Dear Ms. Romero:

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

**Determination of Compliance:**

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

**Compliance with all Conditions of Participation.**

This determination is based on your agency's compliance with CMS waiver assurances at the Condition of Participation level. The attached QMB Report of Findings indicates Standard Level deficiencies identified and requires implementation of a Plan of Correction.
Plan of Correction:
The attached Report of Findings identifies the Standard Level and/or Condition of Participation deficiencies found during your agency’s compliance review. You are required to complete and implement a Plan of Correction. Your agency has a total of 45 business days (10 business days to submit your POC for approval and 35 days to implement your approved Plan of Correction) from the receipt of this letter.

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: Plan of Correction Coordinator
   5301 Central Ave. NE Suite 400 Albuquerque, NM 87108

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.

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 at 505-699-0714 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,

Marti Madrid, LBSW
Marti Madrid, LBSW
Team Lead/Healthcare Surveyor
Division of Health Improvement
Quality Management Bureau
Survey Process Employed:

Entrance Conference Date: May 15, 2012

Present:

**Easter Seals Santa Maria el Mirador**
Barbara Hall, Program Manager
Gary Maestas, Residential Manager

**DOH/DHI/QMB**
Marti Madrid, LBSW, Team Lead/Healthcare Surveyor
Erica Nilsen, BA, Healthcare Surveyor

Exit Conference Date: May 16, 2012

Present:

**Easter Seals Santa Maria el Mirador**
Barbara Hall, Program Manager

**DOH/DHI/QMB**
Marti Madrid, LBSW Team Lead/Healthcare Surveyor
Erica Nilsen, BA, Healthcare Surveyor

Total Homes Visited

Number: 1

- Supported Homes Visited
  Number: 1

Administrative Locations Visited

Number: 1

Total Sample Size

Number: 3

- Jackson Class Members
- Non-Jackson Class Members
- Supported Living
- Independent Living
- Adult Habilitation
- Supported Employment

Persons Served Records Reviewed

Number: 3

Persons Served Interviewed

Number: 2

Persons Served Observed

Number: 1 (Individual was not available during the on-site survey).

Direct Support Personnel Interviewed

Number: 4

Direct Support Personnel Records Reviewed

Number: 23

Service Coordinator Records Reviewed

Number: 2

Administrative Files Reviewed

- Billing Records
- Medical Records
- Incident Management Records
- Personnel Files
- Training Records
• Agency Policy and Procedure
• Caregiver Criminal History Screening Records
• Employee Abuse Registry
• Human Rights Notes and/or Meeting Minutes
• Evacuation Drills
• 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
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 non compliance.

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

If you have questions about the Plan of Correction process, call the QMB Plan of Correction Coordinator at 505-699-0714 or email at scott.good@state.nm.us. Requests for technical assistance must be requested through your DDSD Regional 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 required six CMS core elements to address each deficiency of the POC:

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.

6. The POC must be signed and dated by the agency director or other authorized official.

The following details should be considered when developing your POC:

- 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, Incident Reporting, and Individual-Specific service requirements, etc;
- How accuracy in Billing documentation is assured;
- How health, safety is assured;
- For Case Management Providers, how ISPs 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; 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 should 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 QMB Deputy Chief, Scott Good at 505-699-0714 for assistance.
3. For Technical Assistance (TA) in developing or implementing your POC, contact your local DDSD Regional Office.
4. Submit your POC to Scott Good, QMB Deputy Chief in any of the following ways:
   a. Electronically at scott.good@state.nm.us (preferred method)
   b. Fax to 505-222-8661, or
   c. Mail to POC Coordinator, 5301 Central Avenue SW, Suite 400, Albuquerque, NM 87108
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 “approve” 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.

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. You may submit your documents by postal mail (paper hard copy or on a disc), fax, or 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. For billing deficiencies, you must submit:
   a. Evidence of an internal audit of billing documentation for a sample of individuals and timeframes;
   b. Copies of “void and adjust” forms submitted to correct all over-billed or unjustified units billed identified during your internal audit.

Revisions, Modifications or Extensions to your Plan of Correction (post QMB approval) must be made in writing and submitted to the Deputy Chief at QMB, 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.
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) but may have standard level deficiencies (deficiencies which are not at the condition level) out of compliance. 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.

- **“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 one (1) to three (3) ‘Conditions of Participation.’ 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).

  Providers receiving a repeat determination of ‘Partial-Compliance’ for repeat deficiencies of CoPs may be referred by the Quality Management Bureau to the Internal Review Committee (IRC) for consideration of remedies and possible actions.

- **“Non-Compliant 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 and/or has:
  - Four (4) Conditions of Participation out of compliance.
  - Multiple findings of widespread non-compliance with any standard or regulation with a significant potential for more than minimal harm.
  - Any finding of actual harm or Immediate Jeopardy.

  The Agency may also have standard level deficiencies (deficiencies which are not at the condition level).

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

The following limitations apply to the IRF process:
- The 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 made 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: Easter Seals Santa Maria el Mirador - Northeast Region
Program: Developmental Disabilities Waiver
Service: Community Living Supports (Supported Living & Independent Living) & Community Inclusion Supports (Adult Habilitation & Supported Employment)
Monitoring Type: Routine Survey
Date of Survey: May 15 – 16, 2012

<table>
<thead>
<tr>
<th>Standard of Care</th>
<th>Deficiencies</th>
<th>Agency Plan of Correction, On-going QA/QI &amp; Responsible Party</th>
<th>Date Due</th>
</tr>
</thead>
<tbody>
<tr>
<td>CMS Assurance – 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.</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tag # 1A08 Agency Case File</td>
<td>Standard Level Deficiency</td>
<td></td>
<td></td>
</tr>
<tr>
<td>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.</td>
<td>Based on record review, the Agency failed to maintain at the administrative office a confidential case file for 1 of 3 individuals. Review of the Agency individual case files found the following items were not found, incomplete, and/or not current: • Documentation of Guardianship/Power of Attorney (#2)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Provider:
State your Plan of Correction for the findings in this Tag above this line.

Enter your Quality Assurance/Quality Improvement processes below the line.
requirements:
(1) Emergency contact information, including the individual’s address, telephone number, names and telephone numbers of relatives, or guardian or conservator, physician’s name(s) and telephone number(s), pharmacy name, address and telephone number, and health plan if appropriate;
(2) The individual’s complete and current ISP, with all supplemental plans specific to the individual, and the most current completed Health Assessment Tool (HAT);
(3) Progress notes and other service delivery documentation;
(4) Crisis Prevention/Intervention Plans, if there are any for the individual;
(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;
(6) When applicable, transition plans completed for individuals at the time of discharge from Fort Stanton Hospital or Los Lunas Hospital and Training School; and
(7) Case records belong to the individual receiving services and copies shall be provided to the individual upon request.
(8) The receiving Provider Agency shall be provided at a minimum the following records whenever an individual changes provider agencies:
(a) Complete file for the past 12 months;
(b) ISP and quarterly reports from the current and prior ISP year;
(c) Intake information from original admission to services; and
(d) When applicable, the Individual
Transition Plan at the time of discharge from Los Lunas Hospital and Training School or Ft. Stanton Hospital.

NMAC 8.302.1.17 RECORD KEEPING AND DOCUMENTATION REQUIREMENTS: A provider must maintain all the records necessary to fully disclose the nature, quality, amount and medical necessity of services furnished to an eligible recipient who is currently receiving or who has received services in the past.

B. Documentation of test results: Results of tests and services must be documented, which includes results of laboratory and radiology procedures or progress following therapy or treatment.
Tag # 1A32 & 6L14 ISP Implementation

<table>
<thead>
<tr>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>Based on record review, the Agency failed to 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 1 of 3 individuals. Per Individuals ISP the following was found with regards to the implementation of ISP Outcomes:</td>
</tr>
</tbody>
</table>

**Administrative Files Reviewed:**

**Independent Living Data Collection/Data Tracking/Progress with regards to ISP Outcomes:**

Individual #3
- None found regarding: “…will make successful purchases in the community,” for 4/2012.

<table>
<thead>
<tr>
<th>Provider:</th>
</tr>
</thead>
<tbody>
<tr>
<td>State your Plan of Correction for the findings in this Tag above this line.</td>
</tr>
</tbody>
</table>

Enter your Quality Assurance/Quality Improvement processes below the line.
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]
<table>
<thead>
<tr>
<th>Tag # 6L17 Reporting Requirements (Community Living Quarterly Reports)</th>
<th>Standard Level Deficiency</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</td>
<td>Based on record review, the Agency failed to complete written quarterly status reports for 1 of 2 individuals receiving Community Living Services.</td>
<td></td>
</tr>
</tbody>
</table>
| CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS | **Independent Living Annual Assessment:**  
  - Individual #3 - None found for 10/6/2010 – 10/6/2011. |   |
| D. Community Living Service Provider Agency Reporting Requirements: All Community Living Support providers shall submit written quarterly status reports to the individual’s Case Manager and other IDT Members no later than fourteen (14) days following the end of each ISP quarter. The quarterly reports shall contain the following written documentation:  
  (1) Timely completion of relevant activities from ISP Action Plans  
  (2) Progress towards desired outcomes in the ISP accomplished during the quarter;  
  (3) Significant changes in routine or staffing;  
  (4) Unusual or significant life events;  
  (5) Updates on health status, including medication and durable medical equipment needs identified during the quarter; and  
  (6) Data reports as determined by IDT members. |   |   |

Provider:  
State your Plan of Correction for the findings in this Tag above this line.  

Enter your Quality Assurance/Quality Improvement processes below the line.
<table>
<thead>
<tr>
<th>TAG6L14 Residential Case File</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</td>
<td>Based on record review, the Agency failed to maintain a complete and confidential case file in the residence for 1 of 3 Individuals receiving Supported Living Services.</td>
</tr>
<tr>
<td>CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS</td>
<td>The following was not found, incomplete and/or not current:</td>
</tr>
<tr>
<td>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.</td>
<td></td>
</tr>
<tr>
<td>Provider:</td>
<td></td>
</tr>
<tr>
<td>State your Plan of Correction for the findings in this Tag above this line.</td>
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<tr>
<td>Enter your Quality Assurance/Quality Improvement processes below the line.</td>
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Q.12.4.DDW.D0974.2.001.RTN.1. 171

7
implementation

(6) Progress notes written by direct care staff and by nurses regarding individual health status and physical conditions including action taken in response to identified changes in condition for at least the past month;
(7) Physician’s or qualified health care providers written orders;
(8) Progress notes documenting implementation of a physician’s or qualified health care provider’s order(s);
(9) Medication Administration Record (MAR) for the past three (3) months which includes:
   (a) The name of the individual;
   (b) A transcription of the healthcare practitioners prescription including the brand and generic name of the medication;
   (c) Diagnosis for which the medication is prescribed;
   (d) Dosage, frequency and method/route of delivery;
   (e) Times and dates of delivery;
   (f) Initials of person administering or assisting with medication; and
   (g) An explanation of any medication irregularity, allergic reaction or adverse effect.
   (h) For PRN medication an explanation for the use of the PRN must include:
      (i) Observable signs/symptoms or circumstances in which the medication is to be used, and
      (ii) Documentation of the effectiveness/result of the PRN delivered.
   (i) A MAR is not required for individuals participating in Independent Living Services who self-administer their own medication. However, when medication administration
is provided as part of the Independent Living Service a MAR must be maintained at the individual's home and an updated copy must be placed in the agency file on a weekly basis.

(10) Record of visits to healthcare practitioners including any treatment provided at the visit and a record of all diagnostic testing for the current ISP year; and

(11) Medical History to include: demographic data, current and past medical diagnoses including the cause (if known) of the developmental disability and any psychiatric diagnosis, allergies (food, environmental, medications), status of routine adult health care screenings, immunizations, hospital discharge summaries for past twelve (12) months, past medical history including hospitalizations, surgeries, injuries, family history and current physical exam.
### Standard of Care

**CMS Assurance – Qualified Providers** – The State monitors non-licensed/non-certified providers to assure adherence to waiver requirements. The State implements its policies and procedures for verifying that provider training is conducted in accordance with State requirements and the approved waiver.

<table>
<thead>
<tr>
<th>Tag # 1A22 Agency Personnel Competency</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 <strong>CHAPTER 1 IV. GENERAL REQUIREMENTS FOR PROVIDER AGENCY SERVICE PERSONNEL:</strong> The objective of this section is to establish personnel standards for DD Medicaid Waiver Provider Agencies for the following services: Community Living Supports, Community Inclusion Services, Respite, Substitute Care and Personal Support Companion Services. These standards apply to all personnel who provide services, whether directly employed or subcontracting with the Provider Agency. Additional personnel requirements and qualifications may be applicable for specific service standards. <strong>F. Qualifications for Direct Service Personnel:</strong> The following employment qualifications and competency requirements are applicable to all Direct Service Personnel employed by a Provider Agency: (1) Direct service personnel shall be eighteen (18) years or older. Exception: Adult Habilitation can employ direct care personnel under the age of eighteen 18 years, but the employee shall work directly under a supervisor, who is physically present at all times; (2) Direct service personnel shall have the ability</td>
<td>Based on interview, the Agency failed to ensure that training competencies were met for 2 of 4 Direct Support Personnel. <strong>When DSP were asked if the Individual had a Medical Emergency Response Plans and if so, what the plan(s) covered, the following was reported:</strong> • DSP #43 stated, “Hypertension.” According to the Individual Specific Training section of the ISP the Individual requires Medical Emergency Response Plans for allergies and gastrointestinal. (Individual #1) <strong>When DSP were asked what the individual’s Diagnosis were, the following was reported:</strong> • DSP #45 stated, “Asthma, other than that she is in good health.” According to the individuals ISP she is diagnosed with Down Syndrome and high cholesterol. Staff did not discuss the listed diagnosis. (Individual #3)</td>
</tr>
</tbody>
</table>

Provider:
State your Plan of Correction for the findings in this Tag above this line.

Enter your Quality Assurance/Quality Improvement processes below the line.
to read and carry out the requirements in an ISP;

(3) Direct service personnel shall be available to communicate in the language that is functionally required by the individual or in the use of any specific augmentative communication system utilized by the individual;

(4) Direct service personnel shall meet the qualifications specified by DDSD in the Policy Governing the Training Requirements for Direct Support Staff and Internal Service Coordinators, Serving Individuals with Developmental Disabilities; and

(5) Direct service Provider Agencies of Respite Services, Substitute Care, Personal Support Services, Nutritional Counseling, Therapists and Nursing shall demonstrate basic knowledge of developmental disabilities and have training or demonstrable qualifications related to the role he or she is performing and complete individual specific training as required in the ISP for each individual he or she support.

(6) Report required personnel training status to the DDSD Statewide Training Database as specified in DDSD policies as related to training requirements as follows:

(a) Initial comprehensive personnel status report (name, date of hire, Social Security number category) on all required personnel to be submitted to DDSD Statewide Training Database within the first ninety (90) calendar days of providing services;

(b) Staff who do not wish to use his or her
Social Security Number may request an alternative tracking number; and (c) Quarterly personnel update reports sent to DDSD Statewide Training Database to reflect new hires, terminations, inter-provider Agency position changes, and name changes.

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.
### CMS Assurance – 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.

<table>
<thead>
<tr>
<th>Tag #</th>
<th>Medication Delivery (MAR) - Routine Medication</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A09</td>
<td>Standard of Care</td>
<td>Deficiencies</td>
</tr>
<tr>
<td></td>
<td><strong>CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS:</strong> 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.</td>
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<td></td>
<td><strong>E. Medication Delivery:</strong> 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.</td>
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<td>(2) When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) shall be maintained and include:</td>
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<tr>
<td></td>
<td>(a) The name of the individual, a transcription of the physician's written or licensed health care provider's</td>
<td></td>
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<tr>
<td></td>
<td>Medication Administration Records (MAR) were reviewed for the months of January, February and March 2012</td>
<td></td>
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<tr>
<td></td>
<td>Based on record review, 1 of 3 individuals had Medication Administration Records, which contained missing medications entries and/or other errors:</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Individual #2</td>
<td></td>
</tr>
<tr>
<td></td>
<td>February 2012</td>
<td></td>
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<tr>
<td></td>
<td>Medication Administration Record did not contain the form (i.e. liquid, tablet, capsule, etc.) of medication to be taken for the following:</td>
<td></td>
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<tr>
<td></td>
<td>• Propranolol 80 mg (1 times daily)</td>
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<tr>
<td></td>
<td>March 2012</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Medication Administration Record did not contain the form (i.e. liquid, tablet, capsule, etc.) of medication to be taken for the following:</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Propranolol 80 mg (1 times daily)</td>
<td></td>
</tr>
</tbody>
</table>

Provider:
State your Plan of Correction for the findings in this Tag above this line.

Enter your Quality Assurance/Quality Improvement processes below the line.
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;

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.
<table>
<thead>
<tr>
<th>Tag # 1A15.2 &amp; 5I09 - Healthcare Documentation</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>Based on record review, the Agency failed to maintain the required documentation in the Individuals Agency Record as required per standard for 2 of 3 individual The following were not found, incomplete and/or not current:</td>
<td></td>
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<tr>
<td>- Quarterly Nursing Review of HCP/Crisis Plans:</td>
<td></td>
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<tr>
<td>- None found for 2/2012 - 4/2012 (#2)</td>
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<tr>
<td>- Special Health Care Needs</td>
<td></td>
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<tr>
<td>- Mealtime Plan</td>
<td></td>
</tr>
<tr>
<td>- Individual #2 - As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found.</td>
<td></td>
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<tr>
<td>- Health Care Plans</td>
<td></td>
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<tr>
<td>- Diabetes</td>
<td></td>
</tr>
<tr>
<td>- Individual #2 - As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found.</td>
<td></td>
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<tr>
<td>- Skin Integrity</td>
<td></td>
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<tr>
<td>- Individual #2 - As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found.</td>
<td></td>
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<tr>
<td>- Constipation</td>
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<tr>
<td>- Individual #2 - As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found.</td>
<td></td>
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<tr>
<td>- Gastrointestinal</td>
<td></td>
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<tr>
<td>- Individual #2 - As indicated by the IST</td>
<td></td>
</tr>
</tbody>
</table>

Provider:
State your Plan of Correction for the findings in this Tag above this line.

Enter your Quality Assurance/Quality Improvement processes below the line.
Agency nurses may also complete these assessments in collaboration with the Primary Care Practitioner if they believe such consultation is necessary for an accurate assessment. Family Living Provider Agencies have the option of having the subcontracted caregiver complete the HAT instead of the nurse or PCP, if the caregiver is comfortable doing so. However, the agency nurse must be available to assist the caregiver upon request. (c) For newly allocated individuals, the HAT and the MAAT must be completed within seventy-two (72) hours of admission into direct services or two weeks following the initial ISP, whichever comes first. (d) For individuals already in services, the HAT and the MAAT must be completed at least fourteen (14) days prior to the annual ISP meeting and submitted to all members of the interdisciplinary team. The HAT must also be completed at the time of any significant change in clinical condition and upon return from any hospitalizations. In addition to annually, the MAAT must be completed at the time of any significant change in clinical condition, when a medication regime or route change requires delivery by licensed or certified staff, or when an individual has completed additional training designed to improve their skills to support self-administration (see DDSD Medication Assessment and Delivery Policy). (e) 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 members; objective information including vital signs, physical section of ISP the individual is required to have a plan. No evidence of a plan found.

- Psychiatric
  - Individual #2 - As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found.

- Crisis Plans/Medical Emergency Response Plans
  - Asthma
    - Individual #3 - As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found.

- Diabetes
  - Individual #2 - As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found.

- Gastrointestinal
  - Individual #2 - As indicated by the IST section of ISP the individual is required to have a plan. No evidence of a plan found.
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.

(2) Health related plans
(a) For individuals with chronic conditions that have the potential to exacerbate into a life-threatening situation, a medical crisis prevention and intervention plan must be written by the nurse or other appropriately designated healthcare professional.
(b) Crisis prevention and intervention plans must be written in user-friendly language that is easily understood by those implementing the plan.
(c) The nurse shall also document training regarding the crisis prevention and intervention plan delivered to agency staff and other team members, clearly indicating competency determination for each trainee.
(d) If the individual receives services from separate agencies for community living and community inclusion services, nurses from each agency shall collaborate in the development of and training delivery for crisis prevention and intervention plans to assure maximum consistency across settings.

(3) For all individuals with a HAT score of 4, 5 or 6, the nurse shall develop a comprehensive healthcare plan that includes health related supports identified in the ISP (The healthcare plan is the equivalent of a nursing care plan; two separate documents are not required nor recommended):
(a) Each healthcare plan must include a statement of the person’s healthcare needs and list measurable goals to be achieved.
through implementation of the healthcare plan. Needs statements may be based upon supports needed for the individual to maintain a current strength, ability or skill related to their health, prevention measures, and/or supports needed to remediate, minimize or manage an existing health condition.

(b) Goals must be measurable and shall be revised when an individual has met the goal and has the potential to attain additional goals or no longer requires supports in order to maintain the goal.

(c) Approaches described in the plan shall be individualized to reflect the individual’s unique needs, provide guidance to the caregiver(s) and designed to support successful interactions. Some interventions may be carried out by staff, family members or other team members, and other interventions may be carried out directly by the nurse – persons responsible for each intervention shall be specified in the plan.

(d) Healthcare plans shall be written in language that will be easily understood by the person(s) identified as implementing the interventions.

(e) The nurse shall also document training on the healthcare plan delivered to agency staff and other team members, clearly indicating competency determination for each trainee. If the individual receives services from separate agencies for community living and community inclusion services, nurses from each agency shall collaborate in the development of and training delivery for healthcare plans to assure maximum consistency across settings.

(f) Healthcare plans must be updated to reflect relevant discharge orders whenever an individual returns to services following a hospitalization.
(g) All crisis prevention and intervention plans and healthcare plans shall include the individual’s name and date on each page and shall be signed by the author.
(h) Crisis prevention and intervention plans as well as healthcare plans shall be reviewed by the nurse at least quarterly, and updated as needed.

(4) General Nursing Documentation
(a) The nurse shall complete 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 shall be documented whether they occur by phone or in person.
(b) For individuals with a HAT score of 4, 5 or 6, or who have identified health concerns in their ISP, the nurse shall provide the interdisciplinary team with a quarterly report that indicates current health status and progress to date on health related ISP desired outcomes and action plans as well as progress toward goals in the healthcare plan.

CHAPTER 5 IV. COMMUNITY INCLUSION SERVICES PROVIDER AGENCY REQUIREMENTS
B. IDT Coordination
(1) Community Inclusion Services Provider Agencies shall participate on the IDT as specified in the ISP Regulations (7.26.5 NMAC), and shall ensure direct support staff participation as needed to plan effectively for the individual; and
(2) Coordinate with the IDT to ensure that each individual participating in Community
Inclusion Services who has a score of 4, 5, or 6 on the HAT has a Health Care Plan developed by a licensed nurse, and if applicable, a Crisis Prevention/Intervention Plan.

**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.
<table>
<thead>
<tr>
<th>Tag # 1A29  Complaints / Grievances - Acknowledgement</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
</table>
| **NMAC 7.26.3.6**
A These regulations set out rights that the department expects all providers of services to individuals with developmental disabilities to respect. These regulations are intended to complement the department's Client Complaint Procedures (7 NMAC 26.4) [now 7.26.4 NMAC]. |
| Based on record review, the Agency failed to provide documentation, the complaint procedure had been made available to individuals or their legal guardians for 2 of 3 individuals. |
| • Grievance/Complaint Procedure Acknowledgement (#1 & 3) |
| **NMAC 7.26.3.13 Client Complaint Procedure Available.** A complainant may initiate a complaint as provided in the client complaint procedure to resolve complaints alleging that a service provider has violated a client’s rights as described in Section 10 [now 7.26.3.10 NMAC]. The department will enforce remedies for substantiated complaints of violation of a client’s rights as provided in client complaint procedure. [09/12/94; 01/15/97; Recompiled 10/31/01] |
| **NMAC 7.26.4.13 Complaint Process:**
A. (2). The service provider’s complaint or grievance procedure shall provide, at a minimum, that: (a) the client is notified of the service provider’s complaint or grievance procedure |

Provider:
State your Plan of Correction for the findings in this Tag above this line.

Enter your Quality Assurance/Quality Improvement processes below the line.
### Standard of Care

**Deficiencies**

Based on record review, the Agency failed to provide written or electronic documentation as evidence for each unit billed for Adult Habilitation Services for 1 of 3 individuals.

**Agency Plan of Correction, On-going QA/QI & Responsible Party**

**Date Due**

#### CMS Assurance – Financial Accountability

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 # 5I44 Adult Habilitation Reimbursement

<table>
<thead>
<tr>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>Based on record review, the Agency failed to provide written or electronic documentation as evidence for each unit billed for Adult Habilitation Services for 1 of 3 individuals.</td>
</tr>
</tbody>
</table>

**Individual #2**

- March 2012
- The Agency billed 292 units of Adult Habilitation (T2021 U1) from 3/14/2012 through 03/30/2012. Documentation did not contain the required elements on 3/15/2012 and 03/26/2012. Documentation accounted for 48 units.
  - The signature or authenticated name of staff providing the service.

**Provider:**

State your Plan of Correction for the findings in this Tag above this line.

Enter your Quality Assurance/Quality Improvement processes below the line.

**Provider:**

State your Plan of Correction for the findings in this Tag above this line.

Enter your Quality Assurance/Quality Improvement processes below the line.

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Q.12.4.DDW.D0974.2.001.RTN.1.171

23
DOCUMENTATION REQUIREMENTS:
Providers must maintain all records necessary to fully disclose the extent of the services provided to the Medicaid recipient. Services that have been billed to Medicaid, but are not substantiated in a treatment plan and/or patient records for the recipient are subject to recoupment.


CHAPTER 5 XVI. REIMBURSEMENT

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

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

(2) Adult Habilitation Services can be provided with any other services, insofar as the services are not reported for the same hours on the same day, except that Therapy Services and Case Management may be provided and billed for the same hours.
Date: August 24, 2012

To: Patsy Romero, Chief Operating Officer
Provider: Easter Seals Santa Maria el Mirador
Address: 10 A Van Nu Po
State/Zip: Santa Fe, New Mexico 87508

E-mail Address: promero@eselm.org

Region: Northeast
Survey Date: May 15 – 16, 2012
Program Surveyed: Developmental Disabilities Waiver
Service Surveyed: Community Living Supports (Supported Living & Independent Living) & Community Inclusion Supports (Adult Habilitation & Supported Employment)
Survey Type: Routine

Dear Ms. Romero:

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 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,

Crystal Lopez-Beck
Plan of Correction Coordinator
Quality Management Bureau/DHI