

Date:	October 20, 2016
To: Provider: Address: State/Zip:	Hector Johnson, State Director Community Options, Inc. 2720 San Pedro NE Albuquerque, New Mexico 87110
E-mail Address:	hector.johnson@comop.org
CC: Address: State/Zip:	Robert Stack, President and CEO 16 Farber Road Princeton, New Jersey 08540
E-Mail Address	robert.stack@comop.org
Region: Survey Date: Program Surveyed:	Metro September 16 – 21, 2016 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
Team Leader:	Barbara Kane, BAS, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau
Team Members:	Chris Melon, MPA, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau and Corrina Strain, BSN, RN, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

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.

COUNTED HEALTH DECARDING

DIVISION OF HEALTH IMPROVEMENT 5301 Central Avenue NE, Suite 400 • Albuquerque, New Mexico • 87108 (505) 222-8623 • FAX: (505) 222-8661 • <u>http://www.dhi.health.state.nm.us</u>

QMB Report of Findings - Community Option, Inc. - Metro Region - September 16 - 21, 2016

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 i locess Employed.		
Entrance Conference Date:	September 19,	2016
Present:	Jessica Adamo	n, State Director
	Chris Melon, M	<u>3</u> BAS, Team Lead/Healthcare Surveyor IPA, Healthcare Surveyor BSN, RN, Healthcare Surveyor
Exit Conference Date:	September 21,	2016
Present:	Marsha Ford, F Personnel	n, State Director Program Coordinator/Service Coordinator/Direct Support I, Incident Management Coordinator/Quality
	Chris Melon, M	<u>3</u> BSA, Team Lead/Healthcare Surveyor IPA, Healthcare Surveyor BSN, RN, Healthcare Surveyor
	DDSD - SW Re Scott Doan, Bu	egional Office Ireau Chief (via phone)
Administrative Locations Visited	Number:	1
Total Sample Size	Number:	7
		1 - <i>Jackson</i> Class Member 6 - Non- <i>Jackson</i> Class Members
		 Supported Living Adult Habilitation Supported Employment Customized Community Supports 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

1 (One Individual was not available during the on-site survey)

Survey Process Employed:

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:
 - Individual Service Plans
 - Progress on Identified Outcomes
 - Healthcare Plans
 - Medication Administration Records
 - Medical Emergency Response Plans
 - Therapy Evaluations and Plans
 - Healthcare Documentation Regarding Appointments and Required Follow-Up
 - 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 <u>AmandaE.Castaneda@state.nm.us</u>. 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: <u>Instruction or in-service of staff alone may not be a sufficient plan of correction.</u> 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 <u>AmandaE.Castaneda@state.nm.us</u> 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 <u>AmandaE.Castaneda@state.nm.us</u> (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.
- 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 <u>must be annotated</u>; 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.
- 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.

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 <u>repeat</u> 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 <u>repeat</u> 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.

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 <u>within 10 business days</u> 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: <u>http://dhi.health.state.nm.us/qmb</u>
- 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 <u>Crystal.Lopez-Beck@state.nm.us</u> 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-toface 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)
	Employment)
Monitoring Type:	Routine, Survey
Survey Date:	September 16 – 21, 2016

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Date Due			
	Service Domain: Service Plans: ISP Implementation – Services are delivered in accordance with the service plan, including type, scope, amount, duration and frequency specified in the service plan.					
Tag # 1A08 Agency Case File	Standard Level Deficiency					
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. Provider agency case files for individuals are required to comply with the DDSD 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: 1. Vocational Assessments (if applicable) that are of quality and contain content acceptable to DVR and DDSD.	 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: Annual ISP 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) 	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?): →				

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	Standard Level Deficiency		
Agency Case File - Progress Notes			
Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015 Chapter 5 (CIES) 3. Agency Requirements: 6. Reimbursement A. 1. Provider Agencies must maintain all records necessary to fully disclose the service, qualityThe documentation of the billable time spent with an individual shall be kept on the written or electronic record	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;	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?): →	
Chapter 6 (CCS) 3. Agency Requirements: 4. Reimbursement A. Record Requirements 1. Provider Agencies must maintain all records necessary to fully disclose the service, qualityThe 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. 1Provider Agencies must maintain all records necessary to fully disclose the service, qualityThe documentation of the billable time spent with an individual shall be kept on the written or electronic record	7/09/2016.	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 11 (FL) 3. Agency Requirements: 4. Reimbursement A. 1Provider Agencies must maintain all records necessary to fully disclose the service, qualityThe 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, qualityThe documentation of the billable time spent with an individual shall be kept on the written or electronic record			

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Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Date Due
		ified providers to assure adherence to waive rovider training is conducted in accordance	
Tag # 1A11.1 Transportation Training	Standard Level Deficiency		
 Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy Training Requirements for Direct Service Agency Staff Policy Eff. Date: March 1, 2007 II. POLICY STATEMENTS: Staff providing direct services shall complete safety training within the first thirty (30) days of employment and before working alone with an individual receiving services. The training shall address at least the following: Operating a fire extinguisher Proper lifting procedures General vehicle safety precautions (e.g., pre-trip inspection, removing keys from the ignition when not in the driver's seat) 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) Operating wheelchair lifts (if applicable to the staff's role) Wheelchair tie-down procedures (if applicable to the staff's role) Emergency and evacuation procedures (e.g., roadside emergency, fire emergency) 	 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: Transportation (DSP #202) 	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?): →	

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 polices (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.	
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.	
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;	
Agency Starr Folicy,	
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;	
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Tag # 1A20	Standard Level Deficiency		
Direct Support Personnel TrainingDepartment of Health (DOH) DevelopmentalDisabilities Supports Division (DDSD) Policy- Policy Title: Training Requirements forDirect Service Agency Staff Policy - Eff.March 1, 2007 - II. POLICY STATEMENTS:A. Individuals shall receive services fromcompetent and qualified staff.B. Staff shall complete individual-specific(formerly known as "Addendum B") training	Based on record review, the Agency did not ensure Orientation and Training requirements were met for 1 of 9 Direct Support Personnel training records found no evidence of the following required DOH/DDSD trainings and certification being completed: • Assisting with Medication Delivery (DSP #206)	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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safety training within the first thirty (30) days of	
employment and before working alone with an	
individual receiving service.	
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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.	
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	
Direct Service Agency Stall Fully	
CUADTED 11 (EL) 2 Agaman Daminamenta	
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	
	1

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	
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;		

Detak and an antification the DDOD Daliau T		
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 served, and 4. Start that assists the individual with medication (e.g., setting up		
medication, or reminders) must have completed		
Assisting with Medication Delivery (AWMD)		
Training.		
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		
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and conducted, including training on the	
Individual Service Plan outcomes, actions steps	
and strategies and associated support plans	
(e.g. health care plans, MERP, PBSP and BCIP	
etc), information about the individual's	
preferences with regard to privacy,	
communication style, and routines. Individual	
specific training for therapy related WDSI,	
Healthcare Plans, MERPs, CARMP, PBSP, and	
BCIP must occur at least annually and more	
often if plans change or if monitoring finds	
incorrect implementation. Family Living	
providers must notify the relevant support plan	
author whenever a new DSP is assigned to work	
with an individual, and therefore needs to	
receive training, or when an existing DSP	
requires a refresher. The individual should be	
present for and involved in individual specific	
training whenever possible.	
CHAPTER 12 (SL) 3. Agency Requirements	
B. Living Supports- Supported Living	
Services Provider Agency Staffing	
Requirements: 3. Training:	
A. All Living Supports- Supported Living	
Provider Agencies must ensure staff training in	
accordance with the DDSD Policy T-003: for	
Training Requirements for Direct Service	
Agency Staff. Pursuant to CMS requirements,	
the services that a provider renders may only be	
claimed for federal match if the provider has	
completed all necessary training required by the	
state. All Supported Living provider agencies	
must report required personnel training status to	
the DDSD Statewide Training Database as	
specified in DDSD Policy T-001: Reporting and	
Documentation for DDSD Training	
Requirements.	
B Individual specific training must be arranged	
and conducted, including training on the ISP	
Outcomes, actions steps and strategies,	
associated support plans (e.g. health care plans,	

MERP, PBSP and BCIP, etc), and information		
about the individual's preferences with regard to		
privacy, communication style, and routines.		
Individual specific training for therapy related		
WDSI, Healthcare Plans, MERP, CARMP,		
PBSP, and BCIP must occur at least annually		
and more often if plans change or if monitoring		
finds incorrect implementation. Supported		
Living providers must notify the relevant support		
plan author whenever a new DSP is assigned to		
work with an individual, and therefore needs to		
receive training, or when an existing DSP		
requires a refresher. The individual should be present for and involved in individual specific		
training whenever possible.		
CHAPTER 13 (IMLS) R. 2. Service		
Requirements. Staff Qualifications 2. DSP		
Qualifications. E. Complete training		
requirements as specified in the DDSD Policy T-		
003: Training Requirements for Direct Service		
Agency Staff - effective March 1, 2007. Report		
required personnel training status to the DDSD		
Statewide Training Database as specified in the		
DDSD Policy T-001: Reporting and		
Documentation of DDSD Training Requirements		
Policy;		

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Date Due
	The state, on an ongoing basis, identifies, a	•	
	als shall be afforded their basic human righ	ts. The provider supports individuals to ac	cess
needed healthcare services in a timely ma			
Tag #1A08.2 Healthcare Requirements	Standard Level Deficiency		
NMAC 8.302.1.17 RECORD KEEPING AND	Based on record review, the Agency did not	Provider:	
DOCUMENTATION REQUIREMENTS: A	provide documentation of annual physical	State your Plan of Correction for the	
provider must maintain all the records	examinations and/or other examinations as	deficiencies cited in this tag here (How is the	
necessary to fully disclose the nature, quality,	specified by a licensed physician for 2 of 7	deficiency going to be corrected? This can be specific to each deficiency cited or if possible an	
amount and medical necessity of services	individuals receiving Community Inclusion,	overall correction?): \rightarrow	
furnished to an eligible recipient who is	Living Services and Other Services.		
currently receiving or who has received services in the past.	Review of the administrative individual case files		
Services in the past.	revealed the following items were not found,		
B. Documentation of test results: Results of	incomplete, and/or not current:		
tests and services must be documented, which			
includes results of laboratory and radiology	Community Inclusion Services / Other		
procedures or progress following therapy or	Services Healthcare Requirements		
treatment.	(Individuals Receiving Inclusion / Other	Provider:	
	Services Only):	Enter your ongoing Quality	
DEVELOPMENTAL DISABILITIES SUPPORTS		Assurance/Quality Improvement processes	
DIVISION (DDSD): Director's Release:	Dental Exam	as it related to this tag number here (What is	
Consumer Record Requirements eff. 11/1/2012	 Individual #8 - As indicated by collateral 	going to be done? How many individuals is this	
III. Requirement Amendments(s) or	documentation reviewed, exam was	going to effect? How often will this be completed? Who is responsible? What steps will be taken if	
Clarifications:	completed on 8/26/2013. Follow-up was to	issues are found?): \rightarrow	
A. All case management, living supports,	be completed in 3 years. No evidence of		
customized in-home supports, community	follow-up found.		
integrated employment and customized			
community supports providers must maintain	Vision Exam		
records for individuals served through DD Waiver	 Individual #3 - As indicated by the DDSD file 		
in accordance with the Individual Case File Matrix	matrix Vision Exams are to be conducted		
incorporated in this director's release.	every other year. No evidence of exam was		
	found.		
H. Readily accessible electronic records are accessible, including those stored through the			
Therap web-based system.	Auditory Exam		
inerap web-based system.	 Individual #3 - As indicated by collateral 		
Developmental Disabilities (DD) Waiver Service	documentation reviewed, exam was		
Standards effective 11/1/2012 revised 4/23/2013;	completed on 5/8/2014. Next exam was to		
6/15/2015	be completed in 5/2016. No evidence of		

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.	 exam was found. 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. 	
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.	 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. 	
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)		
lionis)		
Developmental Dischilition (DD) Waisen Comise		
Developmental Disabilities (DD) Waiver Service		
Standards effective 4/1/2007		
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		
i or manadalo who are nowly anotated to the		I

DD Waiver program, the HAT may be	
completed within 2 weeks following the initial	
ISP meeting and submitted with any strategies	
and support plans indicated in the ISP, or	
within 72 hours following admission into direct	
services, whichever comes first.	
(2) Each individual will have a Health Care	
Coordinator, designated by the IDT. When the	
individual's HAT score is 4, 5 or 6 the Health	
Care Coordinator shall be an IDT member,	
other than the individual. The Health Care	
Coordinator shall oversee and monitor health	
care services for the individual in accordance	
with these standards. In circumstances where	
no IDT member voluntarily accepts designation	
as the health care coordinator, the community	
living provider shall assign a staff member to	
this role.	
(3) For each individual receiving Community	
Living Services, the provider agency shall	
ensure and document the following:	
(a)Provision of health care oversight	
consistent with these Standards as	
detailed in Chapter One section III E:	
Healthcare Documentation by Nurses for	
Community Living Services, Community	
Inclusion Services and Private Duty	
Nursing Services.	
b) That each individual with a score of 4, 5,	
or 6 on the HAT, has a Health Care Plan	
developed by a licensed nurse.	
(c)That an individual with chronic	
condition(s) with the potential to	
exacerbate into a life threatening	
condition, has Crisis Prevention/	
Intervention Plan(s) developed by a	
licensed nurse or other appropriate	
professional for each such condition.	
(4) That an average of 3 hours of documented	
nutritional counseling is available annually, if	
recommended by the IDT.	
(5) That the physical property and grounds are	

free of hazards to the individual's health and		
safety.		
(6) In addition, for each individual receiving		
Supported Living or Family Living Services, the		
provider shall verify and document the		
following:		
(a)The individual has a primary licensed		
physician;		
(b)The individual receives an annual		
physical examination and other		
examinations as specified by a licensed		
physician;		
(c)The individual receives annual dental		
check-ups and other check-ups as		
specified by a licensed dentist;		
(d)The individual receives eye examinations		
as specified by a licensed optometrist or		
ophthalmologist; and		
(e)Agency activities that occur as follow-up		
to medical appointments (e.g. treatment,		
visits to specialists, changes in		
medication or daily routine).		
medication of daily routine).		

Tag # 1A09	Standard Level Deficiency		
Medication Delivery			
Routine Medication Administration			
NMAC 16.19.11.8 MINIMUM STANDARDS:	Medication Administration Records (MAR) were	Provider:	
A. MINIMUM STANDARDS FOR THE	reviewed for the month of September 2016.	State your Plan of Correction for the	
DISTRIBUTION, STORAGE, HANDLING AND		deficiencies cited in this tag here (How is the	
RECORD KEEPING OF DRUGS:	Based on record review, 1 of 2 individuals had	deficiency going to be corrected? This can be	
(d) The facility shall have a Medication	Medication Administration Records (MAR),	specific to each deficiency cited or if possible an	
Administration Record (MAR) documenting	which contained missing medications entries	overall correction?): \rightarrow	
medication administered to residents,	and/or other errors:		
including over-the-counter medications.			
This documentation shall include:	Individual #7		
(i) Name of resident;	September 2016		
(ii) Date given;	Medication Administration Records contained		
(iii) Drug product name;	missing entries. No documentation found		
(iv) Dosage and form;	indicating reason for missing entries:	Provider:	
(v) Strength of drug;		Enter your ongoing Quality	
(vi) Route of administration;	Melatonin 5 mg 2 tablets (1 time daily) –	Assurance/Quality Improvement processes	
(vii) How often medication is to be taken;	Blank 9/18 (8:00 PM)	as it related to this tag number here (What is	
(viii) Time taken and staff initials;		going to be done? How many individuals is this	
(ix) Dates when the medication is	• Oxybutynin Chloride 5 mg 1 tablet (3 times	going to effect? How often will this be completed?	
discontinued or changed; (x) The name and initials of all staff	daily) – Blank 9/16 (3:00 PM)	Who is responsible? What steps will be taken if	
		issues are found?): \rightarrow	
administering medications.	Vitamin C 250 mg 1 tablet (2 times daily) –		
Model Custodial Procedure Manual	Blank 9/16 (4:00 PM)		
D. Administration of Drugs			
Unless otherwise stated by practitioner,	• Vitamin D3 400 units 2 tablets (1 time daily)		
patients will not be allowed to administer their	– Blank 9/2 (8:00 AM)		
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.			

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Developmental Disabilities (DD) Waiver Service	
Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015	
CHAPTER 5 (CIES) 1. Scope of Service B.	
Self Employment 8. Providing assistance with	
medication delivery as outlined in the ISP; C.	
Individual Community Integrated	
Employment 3. Providing assistance with	
medication delivery as outlined in the ISP; D.	
Group Community Integrated Employment 4.	
Providing assistance with medication delivery as	
outlined in the ISP; and	
B. Community Integrated Employment	
Agency Staffing Requirements: o. Comply	
with DDSD Medication Assessment and Delivery	
Policy and Procedures;	
CHAPTER 6 (CCS) 1. Scope of Services A.	
Individualized Customized Community	
Supports 19. Providing assistance or supports	
with medications in accordance with DDSD	
Medication Assessment and Delivery policy. C.	
Small Group Customized Community	
Supports 19. Providing assistance or supports	
with medications in accordance with DDSD	
Medication Assessment and Delivery policy. D.	
Group Customized Community Supports 19.	
Providing assistance or supports with	
medications in accordance with DDSD	
Medication Assessment and Delivery policy.	
CHAPTER 11 (FL) 1 SCOPE OF SERVICES	
A. Living Supports- Family Living Services:	
The scope of Family Living Services includes,	
but is not limited to the following as identified by	
the Interdisciplinary Team (IDT):	
19. Assisting in medication delivery, and related	
monitoring, in accordance with the DDSD's	
Medication Assessment and Delivery Policy,	
New Mexico Nurse Practice Act, and Board of	
Pharmacy regulations including skill	

development activities leading to the ability for	
individuals to self-administer medication as	
appropriate; and	
I. Healthcare Requirements for Family Living.	
3. B. Adult Nursing Services for medication	
oversight are required for all surrogate Lining	
Supports- Family Living direct support personnel	
if the individual has regularly scheduled	
medication. Adult Nursing services for	
medication oversight are required for all	
surrogate Family Living Direct Support	
Personnel (including substitute care), if the	
individual has regularly scheduled medication.	
6. Support Living- Family Living Provider	
Agencies must have written policies and	
procedures regarding medication(s) delivery and	
tracking and reporting of medication errors in	
accordance with DDSD Medication Assessment	
and Delivery Policy and Procedures, the New	
Mexico Nurse Practice Act and Board of	
Pharmacy standards and regulations.	
a. All twenty-four (24) hour residential home	
sites serving two (2) or more unrelated	
individuals must be licensed by the Board of	
Pharmacy, per current regulations;	
b. When required by the DDSD Medication	
Assessment and Delivery Policy, Medication	
Administration Records (MAR) must be	
maintained and include:	
i. The name of the individual, a transcription of	
the physician's or licensed health care	
provider's prescription including the brand	
and generic name of the medication, and	
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;	
assisting with the medication delivery,	

iv.Explanation of any medication error;		
v.Documentation of any allergic reaction or		
adverse medication effect; and		
vi.For PRN medication, instructions for the use		
of the PRN medication must include		
observable signs/symptoms or		
circumstances in which the medication is to		
be used, and documentation of effectiveness		
of PRN medication administered.		
c. The Family Living Provider Agency must		
also maintain a signature page that		
designates the full name that corresponds to		
each initial used to document administered		
or assisted delivery of each dose; and		
d. Information from the prescribing pharmacy		
regarding medications must be kept in the		
home and community inclusion service		
locations and must include the expected		
desired outcomes of administering the		
medication, signs and symptoms of adverse		
events and interactions with other		
medications.		
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.	
piovided.	
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	
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 administrating the medication, signs, and symptoms of adverse events and interactions with other medications.		
CHAPTER 13 (IMLS) 2. Service Requirements. B. There must be compliance		
Requirements. D. 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.		
Developmental Disabilities (DD) Waiver		
Service Standards effective 4/1/2007		
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 altergic reaction or adverse medication of flext; 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 affectiveness of PRN medication administered. (f) The Provider Agency shall also maintain a signature page that designates the full name that corresponds to each inful used to document administered or assisted delivery of each dose; (e) MARs are not required for individuals participating in Independent Living who self-administer their own medications shall be kept in the medication is avoid a start of the dores with other medications; (f) Information from the presenting pharmacy regions and shall include the expected desired outcomes of administrating the medications; (f) administrating the medications; 			1
 (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 administrating the medication, signs and symptoms of adverse 			
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regarding medications shall be kept in the home and community inclusion service locations and shall include the expected desired outcomes of administrating the medication, signs and symptoms of adverse	administer their own medications;		
home and community inclusion service locations and shall include the expected desired outcomes of administrating the medication, signs and symptoms of adverse			
locations and shall include the expected desired outcomes of administrating the medication, signs and symptoms of adverse			
desired outcomes of administrating the medication, signs and symptoms of adverse			
medication, signs and symptoms of adverse			
events and interactions with other medications;			
	events and interactions with other medications;		

Tag # 1A31	Standard Level Deficiency		
 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 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. 	 Based on record review, the Agency did not ensure the rights of Individuals was not restricted or limited for 1 of 7 Individuals. A review of Agency Individual files found no documentation of Positive Behavior Plans and/or Positive Behavior Crisis Plans, which contain restrictions being reviewed at least quarterly by the Human Rights Committee. No current Human Rights Approval was found for the following: Physical Restraint Last Review was dated 1/29/2013. (Individual #6) 	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?): →	
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			

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.		
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.		
Department of Health Developmental		

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).		

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Date Due
Service Domain: Medicaid Billing/Rein		QA/QI and Responsible Party kists to assure that claims are coded and party	Due
 b. A description of what occurred during the encounter or service interval; and 			
c. The signature or authenticated name of staff providing the service.			
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.		

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Tag # 5144	Standard Level Deficiency		
Adult Habilitation Reimbursement			
Developmental Disabilities (DD) Waiver	Based on record review, the Agency did not	Provider:	
Service Standards effective 4/1/2007 CHAPTER 1 III. PROVIDER AGENCY DOCUMENTATION OF SERVICE DELIVERY AND LOCATION A. General: All Provider Agencies shall	provide written or electronic documentation as evidence for each unit billed for Adult Habilitation Services for 1 of 1 individual. Individual #7	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?): \rightarrow	
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	 July 2016 The Agency billed 103 units of Adult Habilitation (T2021 U1) from 7/18/2016 through 7/22/2016. Documentation received accounted for 99 units. 	Provider:	
Provider Agency, level of services, and length of a session of service billed.B. Billable Units: The documentation of the		Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is	
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:		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?): \rightarrow	
 Date, start and end time of each service encounter or other billable service interval; 			
 (2) A description of what occurred during the encounter or service interval; and (3) The signature or authenticated name of 			
staff providing the service.			
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	Standard Level Deficiency		
 Reimbursement Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015 CHAPTER 6 (CCS) 4. REIMBURSEMENT A. Required Records: All Provider Agencies must maintain all records necessary to fully disclose the type, quality, quantity and clinical necessity of services furnished to individuals who are currently receiving services. The Provider Agency records must be sufficiently detailed to substantiate the date, time, individual name, servicing Provider Agency, nature of services, and length of a session of service billed. 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: Date, start and end time of each service encounter or other billable service interval; A description of what occurred during the encounter or service. Billable Unit: The billable unit for Individual Customized Community Supports is a fifteen (15) minute unit. 	 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. Individual #2 July 2016 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. Individual #8 August 2016 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. 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 108 units. 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. 	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?): →	

 The billable unit for Group Customized Community Supports is a fifteen (15) minute unit, with the rate category based on the NM DDW group. 		
 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).		
 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;		
 Provided in accordance with the Scope of Services; and 		
 Activities included in billable services, activities or situations. 		
 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	Т	
including administrative processing fee.		
 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.		

QMB Report of Findings – Community Options, Inc. – Metro – September 16 – 21, 2016

SUSANA MARTINEZ, GOVERNOR



LYNN GALLAGHER, SECRETARY DESIGNATE

Date:

January 5, 2017

To: Provider: Address:	Hector Johnson, State Director Community Options, Inc. 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

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