Date: July 27, 2016

To: Patricia Tarin, Director
Provider: Campo Behavioral Health, LLC
Address: 424 N. Mesilla Street
City/State/Zip: Las Cruces, New Mexico 88005

E-mail Address: patsy@campobh.com

Region: Southwest
Survey Date: June 13 - 15, 2016
Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: ***2012:*** *Living Supports* (Supported Living); *Inclusion Supports* (Customized Community Supports)

Survey Type: Routine

Team Leader: Chris Melon, MPA, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Team Members: Barbara Kane, BS, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; Deb Russell, BS, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; Corrina Strain, RN, BSN, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Dear Ms. Tarin;

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

Determination of Compliance:

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

Partial Compliance with Conditions of Participation

The following tags are identified as Condition of Participation Level Deficiencies:

- Tag # 1A28.1 Incident Mgt. System - Personnel Training

This determination is based on noncompliance with one or more CMS waiver assurances at the Condition of Participation level as well as Standard level deficiencies identified in the attached QMB Report of Findings 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.

Submission of your Plan of Correction:
Please submit your agency's Plan of Correction in the space on the two right columns of the Report of Findings. *(See attachment "A" for additional guidance in completing the Plan of Correction)*.

Within 10 business days of receipt of this letter your agency Plan of Correction must be submitted to the parties below:

1. **Quality Management Bureau, Attention: Amanda Castaneda, Plan of Correction Coordinator 1170 North Solano Suite D Las Cruces, New Mexico 88001**

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

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

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

Billing Deficiencies:
If you have deficiencies noted in this report of findings under the *Service Domain: Medicaid Billing/Reimbursement*, you must complete a Void/Adjust claims or remit the identified overpayment via a check within 30 calendar days of the date of this letter to HSD/OIG/PIU, *though this is not the preferred method of payment*. If you choose to pay via check, please include a copy of this letter with the payment. Make the check payable to the New Mexico Human Services Department and mail to:

Attention: Julie Ann Hill-Clapp
HSD/OIG
Program Integrity Unit
P.O. Box 2348
Santa Fe, New Mexico 87504-2348

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

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

Please be advised that there is a one-week lag period for applying payments received by check to Voided/Adjusted claims. During this lag period, your other claim payments may be applied to the amount you owe even though you have sent a refund, reducing your payment amount. For this reason, we recommend that you allow the system to recover the overpayment instead of sending in a check.

Request for Informal Reconsideration of Findings (IRF):
If you disagree with a finding of deficient practice, you have 10 business days upon receipt of this notice to request an IRF. Submit your request for an IRF in writing to:

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

See Attachment "C" for additional guidance in completing the request for Informal Reconsideration of Findings. The request for an IRF will not delay the implementation of your Plan of Correction which must be completed within 45 total business days (10 business days to submit your POC for approval and 35 days to implement your *approved* Plan of Correction). Providers may not appeal the nature or interpretation of the standard or regulation, the team composition or sampling methodology. If the IRF approves the modification or removal of a finding, you will be advised of any changes.

Please call the Plan of Correction Coordinator Amanda Castaneda at 575-373-5716 if you have questions about the Report of Findings or Plan of Correction. Thank you for your cooperation and for the work you perform.

Sincerely,

Chris Melon, MPA

Chris Melon, MPA
Team Lead/Healthcare Surveyor
Division of Health Improvement
Quality Management Bureau

Survey Process Employed:

Entrance Conference Date:	June 13, 2016	
Present:	**Campo Behavioral Health, LLC** Yolanda Costales, Incident Coordinator & Quality Assurance/Quality Improvement Patsy Tarin, Director **DOH/DHI/QMB** Chris Melon, MPA, Team Lead/Healthcare Surveyor Barbara Kane, BS, Healthcare Surveyor Deb Russell, BS, Healthcare Surveyor	
Exit Conference Date:	June 15, 2016	
Present:	**Campo Behavioral Health, LLC** Yolanda Costales, Incident Coordinator & Quality Assurance/Quality Improvement Kristina Rueckner, RN Patsy Tarin, Director **DOH/DHI/QMB** Chris Melon, MPA, Team Lead/Healthcare Surveyor Barbara Kane, BS, Healthcare Surveyor Deb Russell, BS, Healthcare Surveyor Corrina Strain, RN, BSN, Healthcare Surveyor **DDSD - SW Regional Office** Angie Brooks, Generalist	
Administrative Locations Visited	Number:	1
Total Sample Size	Number:	6 0 - *Jackson* Class Members 6 - Non-*Jackson* Class Members 6 - Supported Living 6 - Customized Community Supports
Total Homes Visited	Number:	4
❖ Supported Living Homes Visited	Number:	4 *Note: The following Individuals share a SL residence:* ➢ #1, 6 ➢ #4, 5
Persons Served Records Reviewed	Number:	6
Persons Served Interviewed	Number:	4
Persons Served Observed	Number:	1 (One individual chose not to be interviewed)
Persons Served Not Seen and/or Not Available	Number:	1 (One individual was not available at time of visit)

Direct Support Personnel Interviewed	Number:	9
Direct Support Personnel Records Reviewed	Number:	60
Service Coordinator Records Reviewed	Number:	3

Administrative Processes and Records Reviewed:

- Medicaid Billing/Reimbursement Records for all Services Provided
- Accreditation Records
- Oversight of Individual Funds
- Individual Medical and Program Case Files, including, but not limited to:
    - Individual Service Plans
    - Progress on Identified Outcomes
    - Healthcare Plans
    - Medication Administration Records
    - Medical Emergency Response Plans
    - Therapy Evaluations and Plans
    - Healthcare Documentation Regarding Appointments and Required Follow-Up
    - Other Required Health Information
- Internal Incident Management Reports and System Process / General Events Reports
- Personnel Files, including nursing and subcontracted staff
- Staff Training Records, Including Competency Interviews with Staff
- Agency Policy and Procedure Manual
- Caregiver Criminal History Screening Records
- Consolidated Online Registry/Employee Abuse Registry
- Human Rights Committee Notes and Meeting Minutes
- Evacuation Drills of Residences and Service Locations
- Quality Assurance / Improvement Plan

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

Attachment A

Provider Instructions for Completing the QMB Plan of Correction (POC) Process

Introduction:
After a QMB Compliance Survey, your QMB Report of Findings will be sent to you via e-mail.

Each provider must develop and implement a Plan of Correction (POC) that identifies specific quality assurance and quality improvement activities the agency will implement to correct deficiencies and prevent continued deficiencies and non-compliance.

Agencies must submit their Plan of Correction within ten (10) business days from the date you receive the QMB Report of Findings. (Providers who do not submit a POC within 10 business days may be referred to the Internal Review Committee [IRC] for possible actions or sanctions).

Agencies must fully implement their approved Plan of Correction within 45 business days (10 business days to submit your POC for approval and 35 days to implement your approved Plan of Correction) from the date they receive the QMB Report of Findings (Providers who fail to complete a POC within the 45 business days allowed will be referred to the IRC for possible actions or sanctions.)

If you have questions about the Plan of Correction process, call the Plan of Correction Coordinator at 575-373-5716 or email at AmandaE.Castaneda@state.nm.us. Requests for technical assistance must be requested through your Regional DDSD Office.

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

Instructions for Completing Agency POC:

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

If a deficiency has already been corrected, the plan should state how it was corrected, the completion date (date the correction was accomplished), and how possible recurrence of the deficiency will be prevented.

The Plan of Correction must address the six required Center for Medicare and Medicaid Services (CMS) core elements to address each deficiency cited in the Report of Findings:

1. How the specific and realistic corrective action will be accomplished for individuals found to have been affected by the deficient practice.
2. How the agency will identify other individuals who have the potential to be affected by the same deficient practice, and how the agency will act to protect individuals in similar situations.
3. What QA measures will be put into place or systemic changes made to ensure that the deficient practice will not recur
4. Indicate how the agency plans to monitor its performance to make sure that solutions are sustained. The agency must develop a QA plan for ensuring that correction is achieved and

sustained. This QA plan must be implemented, and the corrective action evaluated for its effectiveness. The plan of correction is integrated into the agency quality assurance system; and

5. Include dates when corrective action will be completed. The corrective action completion dates must be acceptable to the State.

The following details should be considered when developing your Plan of Correction:

- Details about how and when Consumer, Personnel and Residential files are audited by Agency personnel to ensure they contain required documents;
- Information about how Medication Administration Records are reviewed to verify they contain all required information before they are distributed, as they are being used, and after they are completed;
- Your processes for ensuring that all staff are trained in Core Competencies, Abuse, Neglect and Exploitation Reporting, and Individual-Specific service requirements, etc.;
- How accuracy in Billing/Reimbursement documentation is assured;
- How health, safety is assured;
- For Case Management Providers, how Individual Specific Plans are reviewed to verify they meet requirements, how the timeliness of LOC packet submissions and consumer visits are tracked;
- Your process for gathering, analyzing and responding to Quality data indicators; and,
- Details about Quality Targets in various areas, current status, analyses about why targets were not met, and remedies implemented.

Note: **Instruction or in-service of staff alone may not be a sufficient plan of correction.** This is a good first step toward correction, but additional steps must be taken to ensure the deficiency is corrected and will not recur.

Completion Dates

- The plan of correction must include a **completion date** (entered in the far right-hand column) for each finding. Be sure the date is **realistic** in the amount of time your Agency will need to correct the deficiency; not to exceed 45 total business days.
- Direct care issues should be corrected immediately and monitored appropriately.
- Some deficiencies may require a staged plan to accomplish total correction.
- Deficiencies requiring replacement of equipment, etc., may require more time to accomplish correction but should show reasonable time frames.

Initial Submission of the Plan of Correction Requirements

1. The Plan of Correction must be completed on the official QMB Survey Report of Findings/Plan of Correction Form and received by QMB within ten (10) business days from the date you received the report of findings.
2. For questions about the POC process, call the POC Coordinator, Amanda Castaneda at 575-373-5716 or email at AmandaE.Castaneda@state.nm.us for assistance.
3. For Technical Assistance (TA) in developing or implementing your POC, contact your Regional DDSD Office.
4. Submit your POC to Amanda Castaneda, POC Coordinator in any of the following ways:
    a. Electronically at AmandaE.Castaneda@state.nm.us *(preferred method)*
    b. Fax to 575-528-5019, or
    c. Mail to POC Coordinator, 1170 North Solano Ste D, Las Cruces, New Mexico 88001
5. Do not submit supporting documentation (evidence of compliance) to QMB until after your POC has been approved by the QMB.
6. QMB will notify you when your POC has been “approved” or “denied.”

a. During this time, whether your POC is “approved,” or “denied,” you will have a maximum of 45 business days from the date of receipt of your Report of Findings to correct all survey deficiencies.
b. If your POC is denied, it must be revised and resubmitted as soon as possible, as the 45 business day limit is in effect.
c. If your POC is denied a second time your agency may be referred to the Internal Review Committee.
d. You will receive written confirmation when your POC has been approved by QMB and a final deadline for completion of your POC.
e. Please note that all POC correspondence will be sent electronically unless otherwise requested.

7. Failure to submit your POC within 10 business days without prior approval of an extension by QMB will result in a referral to the Internal Review Committee and the possible implementation of monetary penalties and/or sanctions.

POC Document Submission Requirements

Once your POC has been approved by the QMB Plan of Correction Coordinator you must submit copies of documents as evidence that all deficiencies have been corrected, as follows.

1. Your internal documents are due within a ***maximum*** of 45 business days of receipt of your Report of Findings.
2. It is preferred that you submit your documents via USPS or other carrier (scanned and saved to CD/DVD disc, flash drive, etc.). If the documents do not contain protected Health information (PHI) the preferred method is that you submit your documents electronically (scanned and attached to e-mails).
3. All submitted documents ***must be annotated***; please be sure the tag numbers and Identification numbers are indicated on each document submitted. Documents which are not annotated with the Tag number and Identification number may not be accepted.
4. Do not submit original documents; Please provide copies or scanned electronic files for evidence. Originals must be maintained in the agency file(s) per DDSD Standards.
5. In lieu of some documents, you may submit copies of file or home audit forms that clearly indicate cited deficiencies have been corrected, other attestations of correction must be approved by the Plan of Correction Coordinator prior to their submission.
6. When billing deficiencies are cited, you must provide documentation to justify billing and/or void and adjust forms submitted to Xerox State Healthcare, LLC for the deficiencies cited in the Report of Findings.

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

Attachment B

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

The Division of Health Improvement, Quality Management Bureau (QMB) surveys compliance of the Developmental Disabilities Waiver (DDW) standards and state and federal regulations. QMB has grouped the CMS assurances into five Service Domains: Level of Care; Plan of Care; Qualified Providers; Health, Welfare and Safety; and Administrative Oversight (note that Administrative Oversight listed in this document is not the same as the CMS assurance of Administrative Authority. Used in this context it is related to the agency's operational policies and procedures, Quality Management system and Medicaid billing and reimbursement processes.)

The QMB Determination of Compliance process is based on provider compliance or non-compliance with standards and regulations identified in the QMB Report of Findings. All deficiencies (non-compliance with standards and regulations) are identified and cited as either a Standard level deficiency or a Condition of Participation level deficiency in the QMB Reports of Findings. All deficiencies require corrective action when non-compliance is identified.

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

The Determination of Compliance for each service type is based on a provider's compliance with CoPs in the following Service Domains.

Case Management Services *(Four Service Domains)*:

- Plan of Care: ISP Development & Monitoring
- Level of Care
- Qualified Providers
- Health, Safety and Welfare

Community Living Supports / Inclusion Supports *(Three Service Domains)*:

- Service Plans: ISP Implementation
- Qualified Provider
- Health, Safety and Welfare

Conditions of Participation (CoPs)

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

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

significant harm or actual harm. It is then cited as a CoP out of compliance. If the deficiency does not rise to the level of a CoP out of compliance, it is cited as a Standard Level Deficiency.

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

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

Service Domain: Plan of Care ISP Development & Monitoring

Condition of Participation:

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

Condition of Participation:

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

Service Domain: Level of Care

Condition of Participation:

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

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

Service Domain: Qualified Providers

Condition of Participation:

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

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

Service Domain: Service Plan: ISP Implementation

Condition of Participation:

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

Service Domain: Health, Welfare and Safety

Condition of Participation:

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

Condition of Participation:

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

QMB Determinations of Compliance

Compliance with Conditions of Participation

The QMB determination of *Compliance with Conditions of Participation* indicates that a provider is in compliance with all Conditions of Participation, (CoP). The agency has obtained a level of compliance such that there is a minimal potential for harm to individuals' health and safety. To qualify for a determination of Compliance with Conditions of Participation, the provider must be in compliance with all Conditions of Participation in all relevant Service Domains. The agency may also have Standard level deficiencies (deficiencies which are not at the condition level) out of compliance in any of the Service Domains.

Partial-Compliance with Conditions of Participation

The QMB determination of *Partial-Compliance with Conditions of Participation* indicates that a provider is out of compliance with Conditions of Participation in one (1) to two (2) Service Domains. The agency may have one or more Condition level tags within a Service Domain. This partial-compliance, if not corrected, may result in a serious negative outcome or the potential for more than minimal harm to individuals' health and safety. The agency may also have Standard level deficiencies (deficiencies which are not at the condition level) in any of the Service Domains.

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

Non-Compliance with Conditions of Participation

The QMB determination of *Non-Compliance with Conditions of Participation* indicates a provider is significantly out of compliance with Conditions of Participation in multiple Service Domains. The agency may have one or more Condition level tags in each of 3 relevant Service Domains and/or 6 or more total Condition level tags in the Report of Findings. This non-compliance, if not corrected, may result in a serious negative outcome or the potential for more than minimal harm to individuals' health and safety. The agency may also have Standard level deficiencies (deficiencies which are not at the condition level) in any of the Service Domains.

Providers receiving a repeat determination of Non-Compliance will be referred by Quality Management Bureau to the Internal Review Committee (IRC) for consideration of remedies and possible actions or sanctions.

Attachment C

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

Introduction:

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

Instructions:

1. The Informal Reconsideration of the Finding (IRF) request must be received in writing to the QMB Deputy Bureau Chief **within 10 business days** of receipt of the final Report of Findings.
2. The written request for an IRF *must* be completed on the QMB Request for Informal Reconsideration of Finding form available on the QMB website: http://dhi.health.state.nm.us/qmb
3. The written request for an IRF must specify in detail the request for reconsideration and why the finding is inaccurate.
4. The IRF request must include all supporting documentation or evidence.
5. If you have questions about the IRF process, email the IRF Chairperson, Crystal Lopez-Beck at Crystal.Lopez-Beck@state.nm.us for assistance.

The following limitations apply to the IRF process:

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

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

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

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

Agency: **Campo Behavioral Health, LLC - Southwest Region**
Program: Developmental Disabilities Waiver
Service: ***2012:*** *Living Supports* (Supported Living); *Inclusion Supports* (Customized Community Supports)
Monitoring Type: Routine Survey
Survey Date: **June 13 - 15, 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 # 1A32 and LS14 / 6L14 **Individual Service Plan Implementation**	**Standard Level Deficiency**		
NMAC 7.26.5.16.C and D Development of the ISP. Implementation of the ISP. The ISP shall be implemented according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan. C. The IDT shall review and discuss information and recommendations with the individual, with the goal of supporting the individual in attaining desired outcomes. The IDT develops an ISP based upon the individual's personal vision statement, strengths, needs, interests and preferences. The ISP is a dynamic document, revised periodically, as needed, and amended to reflect progress towards personal goals and achievements consistent with the individual's future vision. This regulation is consistent with standards established for individual plan development as set forth by the commission on the accreditation of rehabilitation facilities (CARF) and/or other program accreditation approved and adopted by the developmental disabilities division and the department of health. It is the policy of the developmental disabilities division (DDD), that to the extent permitted by funding, each individual receive supports and	Based on record review, the Agency did not implement the ISP according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan for 4 of 6 individuals. As indicated by Individuals ISP the following was found with regards to the implementation of ISP Outcomes: **Residential Files Reviewed:** **Supported Living Data Collection/Data Tracking/Progress with regards to ISP Outcomes**: Individual #3 • None found regarding: Live Outcome/Action Step: "Will shower thoroughly" for 6/1 – 10, 2016. Action step is to be completed 2 times per week. • None found regarding: Live Outcome/Action Step: "Will comb her hair" for 6/1 – 10, 2016. Action step is to be completed daily.	**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?)***: →**	

services that will assist and encourage independence and productivity in the community and attempt to prevent regression or loss of current capabilities. Services and supports include specialized and/or generic services, training, education and/or treatment as determined by the IDT and documented in the ISP. D. The intent is to provide choice and obtain opportunities for individuals to live, work and play with full participation in their communities. The following principles provide direction and purpose in planning for individuals with developmental disabilities. [05/03/94; 01/15/97; Recompiled 10/31/01]	• None found regarding: Live, Outcome/Action Step: "Will brush her teeth thoroughly" for 6/1 – 10, 2016. Action step is to be completed daily. Individual #4 • None found regarding: Live Outcome/Action Step: "Will take pictures of meals" for 6/1 – 10, 2016. Action step is to be completed weekly. • None found regarding: Live Outcome/Action Step: "Will research, plan, and save money for day trip" for 6/1 – 10, 2016. Action step is to be completed weekly. Individual #5 • None found regarding: Live Outcome/Action Step: "Will complete her hygiene routine" for 6/1 – 10, 2016. Action step is to be completed 4 times per week. • None found regarding: Live Outcome/Action Step: "Will complete her assigned chores" for 6/1 - 10, 2016. Action step is to be completed 3 times per week. Individual #6 • None found regarding: Live Outcome/Action Step: "Will participate in feeding his fish the proper amount of food with staff assistance for safety" for 6/1 – 10, 2016. Action step is to be completed daily.		

Tag # LS14 / 6L14 Residential Case File	Standard Level Deficiency		
Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013 **CHAPTER 11 (FL) 3. Agency Requirements** **C. Residence Case File:** The Agency must maintain in the individual's home a complete and current confidential case file for each individual. Residence case files are required to comply with the DDSD Individual Case File Matrix policy. **CHAPTER 12 (SL) 3. Agency Requirements** **C. Residence Case File:** The Agency must maintain in the individual's home a complete and current confidential case file for each individual. Residence case files are required to comply with the DDSD Individual Case File Matrix policy. **CHAPTER 13 (IMLS) 2. Service Requirements** **B.1. Documents To Be Maintained In The Home:** a. Current Health Passport generated through the e-CHAT section of the Therap website and printed for use in the home in case of disruption in internet access; b. Personal identification; c. Current ISP with all applicable assessments, teaching and support strategies, and as applicable for the consumer, PBSP, BCIP, MERP, health care plans, CARMPs, Written Therapy Support Plans, and any other plans (e.g. PRN Psychotropic Medication Plans) as applicable; d. Dated and signed consent to release information forms as applicable; e. Current orders from health care practitioners; f. Documentation and maintenance of accurate medical history in Therap website; g. Medication Administration Records for the current month; h. Record of medical and dental appointments for the current year, or during the period of stay for short term stays, including any treatment provided;	Based on record review, the Agency did not maintain a complete and confidential case file in the residence for 6 of 6 Individuals receiving Supported Living Services. Review of the residential individual case files revealed the following items were not found, incomplete, and/or not current: • **Current Emergency and Personal Identification Information** ◦ Individual's address not current (#3) ◦ Individual's phone number not current (#3) • Occupational Therapy Plan (#5) • Healthcare Passport (#3, 4, 5, 6) • **Special Health Care Needs** ◦ Nutritional Plan (#6) ◦ Comprehensive Aspiration Risk Management Plan: ➢ Not Current (#1) • **Medical Emergency Response Plans** ◦ Respiratory (#3) • **Progress Notes/Daily Contacts Logs:** ◦ Individual #1 - None found for 6/1 – 10, 2016. ◦ Individual #2 - None found for 6/1 – 10, 2016. ◦ Individual #3 - None found for 6/1– 10, 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?)*: → **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?)*: →	

i. Progress notes written by DSP and nurses; j. Documentation and data collection related to ISP implementation; k. Medicaid card; l. Salud membership card or Medicare card as applicable; and m. A Do Not Resuscitate (DNR) document and/or Advanced Directives as applicable. **DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION (DDSD): Director's Release: Consumer Record Requirements eff. 11/1/2012** **III. Requirement Amendments(s) or Clarifications:** A. All case management, living supports, customized in-home supports, community integrated employment and customized community supports providers must maintain records for individuals served through DD Waiver in accordance with the Individual Case File Matrix incorporated in this director's release. H. Readily accessible electronic records are accessible, including those stored through the Therap web-based system. ***Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007*** **CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS** **A. Residence Case File:** For individuals receiving Supported Living or Family Living, the Agency shall maintain in the individual's home a complete and current confidential case file for each individual. For individuals receiving Independent Living Services, rather than maintaining this file at the individual's home, the complete and current confidential case file for each individual shall be maintained at the agency's administrative site. Each file shall include the following: (1) Complete and current ISP and all supplemental plans specific to the individual;	◦ Individual #4 - None found for 6/1 – 10, 2016. ◦ Individual #5 - None found for 6/1 – 10, 2016 ◦ Individual #6 - None found for 6/1 – 10, 2016		

(2) Complete and current Health Assessment Tool; (3) Current emergency contact information, which includes the individual's address, telephone number, names and telephone numbers of residential Community Living Support providers, relatives, or guardian or conservator, primary care physician's name(s) and telephone number(s), pharmacy name, address and telephone number and dentist name, address and telephone number, and health plan; (4) Up-to-date progress notes, signed and dated by the person making the note for at least the past month (older notes may be transferred to the agency office); (5) Data collected to document ISP Action Plan implementation (6) Progress notes written by direct care staff and by nurses regarding individual health status and physical conditions including action taken in response to identified changes in condition for at least the past month; (7) Physician's or qualified health care providers written orders; (8) Progress notes documenting implementation of a physician's or qualified health care provider's order(s); (9) Medication Administration Record (MAR) for the past three (3) months which includes: (a) The name of the individual; (b) A transcription of the healthcare practitioners prescription including the brand and generic name of the medication; (c) Diagnosis for which the medication is prescribed; (d) Dosage, frequency and method/route of delivery; (e) Times and dates of delivery; (f) Initials of person administering or assisting with medication; and (g) An explanation of any medication irregularity, allergic reaction or adverse effect.			

(h) For PRN medication an explanation for the use of the PRN must include: (i) Observable signs/symptoms or circumstances in which the medication is to be used, and (ii) Documentation of the effectiveness/result of the PRN delivered. (i) A MAR is not required for individuals participating in Independent Living Services who self-administer their own medication. However, when medication administration is provided as part of the Independent Living Service a MAR must be maintained at the individual's home and an updated copy must be placed in the agency file on a weekly basis. (10) Record of visits to healthcare practitioners including any treatment provided at the visit and a record of all diagnostic testing for the current ISP year; and (11) Medical History to include: demographic data, current and past medical diagnoses including the cause (if known) of the developmental disability and any psychiatric diagnosis, allergies (food, environmental, medications), status of routine adult health care screenings, immunizations, hospital discharge summaries for past twelve (12) months, past medical history including hospitalizations, surgeries, injuries, family history and current physical exam.			

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Date Due
Service Domain: Qualified Providers – *The State monitors non-licensed/non-certified providers to assure adherence to waiver requirements. The State implements its policies and procedures for verifying that provider training is conducted in accordance with State requirements and the approved waiver.*			
Tag # 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:** I. Staff providing direct services shall complete safety training within the first thirty (30) days of employment and before working alone with an individual receiving services. The training shall address at least the following: 1. Operating a fire extinguisher 2. Proper lifting procedures 3. General vehicle safety precautions (e.g., pre-trip inspection, removing keys from the ignition when not in the driver's seat) 4. Assisting passengers with cognitive and/or physical impairments (e.g., general guidelines for supporting individuals who may be unaware of safety issues involving traffic or those who require physical assistance to enter/exit a vehicle) 5. Operating wheelchair lifts (if applicable to the staff's role) 6. Wheelchair tie-down procedures (if applicable to the staff's role) 7. Emergency and evacuation procedures (e.g., roadside emergency, fire emergency) **NMAC 7.9.2 F. TRANSPORTATION:** **(1)** Any employee or agent of a regulated facility or agency who is responsible for assisting a resident in boarding or alighting from a motor vehicle must complete a state-approved training program in passenger transportation assistance	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 60 Direct Support Personnel. **No documented evidence was found of the following required training:** • Transportation (DSP #220)	**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?)***:** →	

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 **CHAPTER 5 (CIES) 3. Agency Requirements G. Training Requirements: 1.** All Community Inclusion Providers must provide staff training in accordance with the DDSD policy T-003: Training Requirements for Direct Service Agency Staff Policy. **CHAPTER 6 (CCS) 3. Agency Requirements F. Meet all training requirements as follows: 1.** All Customized Community Supports Providers shall provide staff training in accordance with the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff Policy; **CHAPTER 7 (CIHS) 3. Agency Requirements C. Training Requirements:** The Provider Agency must report required personnel training status to the DDSD Statewide Training Database as specified in the DDSD Policy T-001: Reporting and Documentation of DDSD Training Requirements Policy. The Provider Agency must ensure that the personnel support staff have completed training as specified in the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff Policy **CHAPTER 11 (FL) 3. Agency Requirements B. Living Supports- Family Living Services Provider Agency Staffing Requirements: 3. Training:** A. All Family Living Provider agencies must ensure staff training in accordance with the Training Requirements for Direct Service Agency Staff policy. DSP's or subcontractors delivering substitute care under Family Living must at a minimum comply with the section of the training policy that relates to Respite, Substitute Care, and personal support staff [Policy T-003: for Training Requirements for Direct Service Agency Staff; Sec. II-J, Items 1-4]. Pursuant to the Centers for Medicare and Medicaid Services (CMS)			

requirements, the services that a provider renders may only be claimed for federal match if the provider has completed all necessary training required by the state. All Family Living Provider agencies must report required personnel training status to the DDSD Statewide Training Database as specified in DDSD Policy T-001: Reporting and Documentation for DDSD Training Requirements. **CHAPTER 12 (SL) 3. Agency Requirements B. Living Supports- Supported Living Services Provider Agency Staffing Requirements: 3. Training:** A. All Living Supports- Supported Living Provider Agencies must ensure staff training in accordance with the DDSD Policy T-003: for Training Requirements for Direct Service Agency Staff. Pursuant to CMS requirements, the services that a provider renders may only be claimed for federal match if the provider has completed all necessary training required by the state. All Supported Living provider agencies must report required personnel training status to the DDSD Statewide Training Database as specified in DDSD Policy T-001: Reporting and Documentation for DDSD Training Requirements. **CHAPTER 13 (IMLS) R. 2. Service Requirements. Staff Qualifications 2. DSP Qualifications.** E. Complete training requirements as specified in the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff - effective March 1, 2007. Report required personnel training status to the DDSD Statewide Training Database as specified in the DDSD Policy T-001: Reporting and Documentation of DDSD Training Requirements Policy;			

Tag # 1A20 Direct Support Personnel Training	Standard Level Deficiency		
Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy - Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007 - II. POLICY STATEMENTS: A. Individuals shall receive services from competent and qualified staff. B. Staff shall complete individual-specific (formerly known as "Addendum B") training requirements in accordance with the specifications described in the individual service plan (ISP) of each individual served. C. Staff shall complete training on DOH-approved incident reporting procedures in accordance with 7 NMAC 1.13. D. Staff providing direct services shall complete training in universal precautions on an annual basis. The training materials shall meet Occupational Safety and Health Administration (OSHA) requirements. E. Staff providing direct services shall maintain certification in first aid and CPR. The training materials shall meet OSHA requirements/guidelines. F. Staff who may be exposed to hazardous chemicals shall complete relevant training in accordance with OSHA requirements. G. Staff shall be certified in a DDSD-approved behavioral intervention system (e.g., Mandt, CPI) before using physical restraint techniques. Staff members providing direct services shall maintain certification in a DDSD-approved behavioral intervention system if an individual they support has a behavioral crisis plan that includes the use of physical restraint techniques. H. Staff shall complete and maintain certification in a DDSD-approved medication course in accordance with the DDSD Medication Delivery Policy M-001. I. Staff providing direct services shall complete safety training within the first thirty (30) days of	Based on record review, the Agency did not ensure Orientation and Training requirements were met for 19 of 60 Direct Support Personnel. Review of Direct Support Personnel training records found no evidence of the following required DOH/DDSD trainings and certification being completed: • Pre- Service (DSP #201) • Person-Centered Planning (1-Day) (DSP #220) • First Aid (DSP #202, 204, 205, 222, 246, 252) • CPR (DSP #202, 204, 205, 222, 246, 252) • Assisting With Medication Delivery (DSP #217, 228, 232, 238, 242, 245, 246, 247, 252, 253, 254, 255) • Participatory Communication and Choice Making (DSP #259) • Rights and Advocacy (DSP #259) • Supporting People with Challenging Behaviors (DSP #259) • Teaching and Support Strategies (DSP #259)	**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?)*: →	

employment and before working alone with an individual receiving service. Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013 **CHAPTER 5 (CIES) 3. Agency Requirements G. Training Requirements: 1.** All Community Inclusion Providers must provide staff training in accordance with the DDSD policy T-003: Training Requirements for Direct Service Agency Staff Policy. **CHAPTER 6 (CCS) 3. Agency Requirements F. Meet all training requirements as follows: 1.** All Customized Community Supports Providers shall provide staff training in accordance with the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff Policy; **CHAPTER 7 (CIHS) 3. Agency Requirements C. Training Requirements:** The Provider Agency must report required personnel training status to the DDSD Statewide Training Database as specified in the DDSD Policy T-001: Reporting and Documentation of DDSD Training Requirements Policy. The Provider Agency must ensure that the personnel support staff have completed training as specified in the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff Policy **CHAPTER 11 (FL) 3. Agency Requirements B. Living Supports- Family Living Services Provider Agency Staffing Requirements: 3. Training:** A. All Family Living Provider agencies must ensure staff training in accordance with the Training Requirements for Direct Service Agency Staff policy. DSP's or subcontractors delivering substitute care under Family Living must at a minimum comply with the section of the training policy that relates to Respite, Substitute Care, and personal support staff [Policy T-003: for Training Requirements for Direct Service Agency Staff; Sec.			

II-J, Items 1-4]. Pursuant to the Centers for Medicare and Medicaid Services (CMS) requirements, the services that a provider renders may only be claimed for federal match if the provider has completed all necessary training required by the state. All Family Living Provider agencies must report required personnel training status to the DDSD Statewide Training Database as specified in DDSD Policy T-001: Reporting and Documentation for DDSD Training Requirements. **CHAPTER 12 (SL) 3. Agency Requirements B. Living Supports- Supported Living Services Provider Agency Staffing Requirements: 3. Training:** A. All Living Supports- Supported Living Provider Agencies must ensure staff training in accordance with the DDSD Policy T-003: for Training Requirements for Direct Service Agency Staff. Pursuant to CMS requirements, the services that a provider renders may only be claimed for federal match if the provider has completed all necessary training required by the state. All Supported Living provider agencies must report required personnel training status to the DDSD Statewide Training Database as specified in DDSD Policy T-001: Reporting and Documentation for DDSD Training Requirements. **CHAPTER 13 (IMLS) R. 2. Service Requirements. Staff Qualifications 2. DSP Qualifications.** E. Complete training requirements as specified in the DDSD Policy T-003: Training Requirements for Direct Service Agency Staff - effective March 1, 2007. Report required personnel training status to the DDSD Statewide Training Database as specified in the DDSD Policy T-001: Reporting and Documentation of DDSD Training Requirements Policy;			

Tag # 1A28.1 Incident Mgt. System - Personnel Training	Condition of Participation Level Deficiency		
NMAC 7.1.14 ABUSE, NEGLECT, EXPLOITATION, AND DEATH REPORTING, TRAINING AND RELATED REQUIREMENTS FOR COMMUNITY PROVIDERS **NMAC 7.1.14.9 INCIDENT MANAGEMENT SYSTEM REQUIREMENTS:** **A. General:** All community-based service providers shall establish and maintain an incident management system, which emphasizes the principles of prevention and staff involvement. The community-based service provider shall ensure that the incident management system policies and procedures requires all employees and volunteers to be competently trained to respond to, report, and preserve evidence related to incidents in a timely and accurate manner. **B. Training curriculum:** Prior to an employee or volunteer's initial work with the community-based service provider, all employees and volunteers shall be trained on an applicable written training curriculum including incident policies and procedures for identification, and timely reporting of abuse, neglect, exploitation, suspicious injury, and all deaths as required in Subsection A of 7.1.14.8 NMAC. The trainings shall be reviewed at annual, not to exceed 12-month intervals. The training curriculum as set forth in Subsection C of 7.1.14.9 NMAC may include computer-based training. Periodic reviews shall include, at a minimum, review of the written training curriculum and site-specific issues pertaining to the community-based service provider's facility. Training shall be conducted in a language that is understood by the employee or volunteer. **C. Incident management system training curriculum requirements:** **(1)** The community-based service provider shall conduct training or designate a knowledgeable representative to conduct	After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur. Based on record review and interview, the Agency did not ensure Incident Management Training for 19 of 63 Agency Personnel. **Direct Support Personnel (DSP):** • Incident Management Training (Abuse, Neglect and Exploitation) (DSP# 201, 202, 212, 220, 221, 227, 231, 233, 241, 243, 251, 252, 253, 254, 257) **Service Coordination Personnel (SC):** • Incident Management Training (Abuse, Neglect and Exploitation) (SC #260, 261, 262) **When Direct Support Personnel were asked what State Agency must be contacted when there is suspected Abuse, Neglect and Exploitation, the following was reported:** • DSP #225 stated, "DHS." Staff was not able to identify the State Agency as Division of Health Improvement.	**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?)*: →	

training, in accordance with the written training curriculum provided electronically by the division that includes but is not limited to: **(a)** an overview of the potential risk of abuse, neglect, or exploitation; **(b)** informational procedures for properly filing the division's abuse, neglect, and exploitation or report of death form; **(c)** specific instructions of the employees' legal responsibility to report an incident of abuse, neglect and exploitation, suspicious injury, and all deaths; **(d)** specific instructions on how to respond to abuse, neglect, or exploitation; **(e)** emergency action procedures to be followed in the event of an alleged incident or knowledge of abuse, neglect, exploitation, or suspicious injury. **(2)** All current employees and volunteers shall receive training within 90 days of the effective date of this rule. **(3)** All new employees and volunteers shall receive training prior to providing services to consumers. **D. Training documentation:** All community-based service providers shall prepare training documentation for each employee and volunteer to include a signed statement indicating the date, time, and place they received their incident management reporting instruction. The community-based service provider shall maintain documentation of an employee or volunteer's training for a period of at least three years, or six months after termination of an employee's employment or the volunteer's work. Training curricula shall be kept on the provider premises and made available upon request by the department. Training documentation shall be made available immediately upon a division representative's request. Failure to provide employee and volunteer training documentation			

shall subject the community-based service provider to the penalties provided for in this rule. **Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007 II. POLICY STATEMENTS:** A. Individuals shall receive services from competent and qualified staff. C. Staff shall complete training on DOH-approved incident reporting procedures in accordance with 7 NMAC 1.13.			

Tag #1A40 Provider Requirement Accreditation	Standard Level Deficiency		
NMAC 7.26.6.6 OBJECTIVE: A. These regulations are being promulgated to promote and assure the provision of quality services to persons with developmental disabilities residing in community agencies. B. These regulations are being promulgated as part of a quality assurance initiative requiring all community agencies providing services to persons with developmental disabilities and contracting with the developmental disabilities division to be accredited by the commission on accreditation of rehabilitation facilities (CARF). **7.26.6.14 CARF STANDARDS MANUAL FOR ORGANIZATIONS SERVING PEOPLE WITH DEVELOPMENTAL DISABILITIES:** Community agencies governed by these regulations are required to meet applicable provisions of the most current edition of the "CARF Standards Manual for Organizations Serving People with Disabilities". Sections of the CARF standards may be waived by the Department when deemed not applicable to the services provided by the community agency. **Long Term Services Division Policy -** Accreditation of Long Term Services Division Funded Providers eff. August 30, 2004 **A. Mandate for Accreditation** The Department of Health, Long Term Services Division (hereafter referred to as the Division) will contract only with agencies/organizations accredited in compliance with this policy. 1. Within eighteen (18) months of an initial contract or change in exemption status as defined in this policy, the contractor must provide the Division with written verification of accreditation from the Commission on Accreditation of Rehabilitation Facilities	Based on observation and interview, the Agency did not obtain the Commission on Accreditation of Rehabilitation Facilities (CARF) or the Council on Quality and Leadership in Supports for People with Disabilities (The Council) accreditation or the applicable waiver from the Developmental Disability Support Division. **When #263 was asked if the Agency had evidence of current CARF accreditation or a waiver from DDSD the following was reported:** • #263 stated, "Our last CARF accreditation was in 2010. We have had two consecutive 3 year accreditations (2007 and 2010) so are eligible for an exemption." The aforementioned 2010 CARF accreditation plaque was presented, however no further evidence/documentation of a waiver from DDSD was provided.	**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?)*: →	

(CARF) or the Council on Quality and Leadership in Supports for People with Disabilities (The Council). 2. Except as provided in this policy, the Division may terminate its contract with a contractor that fails to maintain an accreditation status of at least one year, regardless of any appeal process available from CARF or the Council.			

Tag # 1A43 General Events Reporting	Standard Level Deficiency		
Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy: General Events Reporting Effective 1/1/2012 **1. Purpose** To report, track and analyze significant events experiences by adult participants of the DD Waiver program, which do not meet criteria for abuse, neglect or exploitation, or other "reportable incident" as defined by the Incident Management Bureau of the Division of Health Improvement, Department of Health, but which pose a risk to individuals served. Analysis of reported significant events is intended to identify emerging patterns so that preventative actions can be identified at the individual, provider agency, regional and statewide levels. **II. Policy Statements** A. Designated employees of each agency will enter specified information into the General Events Reporting section of the secure website operated under contract by Therap Services within 2 business days of the occurrence or knowledge by the reporting agency of any of the following defined events in which DDSD requires reporting: Chocking, Missing Person, Suicide Attempt or Threat, Restraint related to Behavior, Serious Injury including Skin Breakdown, Fall (with or without injury), Out of Home Placement and Infections…Providers shall utilize the "Significant Events Reporting System Guide" to assure that events are reported correctly for DDSD tracking purposes. At providers' discretion additional events may be tracked within the Therap General Events Reporting	Based on record review the Agency did not follow the General Events Reporting requirements as indicated by the policy. **Agency record review revealed the following incidents were not entered in the General Events Reporting System as required for 2 of 6 individuals:** Individual #1 • Review of agency internal Incident Reports revealed that the individual had falls in the home on 12/19/2015 and 11/20/2015. Review of the agency's General Events Reports produced no evidence of these incidents being entered as GERs. (Injury) Individual #5 • Review of agency internal Incident Reports revealed that the individual had a fall in the home on 12/6/2015. Review of the agency's General Events Reports produced no evidence of this incident being entered as a GER. (Injury)	**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?)*: →	

which are not required by DDSD such as medication errors. B. General Events Reporting does not replace agency obligations to report abuse, neglect, exploitation and other reportable incidents in compliance with policies and procedures issued by the Department's Incident Management Bureau of the Division of Health Improvement. D. On at least a quarterly and annual basis, provider agencies shall analyze general events reporting data at both the individual level and agency wide to identify any patterns which warrant preventative or corrective action. If multiple events are noted for particular individuals, the agency shall consider the need to contact the case manager to convene interdisciplinary team meetings to discuss prevention measures. Agency level data shall be used as part of the agencies continuous quality improvement activities when patterns are noted across the agency or for particular service delivery sites. **New Mexico DDSD General Events Report (GER)** DDSD Revised 10-24-14			

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Date Due
Service Domain: Health and Welfare – *The state, on an ongoing basis, identifies, addresses and seeks to prevent occurrences of abuse, neglect and exploitation. Individuals shall be afforded their basic human rights. The provider supports individuals to access needed healthcare services in a timely manner.*			
Tag #1A08.2 Healthcare Requirements	**Standard Level Deficiency**		
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. **DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION (DDSD): Director's Release: Consumer Record Requirements eff. 11/1/2012** **III. Requirement Amendments(s) or Clarifications:** A. All case management, living supports, customized in-home supports, community integrated employment and customized community supports providers must maintain records for individuals served through DD Waiver in accordance with the Individual Case File Matrix incorporated in this director's release. H. Readily accessible electronic records are accessible, including those stored through the Therap web-based system. Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013 **Chapter 5 (CIES) 3. Agency Requirements**	Based on record review, the Agency did not provide documentation of annual physical examinations and/or other examinations as specified by a licensed physician for 1 of 6 individuals receiving Community Inclusion, Living Services and Other Services. Review of the administrative individual case files revealed the following items were not found, incomplete, and/or not current: ***Community Living Services / Community Inclusion Services (Individuals Receiving Multiple Services):*** • **Dental Exam** ◦ Individual #4 - As indicated by collateral documentation reviewed, exam was completed on 7/15/2015. Follow-up was to be completed in 1 month. The individual refused to attend. No further appointments were made. No evidence of follow-up found.	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)***: →** **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)***: →**	

H. Consumer Records Policy: All Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Consumer Records Policy. **Chapter 6 (CCS) 3. Agency Requirements: G. Consumer Records Policy:** All Provider Agencies shall maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix policy. **Chapter 7 (CIHS) 3. Agency Requirements: E. Consumer Records Policy:** All Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix policy. **Chapter 11 (FL) 3. Agency Requirements: D. Consumer Records Policy:** All Family Living Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix policy. **Chapter 12 (SL) 3. Agency Requirements: D. Consumer Records Policy:** All Living Supports- Supported Living Provider Agencies must maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDSD Individual Case File Matrix policy. **Chapter 13 (IMLS) 2. Service Requirements:**			

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)… 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 providers. The record must also be made available for review when requested by DOH, HSD or federal government representatives for oversight purposes. The individual's case file shall include the following requirements: (5) A medical history, which shall include at least demographic data, current and past medical diagnoses including the cause (if known) of the developmental disability, psychiatric diagnoses, allergies (food, environmental, medications), immunizations, and most recent physical exam; **CHAPTER 6. VI. GENERAL REQUIREMENTS FOR COMMUNITY LIVING** **G. Health Care Requirements for Community Living Services**. (1) The Community Living Service providers shall ensure completion of a HAT for each individual receiving this service. The HAT shall be completed 2 weeks prior to the annual ISP meeting and submitted to the Case Manager and all other IDT Members. A revised HAT is required to also be submitted whenever the individual's health status changes significantly. For individuals who are newly allocated to the			

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

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

Tag # 1A03 CQI System	Standard Level Deficiency		
STATE OF NEW MEXICO DEPARTMENT OF HEALTH DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION PROVIDER AGREEMENT: ARTICLE 17. PROGRAM EVALUATIONS d. PROVIDER shall have a Quality Management and Improvement Plan in accordance with the current MF Waiver Standards and/or the DD Waiver Standards specified by the DEPARTMENT. The Quality Management and Improvement Plan for DD Waiver Providers must describe how the PROVIDER will determine that each waiver assurance and requirement is met**.** The applicable assurances and requirements are: (1) level of care determination; (2) service plan; (3) qualified providers; (4) health and welfare; (5) administrative authority; and, (6) financial accountability. For each waiver assurance, this description must include: i. Activities or processes related to discovery, i.e., monitoring and recording the findings. Descriptions of monitoring/oversight activities that occur at the individual and provider level of service delivery. These monitoring activities provide a foundation for Quality Management by generating information that can be aggregated and analyzed to measure the overall system performance; ii. The entities or individuals responsible for conducting the discovery/monitoring processes; iii. The types of information used to measure performance; and, iv. The frequency with which performance is measured.	Based on record review, the Agency did not implement their Continuous Quality Management System as required by standard. Review of the Agency's CQI Plan revealed the following: • The Agency's CQI Plan did not contain the following components: a. **Implementation of ISPs:** extent to which services are delivered in accordance with ISPs and associated support plans with WDSI including the type, scope, amount, duration and frequency specified in the ISP as well as effectiveness of such implementation as indicated by achievement of outcomes; • Effectiveness and timeliness of implementation of ISPs, associated support plans, and WDSI, including trends in achievement of individual desired outcomes; ***(CCS)*** • Effectiveness and timeliness of implementation of ISPs, including trends in achievement of individual desired outcomes ***(SL)*** b. Compliance with Caregivers Criminal History Screening requirements; c. Compliance with Employee Abuse Registry requirements; d. Compliance with DDSD training requirements; e. Patterns/Trends of reportable incidents;	**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?)***:** →	

Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013 **CHAPTER 5 (CIES) 3. Agency Requirements: J. Quality Assurance/Quality Improvement (QA/QI) Program:** Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QA/QI activities. 1. **Development of a QA/QI plan:** The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working. 2. **Implementing a QA/QI Committee:** The QA/QI committee must convene on at least a quarterly basis and as needed to review service reports, to identify any deficiencies, trends, patterns or concerns as well as opportunities for quality improvement. The QA/QI meeting must be documented. The QA/QI review should address at least the following: a.**Implementation of ISPs:** extent to which services are delivered in accordance with ISPs and associated support plans with WDSI including the type, scope, amount, duration and frequency specified in the ISP as well as	f. Results of improvement actions taken in previous quarters; g. Action taken regarding individual grievances; h. Presence and completeness of required documentation; i. Significant program changes. j. A description of how data collected as part of the agency's QA/QI Plan was used; what quality improvement initiatives were undertaken and what were the results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QA/QI process; and ***(CCS, SL)*** k. Patterns / Trends in medication errors ***(SL)*** • Review of the Agency's Quality Improvement plan did not contain the following Incident Management specific areas: **(1)** community-based service providers shall have current abuse, neglect, and exploitation management policy and procedures in place that comply with the department's requirements; **(2)** community-based service providers providing intellectual and developmental disabilities services must have a designated incident management coordinator in place; and **(3)** community-based service providers providing intellectual and developmental disabilities services must have an incident management committee to identify any deficiencies, trends, patterns, or concerns as well as opportunities for quality improvement, address internal and		

effectiveness of such implementation as indicated by achievement of outcomes; 3. The Provider Agency must complete a QA/QI report annually by February 15th of each calendar year or as otherwise requested by DOH. The report must be kept on file at the agency, made available for review by DOH and upon request from DDSD; the report must be submitted to the relevant DDSD Regional Offices. The report will summarize: a. Analysis of General Events Reports data in Therap; b. Compliance with Caregivers Criminal History Screening requirements; c. Compliance with Employee Abuse Registry requirements; d. Compliance with DDSD training requirements; e. Patterns of reportable incidents; f. Results of improvement actions taken in previous quarters; g. Sufficiency of staff coverage; h. Effectiveness and timeliness of implementation of ISPs, and associated support including trends in achievement of individual desired outcomes; i. Results of General Events Reporting data analysis; j. Action taken regarding individual grievances; k. Presence and completeness of required documentation; **l.** A description of how data collected as part of the agency's QA/QI Plan was used; what quality improvement initiatives were undertaken and what were the results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QA/QI process; and **m.** Significant program changes.	external incident reports for the purpose of examining internal root causes, and to take action on identified issues.		

CHAPTER 6 (CCS) 3. Agency Requirements: I. Quality Assurance/Quality Improvement (QA/QI) Program: Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QI activities. 1. **Development of a QI plan:** The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working. 2. **Implementing a QI Committee:** The QA/QI committee shall convene at least quarterly and as needed to review service reports, to identify any deficiencies, trends, patterns or concerns as well as opportunities for quality improvement. The QA/QI meeting shall be documented. The QA/QI review should address at least the following: a. The extent to which services are delivered in accordance with ISPs, associated support plans and WDSI including the type, scope, amount, duration and frequency specified in the ISP as well as effectiveness of such implementation as indicated by achievement of outcomes; b. Analysis of General Events Reports data;			

c. Compliance with Caregivers Criminal History Screening requirements; d. Compliance with Employee Abuse Registry requirements; e. Compliance with DDSD training requirements; f. Patterns of reportable incidents; and g. Results of improvement actions taken in previous quarters. 3. The Provider Agencies must complete a QA/QI report annually by February 15th of each year, or as otherwise requested by DOH. The report must be kept on file at the agency, made available for review by DOH and upon request from DDSD the report must be submitted to the relevant DDSD Regional Offices. The report will summarize: a. Sufficiency of staff coverage; b. Effectiveness and timeliness of implementation of ISPs, associated support plans, and WDSI, including trends in achievement of individual desired outcomes; c. Results of General Events Reporting data analysis; d. Action taken regarding individual grievances; e. Presence and completeness of required documentation; f. A description of how data collected as part of the agency's QI plan was used; what quality improvement initiatives were undertaken and what were the results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QI process; and g. Significant program changes. **CHAPTER 7 (CIHS) 3. Agency Requirements: G. Quality Assurance/Quality Improvement (QA/QI) Program:** Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This			

includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QA/QI activities. 1. **Development of a QA/QI plan:** The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working. 2. **Implementing a QA/QI Committee:** The QA/QI committee shall convene on at least a quarterly basis and as needed to review monthly service reports, to identify any deficiencies, trends, patterns or concerns as well as opportunities for quality improvement. The QA/QI meeting must be documented. The QA/QI review should address at least the following: a. **Implementation of ISPs:** The extent to which services are delivered in accordance with ISPs and associated support plans and/or WDSI including the type, scope, amount, duration and frequency specified in the ISP as well as effectiveness of such implementation as indicated by achievement of outcomes; b. Analysis of General Events Reports data;			

c. Compliance with Caregivers Criminal History Screening requirements; d. Compliance with Employee Abuse Registry requirements; e. Compliance with DDSD training requirements; f. Patterns of reportable incidents; and g. Results of improvement actions taken in previous quarters. 3. The Provider Agency must complete a QA/QI report annually by February 15th of each calendar year, or as otherwise request by DOH. The report must be kept on file at the agency, made available for review by DOH and, upon request from DDSD the report must be submitted to the relevant DDSD Regional Offices. The report will summarize: a. Sufficiency of staff coverage; b. Effectiveness and timeliness of implementation of ISPs and associated support plans and/or WDSI, including trends in achievement of individual desired outcomes; c. Results of General Events Reporting data analysis; d. Action taken regarding individual grievances; e. Presence and completeness of required documentation; **f.** A description of how data collected as part of the agency's QA/QI plan was used; what quality improvement initiatives were undertaken and what were the results of			

those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QI process; and **g.** Significant program changes. **CHAPTER 11 (FL) 3. Agency Requirements: H. Quality Improvement/Quality Assurance (QA/QI) Program:** Family Living Provider Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QA/QI activities. 1. **Development of a QA/QI plan:** The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working. 2. **Implementing a QA/QI Committee:** The QA/QI committee must convene on at least a quarterly basis and as needed to review monthly service reports, to identify any deficiencies, trends, patterns or concerns as well as opportunities for quality improvement. The QA/QI meeting must be documented. The			

QA/QI review should address at least the following: a. The extent to which services are delivered in accordance with the ISP including the type, scope, amount, duration and frequency specified in the ISP as well as effectiveness of such implementation as indicated by achievement of outcomes; b. Analysis of General Events Reports data; c. Compliance with Caregivers Criminal History Screening requirements; d. Compliance with Employee Abuse Registry requirements; e. Compliance with DDSD training requirements; f. Patterns in reportable incidents; and g. Results of improvement actions taken in previous quarters. 3. The Provider Agency must complete a QA/QI report annually by February 15th of each year, or as otherwise requested by DOH. The report must be kept on file at the agency, made available for review by DOH and upon request from DDSD; the report must be submitted to the relevant DDSD Regional Offices. The report will summarize: a. Sufficiency of staff coverage; b. Effectiveness and timeliness of implementation of ISPs, including trends in achievement of individual desired outcomes; c. Results of General Events Reporting data analysis, Trends in category II significant events; d. Patterns in medication errors; e. Action taken regarding individual grievances; f. Presence and completeness of required documentation; g. A description of how data collected as part of the agency's QI plan was used;			

h. What quality improvement initiatives were undertaken and what were the results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QI process; and i. Significant program changes. **CHAPTER 12 (SL) 3. Agency Requirements: B. Quality Assurance/Quality Improvement (QA/QI) Program:** Supported Living Provider Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QA/QI activities. 1. **Development of a QA/QI plan:** The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working. 2. **Implementing a QA/QI Committee:** The QA/QI committee must convene on at least a quarterly basis and as needed to review monthly service reports, to identify any deficiencies, trends, patterns, or concerns as well as opportunities for quality improvement. The QA/QI meeting must be documented. The			

QA/QI review should address at least the following: a. Implementation of the ISP and the extent to which services are delivered in accordance with the ISP including the type, scope, amount, duration, and frequency specified in the ISP as well as effectiveness of such implementation as indicated by achievement of outcomes; b. Analysis of General Events Reports data; c. Compliance with Caregivers Criminal History Screening requirements; d. Compliance with Employee Abuse Registry requirements; e. Compliance with DDSD training requirements; f. Patterns in reportable incidents; and g. Results of improvement actions taken in previous quarters. 2. The Provider Agency must complete a QA/QI report annually by February 15th of each calendar year, or as otherwise requested by DOH. The report must be kept on file at the agency, made available for review by DOH, and upon request from DDSD the report must be submitted to the relevant DDSD Regional Offices. The report will summarize: a. Sufficiency of staff coverage; b. Effectiveness and timeliness of implementation of ISPs, including trends in achievement of individual desired outcomes; c. Results of General Events Reporting data analysis, Trends in Category II significant events; d. Patterns in medication errors; e. Action taken regarding individual grievances; f. Presence and completeness of required documentation; g. A description of how data collected as part of the agency's QA/QI plan was used, what quality improvement initiatives were			

undertaken, and the results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QI process; and h. Significant program changes. **CHAPTER 13 (IMLS) 3. Service Requirements: F. Quality Assurance/Quality Improvement (QA/QI) Program:** Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QI activities. 1. **Development of a QI plan:** The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working. 2. **Implementing a QA/QI Committee:** The QA/QI committee shall convene on at least on a quarterly basis and as needed to review service reports, to identify any deficiencies, trends, patterns or concerns, as well as opportunities for quality improvement. For Intensive Medical Living providers, at least one nurse shall be a member of this committee. The QA meeting			

shall be documented. The QA review should address at least the following: a. Implementation of the ISPs, including the extent to which services are delivered in accordance with the ISPs and associated support plans and /or WDSI including the type, scope, amount, duration, and frequency specified in the ISPs as well as effectiveness of such implementation as indicated by achievement of outcomes; b. Trends in General Events as defined by DDSD; c. Compliance with Caregivers Criminal History Screening Requirements; d. Compliance with DDSD training requirements; e. Trends in reportable incidents; and f. Results of improvement actions taken in previous quarters. 3. The Provider Agency must complete a QA/QI report annually by February 15th of each calendar year, or as otherwise requested by DOH. The report must be kept on file at the agency, made available for review by DOH and upon request from DDSD; the report must be submitted to the relevant DDSD Regional Offices. The report will summarizes: a. Sufficiency of staff coverage; b. Effectiveness and timeliness of implementation of ISPs and associated Support plans and/or WDSI including trends in achievement of individual desired outcomes; c. Trends in reportable incidents; d. Trends in medication errors; e. Action taken regarding individual grievances; f. Presence and completeness of required documentation; g. How data collected as part of the agency's QA/QI was used, what quality improvement initiatives were undertaken, and what were the results of those efforts, including			

discovery and remediation of any service delivery deficiencies discovered through the QI process; and h. Significant program changes. **CHAPTER 14 (ANS) 3. Service Requirements: N. Quality Assurance/Quality Improvement (QA/QI) Program:** Agencies must develop and maintain an active QA/QI program in order to assure the provision of quality services. This includes the development of a QA/QI plan, data gathering and analysis, and routine meetings to analyze the results of QI activities. 1. **Development of a QI plan:** The quality management plan is used by an agency to continually determine whether the agency is performing within program requirements, achieving desired outcomes and identifying opportunities for improvement. The quality management plan describes the process the Provider Agency uses in each phase of the process: discovery, remediation and improvement. It describes the frequency, the source and types of information gathered, as well as the methods used to analyze and measure performance. The quality management plan should describe how the data collected will be used to improve the delivery of services and methods to evaluate whether implementation of improvements are working. 2. **Implementing a QA/QI Committee:** The QA/QI committee shall convene on at least on a quarterly basis and as needed to review service reports, to identify any deficiencies, trends, patterns or concerns, as well as opportunities for quality improvement. For Intensive Medical Living providers, at least one nurse shall be a member of this committee. The QA meeting shall be documented. The QA review should address at least the following:			

a. Trends in General Events as defined by DDSD; b. Compliance with Caregivers Criminal History Screening Requirements; c. Compliance with DDSD training requirements; d. Trends in reportable incidents; and e. Results of improvement actions taken in previous quarters. 3. The Provider Agency must complete a QA/QI report annually by February 15th of each calendar year, or as otherwise requested by DOH. The report must be kept on file at the agency, made available for review by DOH and upon request from DDSD; the report must be submitted to the relevant DDSD Regional Offices. The report will summarizes: a. Sufficiency of staff coverage; b. Trends in reportable incidents; c. Trends in medication errors; d. Action taken regarding individual grievances; e. Presence and completeness of required documentation; f. How data collected as part of the agency's QA/QI was used, what quality improvement initiatives were undertaken, and what were the results of those efforts, including discovery and remediation of any service delivery deficiencies discovered through the QI process; and g. Significant program changes **NMAC 7.1.14.8 INCIDENT MANAGEMENT SYSTEM REPORTING REQUIREMENTS FOR COMMUNITY-BASED SERVICE PROVIDERS:** **F. Quality assurance/quality improvement program for community-based service providers:** The community-based service provider shall establish and implement a quality improvement program for reviewing alleged			

complaints and incidents of abuse, neglect, or exploitation against them as a provider after the division's investigation is complete. The incident management program shall include written documentation of corrective actions taken. The community-based service provider shall take all reasonable steps to prevent further incidents. The community-based service provider shall provide the following internal monitoring and facilitating quality improvement program: **(1)** community-based service providers shall have current abuse, neglect, and exploitation management policy and procedures in place that comply with the department's requirements; **(2)** community-based service providers providing intellectual and developmental disabilities services must have a designated incident management coordinator in place; and **(3)** community-based service providers providing intellectual and developmental disabilities services must have an incident management committee to identify any deficiencies, trends, patterns, or concerns as well as opportunities for quality improvement, address internal and external incident reports for the purpose of examining internal root causes, and to take action on identified issues.			

Tag # 1A09.1 Medication Delivery PRN Medication Administration	Standard Level Deficiency		
NMAC 16.19.11.8 MINIMUM STANDARDS: **A. MINIMUM STANDARDS FOR THE DISTRIBUTION, STORAGE, HANDLING AND RECORD KEEPING OF DRUGS:** (d) The facility shall have a Medication Administration Record (MAR) documenting medication administered to residents, **including over-the-counter medications.** This documentation shall include: (i) Name of resident; (ii) Date given; (iii) Drug product name; (iv) Dosage and form; (v) Strength of drug; (vi) Route of administration; (vii) How often medication is to be taken; (viii) Time taken and staff initials; (ix) Dates when the medication is discontinued or changed; (x) The name and initials of all staff administering medications. **Model Custodial Procedure Manual** ***D. Administration of Drugs*** Unless otherwise stated by practitioner, patients will not be allowed to administer their own medications. Document the practitioner's order authorizing the self-administration of medications. All PRN (As needed) medications shall have complete detail instructions regarding the administering of the medication. This shall include: ➢ symptoms that indicate the use of the medication, ➢ exact dosage to be used, and ➢ the exact amount to be used in a 24-hour period.	Medication Administration Records (MAR) were reviewed for the months of May, and June, 2016. Based on record review, 1 of 6 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #5 May 2016 No evidence of documented Signs/Symptoms were found for the following PRN medication: • Lorazepam 2mg – PRN – 5/8 (given 1 time) No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Lorazepam 2mg – PRN – 5/8 (given 1 time) June 2016 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Pepto-Bismol 30ml – PRN – 6/12 (given 1 time)	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)*: → **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)*: →	

Department of Health Developmental Disabilities Supports Division (DDSD) Medication Assessment and Delivery Policy - Eff. November 1, 2006 **F. PRN Medication** 3. Prior to self-administration, self-administration with physical assist or assisting with delivery of PRN medications, the direct support staff must contact the agency nurse to describe observed symptoms and thus assure that the PRN medication is being used according to instructions given by the ordering PCP. In cases of fever, respiratory distress (including coughing), severe pain, vomiting, diarrhea, change in responsiveness/level of consciousness, the nurse must strongly consider the need to conduct a face-to-face assessment to assure that the PRN does not mask a condition better treated by seeking medical attention. This does not apply to home based/family living settings where the provider is related by affinity or by consanguinity to the individual. 4. The agency nurse shall review the utilization of PRN medications routinely. Frequent or escalating use of PRN medications must be reported to the PCP and discussed by the Interdisciplinary for changes to the overall support plan (see Section H of this policy). **H. Agency Nurse Monitoring** 1. Regardless of the level of assistance with medication delivery that is required by the individual or the route through which the medication is delivered, the agency nurses must monitor the individual's response to the effects of their routine and PRN medications. The frequency and type of monitoring must be based on the nurse's assessment of the individual and consideration of the individual's diagnoses, health status, stability, utilization of			

PRN medications and level of support required by the individual's condition and the skill level and needs of the direct care staff. Nursing monitoring should be based on prudent nursing practice and should support the safety and independence of the individual in the community setting. The health care plan shall reflect the planned monitoring of the individual's response to medication. **Department of Health Developmental Disabilities Supports Division (DDSD) - Procedure Title:** **Medication Assessment and Delivery** **Procedure Eff Date: November 1, 2006** C. 3. Prior to delivery of the PRN, direct support staff must contact the agency nurse to describe observed symptoms and thus assure that the PRN is being used according to instructions given by the ordering PCP. In cases of fever, respiratory distress (including coughing), severe pain, vomiting, diarrhea, change in responsiveness/level of consciousness, the nurse must strongly consider the need to conduct a face-to-face assessment to assure that the PRN does not mask a condition better treated by seeking medical attention. (References: Psychotropic Medication Use Policy, Section D, page 5 Use of PRN Psychotropic Medications; and, Human Rights Committee Requirements Policy, Section B, page 4 Interventions Requiring Review and Approval – Use of PRN Medications). a. Document conversation with nurse including all reported signs and symptoms, advice given and action taken by staff. 4. Document on the MAR each time a PRN medication is used and describe its effect on the individual (e.g., temperature down, vomiting			

lessened, anxiety increased, the condition is the same, improved, or worsened, etc.). Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013 **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; iv. Explanation of any medication error; v. Documentation of any allergic reaction or adverse medication effect; and vi. For PRN medication, instructions for the use of the PRN medication must include observable signs/symptoms or circumstances in which the medication is to be used, and documentation of effectiveness of PRN medication administered. c. The Family Living Provider Agency must also maintain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose; and d. Information from the prescribing pharmacy regarding medications must be kept in the home and community inclusion service locations and must include the expected desired outcomes of administering the medication, signs and symptoms of adverse events and interactions with other medications. e. Medication Oversight is optional if the individual resides with their biological family (by affinity or consanguinity). If Medication			

Oversight is not selected as an Ongoing Nursing Service, all elements of medication administration and oversight are the sole responsibility of the individual and their biological family. Therefore, a monthly medication administration record (MAR) is not required unless the family requests it and continually communicates all medication changes to the provider agency in a timely manner to insure accuracy of the MAR. i. The family must communicate at least annually and as needed for significant change of condition with the agency nurse regarding the current medications and the individual's response to medications for purpose of accurately completing required nursing assessments. ii. As per the DDSD Medication Assessment and Delivery Policy and Procedure, paid DSP who are not related by affinity or consanguinity to the individual may not deliver medications to the individual unless they have completed Assisting with Medication Delivery (AWMD) training. DSP may also be under a delegation relationship with a DDW agency nurse or be a Certified Medication Aide (CMA). Where CMAs are used, the agency is responsible for maintaining compliance with New Mexico Board of Nursing requirements. iii. If the substitute care provider is a surrogate (not related by affinity or consanguinity) Medication Oversight must be selected and provided. **CHAPTER 12 (SL) 2. Service Requirements L. Training and Requirements: 3. Medication** Delivery: Supported Living Provider Agencies must have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in accordance with DDSD Medication Assessment and Delivery			

Policy and Procedures, New Mexico Nurse Practice Act, and Board of Pharmacy standards and regulations. a. All twenty-four (24) hour residential home sites serving two (2) or more unrelated individuals must be licensed by the Board of Pharmacy, per current regulations; b. When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) must be maintained and include: i. The name of the individual, a transcription of the physician's or licensed health care provider's prescription including the brand and generic name of the medication, and diagnosis for which the medication is prescribed; ii. Prescribed dosage, frequency and method/route of administration, times and dates of administration; iii. Initials of the individual administering or assisting with the medication delivery; iv. Explanation of any medication error; v. Documentation of any allergic reaction or adverse medication effect; and vi. For PRN medication, instructions for the use of the PRN medication must include observable signs/symptoms or circumstances in which the medication is to be used, and documentation of effectiveness of PRN medication administered.			

c. The Supported Living Provider Agency must also maintain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose; and d. Information from the prescribing pharmacy regarding medications must be kept in the home and community inclusion service locations and must include the expected desired outcomes of administrating the medication, signs, and symptoms of adverse events and interactions with other medications. **CHAPTER 13 (IMLS) 2. Service Requirements. B.** There must be compliance with all policy requirements for Intensive Medical Living Service Providers, including written policy and procedures regarding medication delivery and tracking and reporting of medication errors consistent with the DDSD Medication Delivery Policy and Procedures, relevant Board of Nursing Rules, and Pharmacy Board standards and regulations. Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 **CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS:** The objective of these standards is to establish Provider Agency policy, procedure and reporting requirements for DD Medicaid Waiver program. These requirements apply to all such Provider Agency staff, whether directly employed or subcontracting with the Provider Agency. Additional Provider Agency requirements and personnel qualifications may be applicable for specific service standards. **E. Medication Delivery:** Provider Agencies that provide Community Living, Community Inclusion or Private Duty Nursing services shall			

have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in accordance with DDSD Medication Assessment and Delivery Policy and Procedures, the Board of Nursing Rules and Board of Pharmacy standards and regulations. (2) When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) shall be maintained and include: (a) The name of the individual, a transcription of the physician's written or licensed health care provider's prescription including the brand and generic name of the medication, diagnosis for which the medication is prescribed; (b) Prescribed dosage, frequency and method/route of administration, times and dates of administration; (c) Initials of the individual administering or assisting with the medication; (d) Explanation of any medication irregularity; (e) Documentation of any allergic reaction or adverse medication effect; and (f) For PRN medication, an explanation for the use of the PRN medication shall include observable signs/symptoms or circumstances in which the medication is to be used, and documentation of effectiveness of PRN medication administered. (3) The Provider Agency shall also maintain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose;			

(4) MARs are not required for individuals participating in Independent Living who self-administer their own medications; (5) Information from the prescribing pharmacy regarding medications shall be kept in the home and community inclusion service locations and shall include the expected desired outcomes of administrating the medication, signs and symptoms of adverse events and interactions with other medications;			

Tag # 1A33 Board of Pharmacy – Med. Storage	Standard Level Deficiency		
New Mexico Board of Pharmacy Model Custodial Drug Procedures Manual **E. Medication Storage:** 1. Prescription drugs will be stored in a locked cabinet and the key will be in the care of the administrator or designee. 2. Drugs to be taken by mouth will be separate from all other dosage forms. 3. A locked compartment will be available in the refrigerator for those items labeled "Keep in Refrigerator." The temperature will be kept in the 36°F - 46°F range. An accurate thermometer will be kept in the refrigerator to verify temperature. 4. Separate compartments are required for each resident's medication. 5. All medication will be stored according to their individual requirement or in the absence of temperature and humidity requirements, controlled room temperature (68-77°F) and protected from light. Storage requirements are in effect 24 hours a day. 6. Medication no longer in use, unwanted, outdated, or adulterated will be placed in a quarantine area in the locked medication cabinet and held for destruction by the consultant pharmacist. **8. References** A. Adequate drug references shall be available for facility staff **H. Controlled Substances (Perpetual Count Requirement)** 1. Separate accountability or proof-of-use sheets shall be maintained, for each controlled substance, indicating the following information: a. date	Based on observation, the Agency did not to ensure proper storage of medication for 2 of 6 individuals. Observation included: Individual #1 • Zyprexa 10mg: expired 10/2015. Expired medication was not kept separate from other medications as required by Board of Pharmacy Procedures. • Ibuprofen 200mg: expired 4/2016. Expired medication was not kept separate from other medications as required by Board of Pharmacy Procedures. Individual #3 • Ondansetron: expired 9/2015. Expired medication was not kept separate from other medications as required by Board of Pharmacy Procedures.	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)*: → **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)*: →	

b. time administered c. name of patient d. dose e. practitioner's name f. signature of person administering or assisting with the administration the dose g. balance of controlled substance remaining.			

Tag # LS25 / 6L25 Residential Health and Safety (SL)	Standard Level Deficiency		
Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013 **CHAPTER 11 (FL) Living Supports – Family Living Agency Requirements G. Residence Requirements for Living Supports- Family Living Services:** 1.Family Living Services providers must assure that each individual's residence is maintained to be clean, safe and comfortable and accommodates the individuals' daily living, social and leisure activities. In addition the residence must: j. Maintain basic utilities, i.e., gas, power, water and telephone; k. Provide environmental accommodations and assistive technology devices in the residence including modifications to the bathroom (i.e., shower chairs, grab bars, walk in shower, raised toilets, etc.) based on the unique needs of the individual in consultation with the IDT; l. Have a battery operated or electric smoke detectors, carbon monoxide detectors, fire extinguisher, or a sprinkler system; m. Have a general-purpose first aid kit; n. Allow at a maximum of two (2) individuals to share, with mutual consent, a bedroom and each individual has the right to have his or her own bed; o. Have accessible written documentation of actual evacuation drills occurring at least three (3) times a year; p. Have accessible written procedures for the safe storage of all medications with	Based on observation, the Agency did not ensure that each individuals' residence met all requirements within the standard for 2 of 4 Supported Living residences. Review of the residential records and observation of the residence revealed the following items were not found, not current, not functioning or incomplete: **Supported Living Requirements:** • Water temperature in home does not exceed safe temperature (110^0 F) ➢ Water temperature in home measured 112.2^0 F (#3) • General-purpose first aid kit (#1, 6) • Accessible written procedures for the safe storage of all medications with dispensing instructions for each individual that are consistent with the Assisting with Medication Administration training or each individual's ISP (#1, 3, 6) • Accessible written procedures for emergency evacuation of individuals in the event of the residence becomes unsuitable for occupancy. The emergency evacuation procedures shall address, but are not limited to, fire, chemical and/or hazardous waste spills, and flooding (#1, 3, 6) *Note: The following Individuals share a residence:* ➢ #4, 5 ➢ #1, 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?)***:** →	

dispensing instructions for each individual that are consistent with the Assisting with Medication Delivery training or each individual's ISP; and q. Have accessible written procedures for emergency placement and relocation of individuals in the event of an emergency evacuation that makes the residence unsuitable for occupancy. The emergency evacuation procedures must address, but are not limited to, fire, chemical and/or hazardous waste spills, and flooding. **CHAPTER 12 (SL) Living Supports – Supported Living Agency Requirements G. Residence Requirements for Living Supports- Supported Living Services:** 1. Supported Living Provider Agencies must assure that each individual's residence is maintained to be clean, safe, and comfortable and accommodates the individual's daily living, social, and leisure activities. In addition the residence must: a. Maintain basic utilities, i.e., gas, power, water, and telephone; b. Provide environmental accommodations and assistive technology devices in the residence including modifications to the bathroom (i.e., shower chairs, grab bars, walk in shower, raised toilets, etc.) based on the unique needs of the individual in consultation with the IDT; c. Ensure water temperature in home does not exceed safe temperature (110^0 F) ; d. Have a battery operated or electric smoke detectors and carbon monoxide detectors, fire extinguisher, or a sprinkler system;			

e. Have a general-purpose First Aid kit; f. Allow at a maximum of two (2) individuals to share, with mutual consent, a bedroom and each individual has the right to have his or her own bed; g. Have accessible written documentation of actual evacuation drills occurring at least three (3) times a year. For Supported Living evacuation drills must occur at least once a year during each shift; h. Have accessible written procedures for the safe storage of all medications with dispensing instructions for each individual that are consistent with the Assisting with Medication Delivery training or each individual's ISP; and i. Have accessible written procedures for emergency placement and relocation of individuals in the event of an emergency evacuation that makes the residence unsuitable for occupancy. The emergency evacuation procedures must address, but are not limited to, fire, chemical and/or hazardous waste spills, and flooding. **CHAPTER 13 (IMLS) 2. Service Requirements R. Staff Qualifications: 3. Supervisor Qualifications And Requirements:** S Each residence shall include operable safety equipment, including but not limited to, an operable smoke detector or sprinkler system, a carbon monoxide detector if any natural gas appliance or heating is used, fire extinguisher, general purpose first aid kit, written procedures for emergency evacuation due to fire or other emergency and documentation of evacuation drills occurring			

at least annually during each shift, phone number for poison control within line of site of the telephone, basic utilities, general household appliances, kitchen and dining utensils, adequate food and drink for three meals per day, proper food storage, and cleaning supplies. T Each residence shall have a blood borne pathogens kit as applicable to the residents' health status, personal protection equipment, and any ordered or required medical supplies shall also be available in the home. U If not medically contraindicated, and with mutual consent, up to two (2) individuals may share a single bedroom. Each individual shall have their own bed. All bedrooms shall have doors that may be closed for privacy. Individuals have the right to decorate their bedroom in a style of their choosing consistent with safe and sanitary living conditions. V For residences with more than two (2) residents, there shall be at least two (2) bathrooms. Toilets, tubs/showers used by the individuals shall provide for privacy and be designed or adapted for the safe provision of personal care. Water temperature shall be maintained at a safe level to prevent injury and ensure comfort and shall not exceed one hundred ten (110) degrees. Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 **CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS** **L. Residence Requirements for Family Living Services and Supported Living Services**			

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Date Due

Service Domain: Medicaid Billing/Reimbursement – *State financial oversight exists to assure that claims are coded and paid for in accordance with the reimbursement methodology specified in the approved waiver.*

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

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

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

CHAPTER 12 (SL) 2. REIMBURSEMENT

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

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

Billing for ***2012:*** *Living Supports* (Supported Living); *Inclusion Supports* (Customized Community Supports) services was reviewed for 6 of 6 individuals. *Progress notes and billing records supported billing activities for the months of March, April and May 2016.*

Date: August 26, 2016

To: Patricia Tarin, Director
Provider: Campo Behavioral Health, LLC
Address: 424 N. Mesilla Street
City/State/Zip: Las Cruces, New Mexico 88005

E-mail Address: patsy@campobh.com

Region: Southwest
Survey Date: June 13 - 15, 2016
Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: ***2012:*** *Living Supports* (Supported Living); *Inclusion Supports* (Customized Community Supports)

Survey Type: Routine

RE: Request for an Informal Reconsideration of Findings

Dear Ms. Tarin,

Your request for a Reconsideration of Findings was received on 8/11/2016. Your request and the supporting evidence provided have been reviewed. Based on the review of applicable standards and regulations, review of the survey process and the evidence you provided, the following determinations have been made:

Regarding Tag # LS14 / 6L14
Determination: The IRF committee is modifying the original tag in the report of findings. You are required to complete the remainder of your Plan of Correction as previously indicated. Based on documentation provided, the finding for Individual #3, Medical Emergency Response Plan for Respiratory, will be removed. The remaining citations noted in this tag were not disputed.

This concludes the Informal Reconsideration of Finding process. The IRF process is separate and apart from the Informal Dispute Resolution process or the Medicaid Fair Hearing process when DOH sanctions are imposed on a provider.

Thank you.
Respectfully,

Crystal Lopez-Beck

Crystal Lopez-Beck, BA
Deputy Bureau Chief/QMB
Informal Reconsideration of Finding Committee Chair

Date: October 20, 2016

To: Patricia Tarin, Director
Provider: Campo Behavioral Health, LLC
Address: 424 N. Mesilla Street
City/State/Zip: Las Cruces, New Mexico 88005

E-mail Address: patsy@campobh.com

Region: Southwest
Survey Date: June 13 - 15, 2016
Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: ***2012:*** *Living Supports* (Supported Living); *Inclusion Supports* (Customized Community Supports)

Survey Type: Routine

Dear Ms. Tarin;

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

The Plan of Correction process is now complete.

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

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

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

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

Sincerely,

Amanda Castañeda

Amanda Castañeda
Plan of Correction Coordinator
Quality Management Bureau/DHI

Q.16.4.DDW.D1001.3.RTN.09.16.294