities that are:
   a. Provided face to face with the individual;
   b. Described in the individual’s approved ISP;
   c. Provided in accordance with the Scope of Services; and
   d. Activities included in billable services, activities or situations.

2. Purchase of tuition, fees, and/or related materials associated with adult education opportunities as related to the ISP Action
| Plan and Outcomes, not to exceed $550 including administrative processing fee. |

3. Customized Community Supports can be included in ISP and budget with any other services.

MAD-MR: 03-59 Eff 1/1/2004
8.314.1 BI RECORD KEEPING AND DOCUMENTATION REQUIREMENTS:
Providers must maintain all records necessary to fully disclose the extent of the services provided to the Medicaid recipient. Services that have been billed to Medicaid, but are not substantiated in a treatment plan and/or patient records for the recipient are subject to recoupment.
Date: January 5, 2017

To: Hector Johnson, State Director
Provider: Community Options, Inc.
Address: 2720 San Pedro NE
State/Zip: Albuquerque, New Mexico 87110

E-mail Address: hector.johnson@comop.org

CC: Robert Stack, President and CEO
Address: 16 Farber Road
State/Zip: Princeton, New Jersey 08540

E-Mail Address: robert.stack@comop.org

Region: Metro
Survey Date: September 16 – 21, 2016
Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: 2012: Inclusion Supports (Customized Community Supports, Community Integrated Employment Services). 2007: Community Living (Supported Living) and Community Inclusion (Adult Habilitation, Supported Employment)

Survey Type: Routine

Dear Mr. Johnson and Mr. Stack;

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.17.1.DDW.D3124.5.RTN.09.17.005