

Date: April 11, 2018

To: Damian Houfek, President, Chief Executive Officer
 Provider: ENMRSH, Inc.
 Address: 2700 East 7th Street
 State/Zip: Clovis, New Mexico 88101

E-mail Address: dhoufek@enmrsh.org

CC: Cathy Mills, Board Chair
 Board Chair
 Address: 2609 Putnam
 Clovis, New Mexico 88101

Region: Southeast
 Survey Date: January 26 – February 1, 2018
 Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: **2012:** Supported Living, Family Living, Intensive Medical Living, Customized Community Supports, Community Integrated Employment Services and Customized In-Home Supports

Survey Type: Routine

Team Leader: Deb Russell, BS, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Team Members: Michelle Beck, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; Tony Fragua, Program Manager, Division of Health Improvement/Quality Management Bureau; Kandis Gomez, AA; Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; Monica Valdez, BS, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau; Nick Gomez, BS, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Dear Mr. Houfek;

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:

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

QMB Report of Findings – ENMRSH, Inc. – Southeast – January 26 – February 1, 2018

Survey Report #: Q.18.3.DDW.D1808.4.RTN.01.18.101



Compliance with all Conditions of Participation.

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

Plan of Correction:

The attached Report of Findings identifies the Standard Level and/or Condition of Participation deficiencies found during your agency's compliance review. You are required to complete and implement a Plan of Correction. Your agency has a total of 45 business days (10 business days to submit your POC for approval and 35 days to implement your *approved* Plan of Correction) from the receipt of this letter.

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

Corrective Action:

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

On-going Quality Assurance/Quality Improvement Processes:

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

Submission of your Plan of Correction:

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

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

1. **Quality Management Bureau, Attention: Amanda Castaneda, Plan of Correction Coordinator
1170 North Solano Suite D Las Cruces, New Mexico 88001**
2. **Developmental Disabilities Supports Division Regional Office for region of service surveyed**

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

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

Billing Deficiencies:

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

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

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

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

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

Request for Informal Reconsideration of Findings (IRF):

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

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

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

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

Sincerely,

Deb Russell, BS

Deb Russell, BS
Team Lead/Healthcare Surveyor
Division of Health Improvement
Quality Management Bureau

Survey Process Employed:

Administrative Review Start Date: January 26, 2018

Contact: **ENMRSH, Inc.**
Celeste Childers, Director of Quality Development

DOH/DHI/QMB
Deb Russell, BS, Team Lead/Healthcare Surveyor

On-site Entrance Conference Date: January 29, 2018

Present: **ENMRSH, Inc.**
Celeste Childers, Director of Quality Development
Damian Houfek, President, Chief Executive Officer

DOH/DHI/QMB
Deb Russell, Team Lead/Healthcare Surveyor
Tony Fragua, BFA, Program Manager
Michelle Beck, Healthcare Surveyor
Kandis Gomez, AA, Healthcare Surveyor
Monica Valdez, BS, Healthcare Surveyor
Nick Gomez, BS, Healthcare Surveyor

Exit Conference Date: February 1, 2018

Present: **ENMRSH, Inc.**
Celeste Childers, Director of Quality Development
Damian Houfek, President, Chief Executive Officer
Kathy Lynch, RN, Director of Nursing
Janelle Moore, Director of Accounting
Barbara Marion, Vice President, Chief Financial Officer
Valerie Dewbre, Lead Program Manager
Liz Gallegos, Director of Community Living
Theresa Musick, Director of Technology
Christina Chavez, Director of Adult Habilitation
Tony Marion, Director of Client Services

DOH/DHI/QMB
Deb Russell, Team Lead/Healthcare Surveyor
Tony Fragua, BFA, Program Manager
Michelle Beck, Healthcare Surveyor
Kandis Gomez, AA, Healthcare Surveyor
Monica Valdez, BS, Healthcare Surveyor

DDSD – Southeast Regional Office
Juana Bravo, Litigation Management Bureau

Administrative Locations Visited Number: 1

Total Sample Size Number: 20

0 - Jackson Class Members
20- Non-Jackson Class Members

12- Supported Living
1 - Family Living

15 - Customized Community Supports
 7 - Community Integrated Employment Services
 6 - Customized In-Home Supports

Total Homes Visited	Number:	12
❖ Supported Living Homes Visited	Number:	11
		<i>Note: The following Individuals share a SL residence:</i>
		➤ #5, 17
❖ Family Living Homes Visited	Number:	1
Persons Served Records Reviewed	Number:	20
Persons Served Interviewed	Number:	13
Persons Served Observed	Number:	1 (One Individual chose not to participate)
Persons Served Not Seen and/or Not Available	Number:	6
Direct Support Personnel Interviewed	Number:	22 (One DSP also performs dual roles as a Service Coordinator)
Direct Support Personnel Records Reviewed	Number:	130 (One DSP also performs dual roles as a Service Coordinator)
Substitute Care/Respite Personnel Records Reviewed	Number:	3
Service Coordinator Records Reviewed	Number:	5 (One SC also performs dual roles as a DSP)
Administrative Interviews	Number:	1

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

- 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 DDS 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 DDS.

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.

**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: ENMRSH, Inc. - Southeast Region
Program: Developmental Disabilities Waiver
Service: **2012: Living Supports** (Supported Living, Family Living); **Inclusion Supports** (Customized Community Supports, Community Integrated Employment Services) and **Other** (Customized In-Home Supports)
Survey Date: January 26 – February 1, 2018

Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Date Due
Service Domain: Service Plans: ISP Implementation – Services are delivered in accordance with the service plan, including type, scope, amount, duration and frequency specified in the service plan.			
Tag # 1A32 and LS14 / 6L14 Individual Service Plan Implementation	Standard Level Deficiency		
<p>NMAC 7.26.5.16.C and D Development of the ISP. Implementation of the ISP. The ISP shall be implemented according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan.</p> <p>C. The IDT shall review and discuss information and recommendations with the individual, with the goal of supporting the individual in attaining desired outcomes. The IDT develops an ISP based upon the individual's personal vision statement, strengths, needs, interests and preferences. The ISP is a dynamic document, revised periodically, as needed, and amended to reflect progress towards personal goals and achievements consistent with the individual's future vision. This regulation is consistent with standards established for individual plan development as set forth by the commission on the accreditation of rehabilitation facilities (CARF) and/or other program accreditation approved and adopted by the developmental disabilities division and the department of health. It is the policy of the developmental disabilities division (DDD), that to the extent permitted by funding, each individual receive supports and services that will assist and encourage independence and productivity in the community</p>	<p>Based on record review, the Agency did not implement the ISP according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan for 2 of 20 individuals.</p> <p>As indicated by Individuals ISP the following was found with regards to the implementation of ISP Outcomes:</p> <p>Administrative Files Reviewed:</p> <p>Customized Community Supports Data Collection/Data Tracking/Progress with regards to ISP Outcomes:</p> <p>Individual #15</p> <ul style="list-style-type: none"> • According to the Fun Outcome; Action Step for "...will choose a game" is to be completed 2 times per month, evidence found indicated it was not being completed at the required frequency as indicated in the ISP for 12/2017. • According to the Fun Outcome; Action Step for "...will take the lead in gaming event" is to be completed 2 times per month, evidence found indicated it was not being completed at the required frequency as 	<p>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?): →</p> <p>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): →</p>	

<p>and attempt to prevent regression or loss of current capabilities. Services and supports include specialized and/or generic services, training, education and/or treatment as determined by the IDT and documented in the ISP.</p> <p>D. The intent is to provide choice and obtain opportunities for individuals to live, work and play with full participation in their communities. The following principles provide direction and purpose in planning for individuals with developmental disabilities. [05/03/94; 01/15/97; Recompiled 10/31/01]</p>	<p>indicated in the ISP for 12/2017.</p> <p>Residential Files Reviewed:</p> <p>Family Living Data Collection/Data Tracking/Progress with regards to ISP Outcomes:</p> <p>Individual #12</p> <ul style="list-style-type: none"> • None found regarding: Fun Outcome/Action Step: "...will utilize different strategies to save money" for 1/1 - 31, 2018. Action step is to be completed 1 time per month. • None found regarding: Fun Outcome/Action Step: "...will plan for her trip to Utah" for 1/1 – 31, 2018. Action step is to be completed 1 time per month. 		
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Tag # LS14 / 6L14 Residential Case File	Standard Level Deficiency		
<p>Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015</p> <p>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 DDS Individual Case File Matrix policy.</p> <p>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 DDS Individual Case File Matrix policy.</p> <p>CHAPTER 13 (IMLS) 2. Service Requirements B.1. Documents to Be Maintained in The Home:</p> <ol style="list-style-type: none"> 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; Personal identification; 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; Dated and signed consent to release information forms as applicable; Current orders from health care practitioners; Documentation and maintenance of accurate medical history in Therap website; Medication Administration Records for the current month; Record of medical and dental appointments for the current year, or during the period of stay for short term stays, including any treatment 	<p>Based on record review, the Agency did not maintain a complete and confidential case file in the residence for 2 of 13 Individuals receiving Family Living Services and Supported Living Services.</p> <p>Review of the residential individual case files revealed the following items were not found, incomplete, and/or not current:</p> <p>Speech Therapy Plan:</p> <ul style="list-style-type: none"> ◦ Not Found (#12) <p>Special Health Care Needs</p> <ul style="list-style-type: none"> ◦ Comprehensive Aspiration Risk Management Plan: <ul style="list-style-type: none"> ➢ Not Current (#14) 	<p>Provider: State your Plan of Correction for the deficiencies cited in this tag here <i>(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?):</i> →</p> <p>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here <i>(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> →</p>	<p> </p>

<p>provided;</p> <ul style="list-style-type: none"> 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. <p>DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION (DDSD): Director's Release: Consumer Record Requirements eff. 11/1/2012</p> <p>III. Requirement Amendments(s) or Clarifications:</p> <p>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.</p> <p>H. Readily accessible electronic records are accessible, including those stored through the Therap web-based system.</p> <p><i>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</i></p> <p>CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS</p> <p>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:</p> <ul style="list-style-type: none"> (1) Complete and current ISP and all 			
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<p>supplemental plans specific to the individual;</p> <p>(2) Complete and current Health Assessment Tool;</p> <p>(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;</p> <p>(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);</p> <p>(5) Data collected to document ISP Action Plan implementation</p> <p>(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;</p> <p>(7) Physician's or qualified health care providers written orders;</p> <p>(8) Progress notes documenting implementation of a physician's or qualified health care provider's order(s);</p> <p>(9) Medication Administration Record (MAR) for the past three (3) months which includes:</p> <ul style="list-style-type: none"> (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 			
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<p>(g) An explanation of any medication irregularity, allergic reaction or adverse effect.</p> <p>(h) For PRN medication an explanation for the use of the PRN must include:</p> <ul style="list-style-type: none"> (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. <p>(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.</p> <p>(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</p> <p>(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.</p>			
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Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Date Due
Service Domain: Qualified Providers – The State monitors non-licensed/non-certified providers to assure adherence to waiver requirements. The State implements its policies and procedures for verifying that provider training is conducted in accordance with State requirements and the approved waiver.			
Tag # 1A43.1 General Events Reporting – Individual Approval	Standard Level Deficiency		
<p>Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy: General Events Reporting Effective 1/1/2012</p> <p>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.</p> <p>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</p>	<p>Based on record review the Agency did not follow the General Events Reporting requirements as indicated by the policy for 5 of 20 individuals.</p> <p>The following General Events Reporting records contained evidence that indicated the General Events Report was not entered and approved within 2 business days:</p> <p>Individual #3</p> <ul style="list-style-type: none"> General Events Report (GER) indicates on 6/11/2017 the Individual’s non-AWMD staff did not notify the nurse at least 1 hour prior to when medication was due to be administered. (Medication Error). GER was approved 8/16/2017. <p>Individual #7</p> <ul style="list-style-type: none"> General Events Report (GER) indicates on 2/15/2017 the Individual’s was taken to the emergency room. (Emergency Services). GER was approved 2/22/2017. <p>Individual #14</p> <ul style="list-style-type: none"> General Events Report (GER) indicates on 6/25/2017 the Individual’s non-AWMD staff did not notify the nurse at least 1 hour prior to when medication was due to be administered. (Medication Error). GER was approved 8/14/2017. <p>Individual #16</p> <ul style="list-style-type: none"> General Events Report (GER) indicates on 	<p>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?): →</p> <p>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): →</p>	

<p>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.</p> <p>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.</p>	<p>8/18/2017 the Individual’s was taken to the emergency room. (Emergency Services). GER was approved 8/24/2017.</p> <ul style="list-style-type: none"> • General Events Report (GER) indicates on 7/7/2017 the Individual’s was taken to the emergency room. (Emergency Services). GER was approved 7/11/2017. <p>Individual #17</p> <ul style="list-style-type: none"> • General Events Report (GER) indicates on 2/11/2017 the Individual’s was taken to the emergency room. (Emergency Services). GER was approved 3/7/2017. 		
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Standard of Care	Deficiencies	Agency Plan of Correction, On-going QA/QI and Responsible Party	Date Due
Service Domain: Health and Welfare – The state, on an ongoing basis, identifies, addresses and seeks to prevent occurrences of abuse, neglect and exploitation. Individuals shall be afforded their basic human rights. The provider supports individuals to access needed healthcare services in a timely manner.			
Tag #1A08.2 Healthcare Requirements	Standard Level Deficiency		
<p>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.</p> <p>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.</p> <p>DEVELOPMENTAL DISABILITIES SUPPORTS DIVISION (DDSD): Director’s Release: Consumer Record Requirements eff. 11/1/2012</p> <p>III. Requirement Amendments(s) or Clarifications:</p> <p>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.</p> <p>H. Readily accessible electronic records are accessible, including those stored through the Therap web-based system.</p> <p>Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015</p> <p>Chapter 5 (CIES) 3. Agency Requirements</p>	<p>Based on record review, the Agency did not provide documentation of annual physical examinations and/or other examinations as specified by a licensed physician for 2 of 20 individuals receiving Community Inclusion, Living Services and Other Services.</p> <p>Review of the administrative individual case files revealed the following items were not found, incomplete, and/or not current:</p> <p>Community Inclusion Services / Other Services Healthcare Requirements (Individuals Receiving Inclusion / Other Services Only):</p> <ul style="list-style-type: none"> • Vision Exam <ul style="list-style-type: none"> ◦ Individual #16 - As indicated by the DDSD file matrix Vision Exams are to be conducted every other year. No evidence of exam was found. • Blood Levels <ul style="list-style-type: none"> ◦ Individual #20 - As indicated by collateral documentation reviewed, lab work (lipids) was ordered on 7/19/2017. No evidence of lab results was found. 	<p>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?): →</p> <p>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (What is going to be done? How many individuals is this going to effect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?): →</p>	

<p>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 DDS Consumer Records Policy.</p> <p>Chapter 6 (CCS) 3. Agency Requirements: G. Consumer Records Policy: All Provider Agencies shall maintain at the administrative office a confidential case file for each individual. Provider agency case files for individuals are required to comply with the DDS Individual Case File Matrix policy.</p> <p>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 DDS Individual Case File Matrix policy.</p> <p>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 DDS Individual Case File Matrix policy.</p> <p>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 DDS Individual Case File Matrix policy.</p> <p>Chapter 13 (IMLS) 2. Service Requirements:</p>			
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<p>C. Documents to be maintained in the agency administrative office, include: (This is not an all-inclusive list refer to standard as it includes other items)...</p> <p>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</p> <p>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:</p> <p>(5) A medical history, which shall include at least demographic data, current and past medical diagnoses including the cause (if known) of the developmental disability, psychiatric diagnoses, allergies (food, environmental, medications), immunizations, and most recent physical exam;</p> <p>CHAPTER 6. VI. GENERAL REQUIREMENTS FOR COMMUNITY LIVING</p> <p>G. Health Care Requirements for Community Living Services.</p> <p>(1) The Community Living Service providers shall ensure completion of a HAT for each individual receiving this service. The HAT shall be completed 2 weeks prior to the annual ISP meeting and submitted to the Case Manager and all other IDT Members. A revised HAT is required to also be submitted whenever the individual's health status changes significantly.</p>			
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<p>For individuals who are newly allocated to the DD Waiver program, the HAT may be completed within 2 weeks following the initial ISP meeting and submitted with any strategies and support plans indicated in the ISP, or within 72 hours following admission into direct services, whichever comes first.</p> <p>(2) Each individual will have a Health Care Coordinator, designated by the IDT. When the individual's HAT score is 4, 5 or 6 the Health Care Coordinator shall be an IDT member, other than the individual. The Health Care Coordinator shall oversee and monitor health care services for the individual in accordance with these standards. In circumstances where no IDT member voluntarily accepts designation as the health care coordinator, the community living provider shall assign a staff member to this role.</p> <p>(3) For each individual receiving Community Living Services, the provider agency shall ensure and document the following:</p> <ul style="list-style-type: none"> (a) Provision of health care oversight consistent with these Standards as detailed in Chapter One section III E: Healthcare Documentation by Nurses For Community Living Services, Community Inclusion Services and Private Duty Nursing Services. b) That each individual with a score of 4, 5, or 6 on the HAT, has a Health Care Plan developed by a licensed nurse. (c) That an individual with chronic condition(s) with the potential to exacerbate into a life threatening condition, has Crisis Prevention/ Intervention Plan(s) developed by a licensed nurse or other appropriate professional for each such condition. <p>(4) That an average of 3 hours of documented nutritional counseling is available annually, if</p>			
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<p>recommended by the IDT.</p> <p>(5) That the physical property and grounds are free of hazards to the individual's health and safety.</p> <p>(6) In addition, for each individual receiving Supported Living or Family Living Services, the provider shall verify and document the following:</p> <ul style="list-style-type: none"> (a) The individual has a primary licensed physician; (b) The individual receives an annual physical examination and other examinations as specified by a licensed physician; (c) The individual receives annual dental check-ups and other check-ups as specified by a licensed dentist; (d) The individual receives eye examinations as specified by a licensed optometrist or ophthalmologist; and (e) Agency activities that occur as follow-up to medical appointments (e.g. treatment, visits to specialists, changes in medication or daily routine). 			
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Tag # 1A15.1 Nurse Availability	Standard Level Deficiency		
<p>Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015</p> <p>CHAPTER 6 (CCS) 3. Agency Requirements</p> <p>C. Employ or subcontract with at least one RN to comply with services under “Nursing and Medical Oversight Services as needed” that is detailed in the Scope of Services above for Group Customized Community Supports Services. If the size of the provider warrants more than one nurse, a RN must supervise LPNs.</p> <p>1. Ensure compliance with the New Mexico Nurse Practice Act and DDSD Policies and Procedures regarding Delegation of Specific Nursing Functions, including:</p> <p>i. Provider agencies (Small group and Group services) must develop and implement policies and procedures regarding delegation which must comply with relevant DDSD Policies and Procedures, and the New Mexico Nurse Practice Act. Agencies must ensure that all nurses they employ or contract with are knowledgeable of all these requirements;</p> <p>CHAPTER 11. 2. Service Requirements I. Health Care Requirements for Family Living:</p> <p>9. Family Living Provider Agencies are required to be an Adult Nursing provider and have a Registered Nurse (RN) licensed by the State of New Mexico on staff and residing in New Mexico or bordering towns see: Adult Nursing requirements. The agency nurse may be an employee or a sub-contractor.</p> <p>A. The Family Living Provider Agency must not</p>	<p>Based on interview, the Agency did not ensure nursing services were available as needed for 1 of 20 individuals.</p> <p>When Direct Service Professionals (DSP) were asked if there is a nurse available to the individual at all times, the following was reported:</p> <ul style="list-style-type: none"> DSP #605 stated, “No. It’s spotty on weekends.” 	<p>Provider: State your Plan of Correction for the deficiencies cited in this tag here (<i>How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?</i>): →</p> <p>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (<i>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>): →</p>	

<p>use a LPN without a RN supervisor. The RN must provide face to face supervision required by the New Mexico Nurse Practice Act and these services standards for LPNs, CMAs, and direct support personnel who have been delegated nursing tasks.</p> <p>B. On-call nursing services: An on-call nurse must be available to surrogate or host families DSP for medication oversight. It is expected that no single nurse carry the full burden of on-call duties for the agency.</p> <p>CHAPTER 12. 2. Service Requirements. L. Training and Requirement: 6. Nursing Requirements and Roles:</p> <p>A. Supported Living Provider Agencies are required to have a RN licensed by the State of New Mexico on staff. The agency nurse may be an employee or a sub-contractor.</p> <p>CHAPTER 13. 1. SCOPE OF SERVICE. A. Living Supports- Intensive Medical Living Service includes the following:</p> <p>1. Provide appropriate levels of supports: Agency nurses and Direct Support Personnel (DSP) provide individualized support based upon assessed need. Assessment shall include use of required health-related assessments, eligibility parameters issued by the Developmental Disabilities Support Division (DDSD), other pertinent assessments completed by the nurse, and the nurse’s professional judgment.</p> <p>2. Provide daily nursing visits:</p> <p>a. A daily, face to face nursing visit must be made by a Registered Nurse (RN) or Licensed Practical Nurse (LPN) in order to deliver required direct nursing care, monitor each individual’s status, and oversee DSP delivery of health related care and</p>			
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<p>interventions. Face to face nursing visits may not be delegated to non-licensed staff.</p> <p>b. Although a nurse may be present in the home for extended periods of time, a nurse is not required to be present in the home during periods of time when direct nursing services are not needed.</p> <p>NEW MEXICO NURSING PRACTICE ACT CHAPTER 61, ARTICLE 3</p> <p>I. "licensed practical nursing" means the practice of a directed scope of nursing requiring basic knowledge of the biological, physical, social and behavioral sciences and nursing procedures, which practice is at the direction of a registered nurse, physician or dentist licensed to practice in this state. This practice includes but is not limited to:</p> <p>(1) contributing to the assessment of the health status of individuals, families and communities;</p> <p>(2) participating in the development and modification of the plan of care;</p> <p>(3) implementing appropriate aspects of the plan of care commensurate with education and verified competence;</p> <p>(4) collaborating with other health care professionals in the management of health care; and</p> <p>(5) participating in the evaluation of responses to interventions;</p>			
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Tag # 1A31 Client Rights/Human Rights	Standard Level Deficiency		
<p>7.26.3.11 RESTRICTIONS OR LIMITATION OF CLIENT'S RIGHTS:</p> <p>A. A service provider shall not restrict or limit a client's rights except:</p> <p>(1) where the restriction or limitation is allowed in an emergency and is necessary to prevent imminent risk of physical harm to the client or another person; or</p> <p>(2) where the interdisciplinary team has determined that the client's limited capacity to exercise the right threatens his or her physical safety; or</p> <p>(3) as provided for in Section 10.1.14 [now Subsection N of 7.26.3.10 NMAC].</p> <p>B. Any emergency intervention to prevent physical harm shall be reasonable to prevent harm, shall be the least restrictive intervention necessary to meet the emergency, shall be allowed no longer than necessary and shall be subject to interdisciplinary team (IDT) review. The IDT upon completion of its review may refer its findings to the office of quality assurance. The emergency intervention may be subject to review by the service provider's behavioral support committee or human rights committee in accordance with the behavioral support policies or other department regulation or policy.</p> <p>C. The service provider may adopt reasonable program policies of general applicability to clients served by that service provider that do not violate client rights. [09/12/94; 01/15/97; Recompiled 10/31/01]</p> <p>Long Term Services Division Policy Title: Human Rights Committee Requirements Eff Date: March 1, 2003</p>	<p>Based on record review, the Agency did not ensure the rights of Individuals were not restricted or limited for 1 of 20 Individuals.</p> <p>A review of Agency Individual files found no documentation of Positive Behavior Plans and/or Positive Behavior Crisis Plans, which contain restrictions being reviewed at least quarterly by the Human Rights Committee.</p> <p>No current Human Rights Approval was found for the following:</p> <ul style="list-style-type: none"> Room Searches. Last Review was dated 7/17/2017. (Individual #8) 	<p>Provider: State your Plan of Correction for the deficiencies cited in this tag here (<i>How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?</i>): →</p> <p>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (<i>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>): →</p>	<p> </p>

<p>IV. POLICY STATEMENT - Human Rights Committees are required for residential service provider agencies. The purpose of these committees with respect to the provision of Behavior Supports is to review and monitor the implementation of certain Behavior Support Plans.</p> <p>Human Rights Committees may not approve any of the interventions specifically prohibited in the following policies:</p> <ul style="list-style-type: none"> • Aversive Intervention Prohibitions • Psychotropic Medications Use • Behavioral Support Service Provision. <p>A Human Rights Committee may also serve other agency functions as appropriate, such as the review of internal policies on sexuality and incident management follow-up.</p> <p>A. HUMAN RIGHTS COMMITTEE ROLE IN BEHAVIOR SUPPORTS</p> <p>Only those Behavior Support Plans with an aversive intervention included as part of the plan or associated Crisis Intervention Plan need to be reviewed prior to implementation. Plans not containing aversive interventions do not require Human Rights Committee review or approval.</p> <p>2. The Human Rights Committee will determine and adopt a written policy stating the frequency and purpose of meetings. Behavior Support Plans approved by the Human Rights Committee will be reviewed at least quarterly.</p> <p>3. Records, including minutes of all meetings will be retained at the agency with primary responsibility for implementation for at least five years from the completion of each individual's Individual Service Plan.</p>			
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**Department of Health Developmental
Disabilities Supports Division (DDSD) -
Procedure Title:**

**Medication Assessment and Delivery
Procedure Eff Date: November 1, 2006**

B. 1. e. If the PRN medication is to be used in response to psychiatric and/or behavioral symptoms in addition to the above requirements, obtain current written consent from the individual, guardian or surrogate health decision maker and submit for review by the agency's Human Rights Committee (References: Psychotropic Medication Use Policy, Section D, page 5 Use of PRN Psychotropic Medications; and, Human Rights Committee Requirements Policy, Section B, page 4 Interventions Requiring Review and Approval – Use of PRN Medications).

Tag # LS25 / 6L25 Residential Health and Safety (SL/FL)	Standard Level Deficiency		
<p>Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015</p> <p>CHAPTER 11 (FL) Living Supports – Family Living Agency Requirements G. Residence Requirements for Living Supports- Family Living Services: 1. Family Living Services providers must assure that each individual’s residence is maintained to be clean, safe and comfortable and accommodates the individuals’ daily living, social and leisure activities. In addition, the residence must:</p> <p>a. Maintain basic utilities, i.e., gas, power, water and telephone;</p> <p>b. Provide environmental accommodations and assistive technology devices in the residence including modifications to the bathroom (i.e., shower chairs, grab bars, walk in shower, raised toilets, etc.) based on the unique needs of the individual in consultation with the IDT;</p> <p>c. Have a battery operated or electric smoke detectors, carbon monoxide detectors, fire extinguisher, or a sprinkler system;</p> <p>d. Have a general-purpose first aid kit;</p> <p>e. Allow at a maximum of two (2) individuals to share, with mutual consent, a bedroom and each individual has the right to have his or her own bed;</p> <p>f. Have accessible written documentation of actual evacuation drills occurring at least three (3) times a year;</p>	<p>Based on observation, the Agency did not ensure that each individuals’ residence met all requirements within the standard for 1 of 1 Family Living residences.</p> <p>Review of the residential records and observation of the residence revealed the following items were not found, not functioning or incomplete:</p> <p>Family Living Requirements:</p> <ul style="list-style-type: none"> • General-purpose first aid kit (#12) 	<p>Provider: State your Plan of Correction for the deficiencies cited in this tag here (<i>How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?</i>): →</p> <p>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (<i>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>): →</p>	<p> </p>

<p>g. Have accessible written procedures for the safe storage of all medications with dispensing instructions for each individual that are consistent with the Assisting with Medication Delivery training or each individual's ISP; and</p> <p>h. Have accessible written procedures for emergency placement and relocation of individuals in the event of an emergency evacuation that makes the residence unsuitable for occupancy. The emergency evacuation procedures must address, but are not limited to, fire, chemical and/or hazardous waste spills, and flooding.</p> <p>CHAPTER 12 (SL) Living Supports – Supported Living Agency Requirements G. Residence Requirements for Living Supports- Supported Living Services: 1. Supported Living Provider Agencies must assure that each individual's residence is maintained to be clean, safe, and comfortable and accommodates the individual's daily living, social, and leisure activities. In addition, the residence must:</p> <p>a. Maintain basic utilities, i.e., gas, power, water, and telephone;</p> <p>b. Provide environmental accommodations and assistive technology devices in the residence including modifications to the bathroom (i.e., shower chairs, grab bars, walk in shower, raised toilets, etc.) based on the unique needs of the individual in consultation with the IDT;</p> <p>c. Ensure water temperature in home does not exceed safe temperature (110° F);</p>			
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<p>d. Have a battery operated or electric smoke detectors and carbon monoxide detectors, fire extinguisher, or a sprinkler system;</p> <p>e. Have a general-purpose First Aid kit;</p> <p>f. Allow at a maximum of two (2) individuals to share, with mutual consent, a bedroom and each individual has the right to have his or her own bed;</p> <p>g. Have accessible written documentation of actual evacuation drills occurring at least three (3) times a year. For Supported Living evacuation drills must occur at least once a year during each shift;</p> <p>h. Have accessible written procedures for the safe storage of all medications with dispensing instructions for each individual that are consistent with the Assisting with Medication Delivery training or each individual's ISP; and</p> <p>i. Have accessible written procedures for emergency placement and relocation of individuals in the event of an emergency evacuation that makes the residence unsuitable for occupancy. The emergency evacuation procedures must address, but are not limited to, fire, chemical and/or hazardous waste spills, and flooding.</p> <p>CHAPTER 13 (IMLS) 2. Service Requirements R. Staff Qualifications: 3. Supervisor Qualifications And Requirements: S Each residence shall include operable safety equipment, including but not limited to, an operable smoke detector or sprinkler system, a carbon monoxide detector if any natural gas appliance or heating is used, fire</p>			
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<p>extinguisher, general purpose first aid kit, written procedures for emergency evacuation due to fire or other emergency and documentation of evacuation drills occurring at least annually during each shift, phone number for poison control within line of site of the telephone, basic utilities, general household appliances, kitchen and dining utensils, adequate food and drink for three meals per day, proper food storage, and cleaning supplies.</p> <p>T Each residence shall have a blood borne pathogens kit as applicable to the residents' health status, personal protection equipment, and any ordered or required medical supplies shall also be available in the home.</p> <p>U If not medically contraindicated, and with mutual consent, up to two (2) individuals may share a single bedroom. Each individual shall have their own bed. All bedrooms shall have doors that may be closed for privacy. Individuals have the right to decorate their bedroom in a style of their choosing consistent with safe and sanitary living conditions.</p> <p>V For residences with more than two (2) residents, there shall be at least two (2) bathrooms. Toilets, tubs/showers used by the individuals shall provide for privacy and be designed or adapted for the safe provision of personal care. Water temperature shall be maintained at a safe level to prevent injury and ensure comfort and shall not exceed one hundred ten (110) degrees.</p>			
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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 # LS26 / 6L26 Supported Living Reimbursement	Standard Level Deficiency		
<p>Developmental Disabilities (DD) Waiver Service Standards effective 11/1/2012 revised 4/23/2013; 6/15/2015</p> <p>CHAPTER 12 (SL) 4. REIMBURSEMENT</p> <p>A. Supported Living Provider Agencies must maintain all records necessary to fully disclose the type, quality, quantity, and clinical necessity of services furnished to individuals who are currently receiving services. The Supported Living Provider Agency records must be sufficiently detailed to substantiate the date, time, individual name, servicing provider, nature of services, and length of a session of service billed. Providers are required to comply with the Human Services Department Billing Regulations.</p> <p>a. The rate for Supported Living is based on categories associated with each individual's NM DDW Group; and</p> <p>b. A non-ambulatory stipend is available for those who meet assessed need requirements.</p> <p>B. Billable Units:</p> <p>1. The billable unit for Supported Living is based on a daily rate. A day is considered 24 hours from midnight to midnight. If 12 or less hours of service are provided then one half unit shall be billed. A whole unit can be billed if more than 12 hours of service is provided during a 24 hour period.</p> <p>2. The maximum allowable billable units</p>	<p>Based on record review, the Agency did not provide written or electronic documentation as evidence for each unit billed for Supported Living Services for 1 of 12 individuals.</p> <p>Individual #2 October 2017</p> <ul style="list-style-type: none"> The Agency billed 1 unit of Supported Living (T2016 HB U5) on 10/17/2017. Documentation received accounted for .5 units. 	<p>Provider: State your Plan of Correction for the deficiencies cited in this tag here (<i>How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?</i>): →</p> <p>Provider: Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here (<i>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>): →</p>	

<p>cannot exceed three hundred forty (340) calendar days per ISP year or one hundred seventy (170) calendar days per six (6) months.</p> <p>C. Billable Activities:</p> <p>1. Billable activities shall include any activities which DSP provides in accordance with the Scope of Services for Living Supports which are not listed in non-billable services, activities, or situations below.</p> <p>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</p>			
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<p>to any of the following for a period of at least six years from the payment date:</p> <ul style="list-style-type: none"> (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. <p>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</p> <p>CHAPTER 1 III. PROVIDER AGENCY DOCUMENTATION OF SERVICE DELIVERY AND LOCATION</p> <p>A. General: All Provider Agencies shall maintain all records necessary to fully disclose the service, quality, quantity and clinical necessity furnished to individuals who are currently receiving services. The Provider Agency records shall be sufficiently detailed to substantiate the date, time, individual name, servicing Provider Agency, level of services, and length of a session of service billed.</p> <p>B. Billable Units: The documentation of the billable time spent with an individual shall be kept on the written or electronic record that is prepared prior to a request for reimbursement from the HSD. For each unit billed, the record shall contain the following:</p> <ul style="list-style-type: none"> (1) Date, start and end time of each service encounter or other billable service interval; (2) A description of what occurred during the encounter or service interval; and (3) The signature or authenticated name of staff providing the service. <p>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</p>			
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<p>CHAPTER 6. IX. REIMBURSEMENT FOR COMMUNITY LIVING SERVICES</p> <p>A. Reimbursement for Supported Living Services</p> <p>(1) Billable Unit. The billable Unit for Supported Living Services is based on a daily rate. The daily rate cannot exceed 340 billable days a year.</p> <p>(2) Billable Activities</p> <p>(a) Direct care provided to an individual in the residence any portion of the day.</p> <p>(b) Direct support provided to an individual by community living direct service staff away from the residence, e.g., in the community.</p> <p>(c) Any activities in which direct support staff provides in accordance with the Scope of Services.</p> <p>(3) Non-Billable Activities</p> <p>(a) The Supported Living Services provider shall not bill DD Waiver for Room and Board.</p> <p>(b) Personal care, respite, nutritional counseling and nursing supports shall not be billed as separate services for an individual receiving Supported Living Services.</p> <p>(c) The provider shall not bill when an individual is hospitalized or in an institutional care setting.</p>			
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Date: June 18, 2018

To: Damian Houfek, President, Chief Executive Officer
Provider: ENMRSH, Inc.
Address: 2700 East 7th Street
State/Zip: Clovis, New Mexico 88101

E-mail Address: dhoufek@enmrsh.org

CC: Cathy Mills, Board Chair
Board Chair
Address: 2609 Putnam
Clovis, New Mexico 88101

Region: Southeast
Survey Date: January 26 – February 1, 2018
Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: **2012:** Supported Living, Family Living, Intensive Medical Living, Customized Community Supports, Community Integrated Employment Services and Customized In-Home Supports

Survey Type: Routine

Dear Mr. Houfek;

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.D1808.4.RTN.11.18.169