Date:	April 11, 2018
To:	Anna Marie Carrillo, Service Coordinator / Co-Director
Provider:	Carlacare, Inc.
Address:	1988 Crescent Drive
State/Zip:	Las Cruces, New Mexico 88005
E-mail Address:	carrillr0943@comcast.net
Region:	Southwest Region
Survey Date:	January 26 - 30, 2018
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	**2012:** Supported Living
Survey Type:	Routine Survey
Team Leader:	Amanda Castaneda, MPA, Plan of Correction Coordinator, Division of Health Improvement/Quality Management Bureau
Team Members:	Richard Gomez, BS, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Dear Ms. Anna Marie Carrillo;

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 # 1A22 Agency Personnel Competency
- Tag # 1A15.2 and IS09 / 5I09 Healthcare Documentation

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.

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

Corrective Action:

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

On-going Quality Assurance/Quality Improvement Processes:

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

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

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

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

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

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

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

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

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

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

Attention: Lisa Medina-Lujan
HSD/OIG
Program Integrity Unit
1474 Rodeo Road
Santa Fe, New Mexico 87505

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

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

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

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

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

Sincerely,

Amanda Castaneda, MPA

Amanda Castaneda, MPA
Team Lead/Plan of Correction Coordinator
Division of Health Improvement
Quality Management Bureau

Survey Process Employed:

Administrative Review Start Date:	January 26, 2018
Entrance Conference Date:	January 29, 2018
Present:	**Carlacare, Inc.** Richard Carrillo, Incident Management Coordinator / Co-Director Anna Marie Carrillo, Service Coordinator / Co-Director **DOH/DHI/QMB** Amanda Castaneda, MPA, Team Lead/Plan of Correction Coordinator Richard Gomez, BS, Healthcare Surveyor
Exit Conference Date:	January 30, 2018
Present:	**Carlacare, Inc.** Richard Carrillo, Incident Management Coordinator / Co-Director Anna Marie Carrillo, Service Coordinator / Co-Director **DOH/DHI/QMB** Amanda Castaneda, MPA, Team Lead/Plan of Correction Coordinator **DDSD SW Regional Office** Dave Brunson, Community Inclusion Coordinator
Administrative Locations Visited	1
Total Sample Size	1 0 - *Jackson* Class Members 1 - Non-*Jackson* Class Members 1 - Supported Living
Total Homes Visited ❖ Supported Living Homes Visited	1 1
Persons Served Records Reviewed	1
Persons Served Interviewed	1
Direct Support Personnel Interviewed	1
Direct Support Personnel Records Reviewed	3
Service Coordinator Records Reviewed	1
Administrative Interviews	2

Administrative Processes and Records Reviewed:

- Medicaid Billing/Reimbursement Records for all Services Provided
- Accreditation Records
- Oversight of Individual Funds
- Individual Medical and Program Case Files, including, but not limited to:
    - Individual Service Plans

    - Progress on Identified Outcomes
    - Healthcare Plans
    - Medication Administration Records
    - Medical Emergency Response Plans
    - Therapy Evaluations and Plans
    - Healthcare Documentation Regarding Appointments and Required Follow-Up
    - Other Required Health Information
- Internal Incident Management Reports and System Process / General Events Reports
- Personnel Files, including nursing and subcontracted staff
- Staff Training Records, Including Competency Interviews with Staff
- Agency Policy and Procedure Manual
- Caregiver Criminal History Screening Records
- Consolidated Online Registry/Employee Abuse Registry
- Human Rights Committee Notes and Meeting Minutes
- Evacuation Drills of Residences and Service Locations
- Quality Assurance / Improvement Plan

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

Attachment A

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

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

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

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

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

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

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

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

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

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

1. How the specific and realistic corrective action will be accomplished for individuals found to have been affected by the deficient practice.
2. How the agency will identify other individuals who have the potential to be affected by the same deficient practice, and how the agency will act to protect individuals in similar situations.
3. What QA measures will be put into place or systemic changes made to ensure that the deficient practice will not recur
4. Indicate how the agency plans to monitor its performance to make sure that solutions are sustained. The agency must develop a QA plan for ensuring that correction is achieved and sustained. This QA plan must be implemented, and the corrective action evaluated for its effectiveness. The plan of correction is integrated into the agency quality assurance system; and
5. Include dates when corrective action will be completed. The corrective action completion dates must be acceptable to the State.

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 Condition of Participation level deficiencies overall, as well as widespread Standard level deficiencies identified in the attached QMB Report of Findings and requires implementation of a Plan of Correction.

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: https://nmhealth.org/about/dhi/cbp/irf/
3. The written request for an IRF must specify in detail the request for reconsideration and why the finding is inaccurate.
4. The IRF request must include all supporting documentation or evidence.
5. If you have questions about the IRF process, email the IRF Chairperson, 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: **Carlacare, Inc. - Southwest Region**
Program: Developmental Disabilities Waiver
Service: **2012:** Supported Living
Survey Type: Routine Survey
Survey Date: **January 26 - 30, 2018**

Standard of Care	*Deficiencies*	Agency Plan of Correction, On-going QA/QI & Responsible Party	Date Due
Service Domain: Service Plans: ISP Implementation - *Services are delivered in accordance with the service plan, including type, scope, amount, duration and frequency specified in the service plan.*			
Tag # 1A08.1 Agency Case File - Progress Notes	**Standard Level Deficiency**		
Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015 **Chapter 5 (CIES) 3. Agency Requirements: 6. Reimbursement A. 1.** …Provider Agencies must maintain all records necessary to fully disclose the service, quality…The documentation of the billable time spent with an individual shall be kept on the written or electronic record… **Chapter 6 (CCS) 3. Agency Requirements: 4. Reimbursement A. Record Requirements 1.** …Provider Agencies must maintain all records necessary to fully disclose the service, quality…The documentation of the billable time spent with an individual shall be kept on the written or electronic record… **Chapter 7 (CIHS) 3. Agency Requirements: 4. Reimbursement A. 1.**…Provider Agencies must maintain all records necessary to fully disclose the service, quality…The documentation of the billable time spent with an individual shall be kept on the written or electronic record… **Chapter 11 (FL) 3. Agency Requirements: 4. Reimbursement A. 1.**…Provider Agencies must maintain all records necessary to fully disclose the service, quality…The documentation of the billable time spent with an individual shall be kept on the written or electronic record…	Based on record review, the Agency did not maintain progress notes and other service delivery documentation for 1 of 1 Individuals. Review of the Agency individual case files revealed the following items were not found: **Supported Living Individual Intensive Behavioral Services Notes/Daily Contact Logs:** Individual #1 October 2017 • Review of progress notes indicate separate progress notes were not kept for Individual Intensive Behavioral Support services for 10/1/2017 - 10/31/2017. November 2017 • Review of progress notes indicate separate progress notes were not kept for Individual Intensive Behavioral Support services for 11/1/2017 - 11/30/2017. December 2017 • Review of progress notes indicate separate progress notes were not kept for Individual Intensive Behavioral Support services for 12/1/2017 - 12/31/2017.	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?): →* **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): →*	

Chapter 12 (SL) 3. Agency Requirements: 2. Reimbursement A. 1. Provider Agencies must maintain all records necessary to fully disclose the service, quality…The documentation of the billable time spent with an individual shall be kept on the written or electronic record… **Chapter 13 (IMLS) 3. Agency Requirements: 4. Reimbursement A. 1.**…Provider Agencies must maintain all records necessary to fully disclose the service, quality…The documentation of the billable time spent with an individual shall be kept on the written or electronic record… **Chapter 15 (ANS) 4. Reimbursement A. 1.** …Provider Agencies must maintain all records necessary to fully disclose the service, quality…The documentation of the billable time spent with an individual shall be kept on the written or electronic record… 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: (3) Progress notes and other service delivery documentation;			

Tag # LS14 / 6L14 Residential Case File	Standard Level Deficiency		
Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015 **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;	Based on record review, the Agency did not maintain a complete and confidential case file in the residence for 1 of 1 Individual receiving Supported Living Services. Review of the residential individual case files revealed the following items were not found, incomplete, and/or not current: **Healthcare Passport:** • Not found (#1) **Comprehensive Risk Management Plan:** • Not Found (#1) **Health Care Plans:** • Fluid Restriction (#1) • Hydration (#1) • Intake/Output Monitoring (#1) • Communication/Vision/Hearing (#1) • Observed/Reported Expressions of Pain (#1) • Pain Medication (#1)	**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. Record of medical and dental appointments for the current year, or during the period of stay for short term stays, including any treatment provided; i. Progress notes written by DSP and nurses; j. Documentation and data collection related to ISP implementation; k. Medicaid card; l. Salud membership card or Medicare card as applicable; and m. A Do Not Resuscitate (DNR) document and/or Advanced Directives as applicable. **DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION (DDSD): Director's Release: Consumer Record Requirements eff. 11/1/2012** **III. Requirement Amendments(s) or Clarifications:** A. All case management, living supports, customized in-home supports, community integrated employment and customized community supports providers must maintain records for individuals served through DD Waiver in accordance with the Individual Case File Matrix incorporated in this director's release. H. Readily accessible electronic records are accessible, including those stored through the Therap web-based system. 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; (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 practitioner's 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 & 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 # 1A22 Agency Personnel Competency	**Condition of Participation 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) for each individual serviced. Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015 **CHAPTER 5 (CIES) 3. Agency Requirements G. Training Requirements: 1.** All Community Inclusion Providers must provide staff training in accordance with the DDSD policy T-003: Training Requirements for Direct Service Agency Staff Policy. 3. Ensure direct service personnel receives Individual Specific Training as outlined in each individual ISP, including aspects of support plans (healthcare and behavioral) or WDSI that pertain to the employment environment. **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	After an analysis of the evidence there is a significant potential for a negative outcome to occur. Based on interviews, the Agency did not ensure training competencies were met for 1 of 1 Direct Support Personnel. **When DSP were asked if they received training on the individual's Health Care Plans and if so, what the plan(s) covered, the following was reported:** • DSP #502 stated, "Yes she does, call mom." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual requires Health Care Plans for Fluid Restriction, Supports for Hydration or Risk of Dehydration, Intake and Output monitoring, Aspiration Risk, Signs/Symptoms of Reflux, Constipation Management, Communication/Vision/Hearing, Observed or Reported Expressions of Pain, and Pain Medication. (Individual #1) **When DSP were asked if they received training on the individual's Medical Emergency Response Plans and if so, what the plan(s) covered, the following was reported:** • DSP #502 stated, "I don't know to be quite honest with you." As indicated by the Electronic Comprehensive Health Assessment Tool, the Individual requires Medical Emergency Response Plans for	**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?): →*	

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. 3. Staff shall complete individual specific training requirements in accordance with the specifications described in the ISP of each individual served; and 4. Staff that assists the individual with medication (e.g., setting up medication, or reminders) must have completed Assisting with Medication Delivery (AWMD) Training. **CHAPTER 11 (FL) 3. Agency Requirements B. Living Supports- Family Living Services Provider Agency Staffing Requirements: 3. Training:** A. All Family Living Provider agencies must ensure staff training in accordance with the Training Requirements for Direct Service Agency Staff policy. DSP's or subcontractors delivering substitute care under Family Living must at a minimum comply with the section of the training policy that relates to Respite, Substitute Care, and personal support staff [Policy T-003: for Training Requirements for Direct Service Agency Staff; Sec. II-J, Items 1-4]. Pursuant to the Centers for Medicare and Medicaid Services (CMS) requirements, the services that a provider renders may only be claimed for federal match if the provider has completed all necessary training required by the state. All Family Living Provider agencies must report required personnel training status to the DDSD Statewide Training Database as specified in DDSD Policy T-001: Reporting and	Aspiration Risk and Signs/Symptoms of Reflux. (Individual #1)		

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

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

Tag # 1A28.1 Incident Mgt. System - Personnel Training	Standard 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 training, in accordance	Based on interview, the Agency did not ensure Incident Management Training for 1 of 1 Agency Personnel. **When Direct Support Personnel were asked what State agency must be contacted when there is suspected abuse, neglect or exploitation, the following was reported:** • DSP #502 stated, "I usually call parents for everything." 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?): →*	

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 # 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 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.	Based on record review and interview, the Agency did not follow the General Events Reporting requirements as indicated by the policy. **Agency record review revealed the following:** • During record review, multiple internal incident reports were found which were required to be entered into the General Events Reporting System, including the use of PRN medications. **When Surveyors asked the Agency administrative personnel to explain and show how they utilize the General Events Reporting System the following was reported:** • #504 stated, "We don't use GER."	**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?): →*	

Standard of Care	*Deficiencies*	Agency Plan of Correction, On-going QA/QI & 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 # 1A09 Medication Delivery Routine 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 December 2017 and January 2018. Based on record review, 1 of 1 individual had Medication Administration Records (MAR), which contained missing medications entries and/or other errors: Individual #1 January 2018 • As indicated by the Medication Administration Records the individual is to take Coenzyme 100mg (1 time daily). According to the Medication Label, Coenzyme 200mg is to be taken 1 time daily. Medication Administration Record and Medication Label do not match. • As indicated by the Medication Administration Records the individual is to take Vitamin D3 5,000 IU (1 time daily). According to the Medication Label, Vitamin D3 2,000 IU is to be taken 1 time daily. Medication Administration Record and Medication Label do not match.	**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; 6/15/2015 **CHAPTER 5 (CIES) 1. Scope of Service B. Self Employment 8.** Providing assistance with medication delivery as outlined in the ISP; **C. Individual Community Integrated Employment 3.** Providing assistance with medication delivery as outlined in the ISP; **D. Group Community Integrated Employment 4.** Providing assistance with medication delivery as outlined in the ISP; and **B. Community Integrated Employment Agency Staffing Requirements: o.** Comply with DDSD Medication Assessment and Delivery Policy and Procedures; **CHAPTER 6 (CCS) 1. Scope of Services A. Individualized Customized Community Supports 19.** Providing assistance or supports with medications in accordance with DDSD Medication Assessment and Delivery policy. **C. Small Group Customized Community Supports 19.** Providing assistance or supports with medications in accordance with DDSD Medication Assessment and Delivery policy. **D. Group Customized Community Supports 19.** Providing assistance or supports with medications in accordance with DDSD Medication Assessment and Delivery policy. **CHAPTER 11 (FL) 1 SCOPE OF SERVICES A. Living Supports- Family Living Services:** The scope of Family Living Services includes, but is not limited to the following as identified by the Interdisciplinary Team (IDT): 19. Assisting in medication delivery, and related monitoring, in accordance with the DDSD's Medication Assessment and Delivery Policy, New Mexico Nurse Practice Act, and Board of Pharmacy regulations including skill development activities leading to the ability for individuals to self-administer medication as appropriate; and **I. Healthcare Requirements for Family Living. 3. B.** Adult Nursing Services for medication oversight			

are required for all surrogate Living Supports-Family Living direct support personnel if the individual has regularly scheduled medication. Adult Nursing services for medication oversight are required for all surrogate Family Living Direct Support Personnel (including substitute care), if the individual has regularly scheduled medication. 6. Support Living- Family Living Provider Agencies must have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in accordance with DDSD Medication Assessment and Delivery Policy and Procedures, the New Mexico Nurse Practice Act and Board of Pharmacy standards and regulations. a. All twenty-four (24) hour residential home sites serving two (2) or more unrelated individuals must be licensed by the Board of Pharmacy, per current regulations; b. When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) must be maintained and include: i. The name of the individual, a transcription of the physician's or licensed health care provider's prescription including the brand and generic name of the medication, and 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 K. Training and Requirements: 3.** 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. When PRN medications are used, there must be clear documentation that the DSP contacted the agency nurse prior to assisting with the medication. d. 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 e. 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: 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. (1) All twenty-four (24) hour residential home sites serving two (2) or more unrelated individuals shall be licensed by the Board of Pharmacy, per current 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 # 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 December 2017 and January 2018. Based on record review, 1 of 1 individual had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #1 December 2017 Medication Administration Records did not contain the exact amount to be used in a 24 - hour period: • Tylenol 325mg (PRN)	**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; 6/15/2015 **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. f. All twenty-four (24) hour residential home sites serving two (2) or more unrelated individuals must be licensed by the Board of Pharmacy, per current regulations; g. When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) must be maintained and include:			

i. The name of the individual, a transcription of the physician's or licensed health care provider's prescription including the brand and generic name of the medication, and diagnosis for which the medication is prescribed; ii. Prescribed dosage, frequency and method/route of administration, times and dates of administration; iii. Initials of the individual administering or assisting with the medication delivery; iv. Explanation of any medication error; v. Documentation of any allergic reaction or adverse medication effect; and vi. For PRN medication, instructions for the use of the PRN medication must include observable signs/symptoms or circumstances in which the medication is to be used, and documentation of effectiveness of PRN medication administered. h. The Family Living Provider Agency must also maintain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose; and i. 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. j. 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. iv. 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. v. As per the DDSD Medication Assessment and Delivery Policy and Procedure, paid DSP who are not related by affinity or consanguinity to the individual may not deliver medications to the individual unless they have completed Assisting with Medication Delivery (AWMD) training. DSP may also be under a delegation relationship with a DDW agency nurse or be a Certified Medication Aide (CMA). Where CMAs are used, the agency is responsible for maintaining compliance with New Mexico Board of Nursing requirements. vi. If the substitute care provider is a surrogate (not related by affinity or consanguinity) Medication Oversight must be selected and provided. **CHAPTER 12 (SL) 2. Service Requirements K. Training and Requirements: 3.** 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 # 1A15.2 and IS09 / 5I09 Healthcare Documentation	Condition of Participation Level Deficiency		
Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015 **Chapter 5 (CIES) 3. Agency Requirements 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) 2. Service Requirements. E.** The agency nurse(s) for Customized Community Supports providers must provide the following services: 1. Implementation of pertinent PCP orders; ongoing oversight and monitoring of the individual's health status and medically related supports when receiving this service; **3. Agency Requirements: 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.	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, the Agency did not maintain a complete and confidential case file at the administrative office for 1 of 1 individual. Review of the Agency individual case files revealed the following items were not found, incomplete, and/or not current: **Health Care Plans:** • *Communication/Vision/Hearing* ◦ Individual #1 - According to Electronic Comprehensive Health Assessment Tool the individual is required to have a plan. No evidence of a plan found. • *Fluid Restriction* ◦ Individual #1 - According to Electronic Comprehensive Health Assessment Tool the individual is required to have a plan. No evidence of a plan found. • *Supports for Hydration or Risk of Dehydration* ◦ Individual #1 - According to Electronic Comprehensive Health Assessment Tool the individual is required to have a plan. No evidence of a plan found. • *Intake and Output Monitoring* ◦ Individual #1 - According to Electronic Comprehensive Health Assessment Tool the individual is required to have a plan. No evidence of a plan found. • *Observed or Reported Expressions of Pain*	**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. Health Care Requirements for Family Living: 5. A nurse employed or contracted by the Family Living Supports provider must complete the e-CHAT, the Aspiration Risk Screening Tool, (ARST), and the Medication Administration Assessment Tool (MAAT) and any other assessments deemed appropriate on at least an annual basis for each individual served, upon significant change of clinical condition and upon return from any hospitalizations. In addition, the MAAT must be updated for any significant change of medication regime, change of route that requires delivery by licensed or certified staff, or when an individual has completed training designed to improve their skills to support self-administration. a. For newly-allocated or admitted individuals, assessments are required to be completed within three (3) business days of admission or two (2) weeks following the initial ISP meeting, whichever comes first. b. For individuals already in services, the required assessments are to be completed no more than forty-five (45) calendar days and at least fourteen (14) calendar days prior to the annual ISP meeting. c. Assessments must be updated within three (3) business days following any significant change of clinical condition and within three (3) business days following return from hospitalization. **d.** Other nursing assessments conducted to determine current health status or to evaluate a change in clinical condition must be documented in a signed progress note that includes time and date as well as subjective information including the individual complaints, signs and symptoms noted by staff, family members or other team	° Individual #1 - According to Electronic Comprehensive Health Assessment Tool the individual is required to have a plan. No evidence of a plan found.		

members; objective information including vital signs, physical examination, weight, and other pertinent data for the given situation (e.g., seizure frequency, method in which temperature taken); assessment of the clinical status, and plan of action addressing relevant aspects of all active health problems and follow up on any recommendations of medical consultants. **e.** Develop any urgently needed interim Healthcare Plans or MERPs per DDSD policy pending authorization of ongoing Adult Nursing services as indicated by health status and individual/guardian choice. **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. **2. Service Requirements. L. Training and Requirements. 5. Health Related Documentation:** For each individual receiving Living Supports- Supported Living, the provider agency must ensure and document the following: a. That an individual with chronic condition(s) with the potential to exacerbate into a life threatening condition, has a MERP developed by a licensed nurse or other appropriate professional according to the DDSD Medical Emergency Response Plan Policy, that DSP have been trained to implement such plan(s), and ensure that a copy of such plan(s) are readily available to DSP in the home;			

b. That an average of five (5) hours of documented nutritional counseling is available annually, if recommended by the IDT and clinically indicated; c. That the nurse has completed legible and signed progress notes with date and time indicated that describe all interventions or interactions conducted with individuals served, as well as all interactions with other healthcare providers serving the individual. All interactions must be documented whether they occur by phone or in person; and d. Document for each individual that: i. The individual has a Primary Care Provider (PCP); ii. The individual receives an annual physical examination and other examinations as specified by a PCP; iii. The individual receives annual dental check-ups and other check-ups as specified by a licensed dentist; iv. The individual receives a hearing test as specified by a licensed audiologist; v. The individual receives eye examinations as specified by a licensed optometrist or ophthalmologist; and vi. Agency activities occur as required for follow-up activities to medical appointments (e.g. treatment, visits to specialists, and changes in medication or daily routine). vii. The agency nurse will provide the individual's team with a semi-annual nursing report that discusses the services provided			

and the status of the individual in the last six (6) months. This may be provided electronically or in paper format to the team no later than (2) weeks prior to the ISP and semi-annually. **f.** The Supported Living Provider Agency must ensure that activities conducted by agency nurses comply with the roles and responsibilities identified in these standards. **Chapter 13 (IMLS) 2. Service Requirements:** C. Documents to be maintained in the agency administrative office, include: A. All assessments completed by the agency nurse, including the Intensive Medical Living Eligibility Parameters tool; for e-CHAT a printed copy of the current e-CHAT summary report shall suffice; F. Annual physical exams and annual dental exams (not applicable for short term stays); G. Tri-annual vision exam (Not applicable for short term stays. See Medicaid policy 8.310.6 for allowable exceptions for more frequent vision exam); H. Audiology/hearing exam as applicable (Not applicable for short term stays; See Medicaid policy 8.324.6 for applicable requirements); I. All other evaluations called for in the ISP for which the Services provider is responsible to arrange; J. Medical screening, tests and lab results (for short term stays, only those which occur during the period of the stay); L. Record of medical and dental appointments, including any treatment provided (for short term stays, only those appointments that occur during the stay); O. Semi-annual ISP progress reports and MERP reviews (not applicable for short term stays); P. Quarterly nursing summary reports (not applicable for short term stays);			

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. **Department of Health Developmental Disabilities Supports Division Policy. Medical Emergency Response Plan Policy MERP-001 eff.8/1/2010** F. The MERP shall be written in clear, jargon free language and include at a minimum the following information: 1. A brief, simple description of the condition or illness. 2. A brief description of the most likely life threatening complications that might occur and what those complications may look like to an observer. 3. A concise list of the most important measures that may prevent the life threatening complication from occurring (e.g., avoiding allergens that trigger an asthma attack or making sure the person with diabetes has snacks with them to avoid hypoglycemia). 4. Clear, jargon free, step-by-step instructions regarding the actions to be taken by direct support personnel (DSP) and/or others to intervene in the emergency, including criteria for when to call 911. 5. Emergency contacts with phone numbers. 6. Reference to whether the individual has advance directives or not, and if so, where the advance directives are located.			

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)**			
NMAC 8.302.1.17 Effective Date 9-15-08 **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. **Detail Required in Records -** Provider Records must be sufficiently detailed to substantiate the date, time, eligible recipient name, rendering, attending, ordering or prescribing provider; level and quantity of services, length of a session of service billed, diagnosis and medical necessity of any service . . . Treatment plans or other plans of care must be sufficiently detailed to substantiate the level of need, supervision, and direction and service(s) needed by the eligible recipient. **Services Billed by Units of Time -** Services billed on the basis of time units spent with an eligible recipient must be sufficiently detailed to document the actual time spent with the eligible recipient and the services provided during that time unit. **Records Retention -** A provider who receives payment for treatment, services or goods must retain all medical and business records relating to any of the following for a period of at least six years from the payment date: (1) treatment or care of any eligible recipient (2) services or goods provided to any eligible recipient (3) amounts paid by MAD on behalf of any eligible recipient; and (4) any records required by MAD for the administration of Medicaid. Billing for ***2012:*** *Living Supports* (Supported Living) services was reviewed for 1 of 1 individuals. *Progress notes and billing records supported billing activities for the months of October, November, and December 2018.*			

Date: May 24, 2018

To: Anna Marie Carrillo, Service Coordinator / Co-Director
Provider: Carlacare, Inc.
Address: 1988 Crescent Drive
State/Zip: Las Cruces, New Mexico 88005

E-mail Address: carrillr0943@comcast.net

Region: Southwest Region
Survey Date: January 26 - 30, 2018
Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: **2012:** Supported Living

Survey Type: Routine Survey

Team Leader: Amanda Castaneda, MPA, Plan of Correction Coordinator, Division of Health Improvement/Quality Management Bureau

Team Members: Richard Gomez, BS, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Dear Ms. Anna Marie Carrillo;

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.18.3.DDW.D3656.3.RTN.09.18.144