Dear Ms. Rishel Brakey,

The Division of Health Improvement/Quality Management Bureau has completed a quality review 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.

Quality Management Approval Rating:
The Division of Health Improvement is pleased to issue your new agency a “PROVISIONAL” rating for compliance with DDSD Standards and regulations. As part of your Provisional rating, QMB will conduct an additional annual review prior to the end of your current provider agreement. The outcome of that review will be used in determining future DHI ratings.

Plan of Correction:
The attached Report of Findings identifies deficiencies found during your agency’s survey. You are required to complete and implement a Plan of Correction (POC). Please submit your agency’s Plan of Correction (POC) in the
space on the two right columns of the Report of Findings. See attachment A for additional guidance in completing the POC. The response is due to the parties below within 10 working days of the receipt of this letter:

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 within 45 working days. If your plan of correction is denied, you must resubmit a revised plan ASAP for approval. All remedies must still be completed within 45 working days of the original submission.

Failure to submit, complete or implement your POC within the required time frames will 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 determination of noncompliance (finding) you have 10 working 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

A request for an IRF will not delay the implementation of your Plan of Correction which must be completed within 45 working days. Providers may not appeal the nature or interpretation of the standard or regulation, the team composition, sampling methodology or the Scope and Severity of the finding.

If the IRF approves the change or removal of a finding, you will be advised of any changes.

This IRF process is separate and apart from the Informal Dispute Resolution (IDR) and Fair Hearing Process for Sanctions from DOH.

Please call the Team Leader at 505-699-9356, if you have questions about the survey or the report. Thank you for your cooperation and for the work you perform.

Sincerely,

Cristal Lopez-Beck, BA
Team Lead/Healthcare Surveyor
Division of Health Improvement
Quality Management Bureau
Survey Process Employed:

Entrance Conference Date: December 29, 2009
Present:

Mandy’s Special Farm
Heidi Rishel Brakey, Executive Director

DOH/DHI/QMB
Crystal Lopez-Beck, BA, Team Lead/Healthcare Surveyor
Tony Fragua, BFA, Healthcare Surveyor

Exit Conference Date: December 30, 2009
Present:

Mandy’s Special Farm
Heidi Rishel Brakey, Executive Director
Griselda Valenzuela, Service Coordinator

DOH/DHI/QMB
Crystal Lopez-Beck, BA, Team Lead/Healthcare Surveyor
Nadine Romero, LBSW, Healthcare Surveyor

Homes Visited Number: 1
Administrative Locations Visited Number: 1
Total Sample Size Number: 3
0 - Jackson Class Members
3 - Non-Jackson Class Members
3 - Supported Living

Persons Served Interviewed Number: 1
Persons Served Observed Number: 2 (2 Individuals were unavailable during the on-site survey)

Records Reviewed (Persons Served) Number: 3

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
- Nursing personnel files
- Evacuation Drills
- Quality Improvement/Quality Assurance Plan

CC: Distribution List: DOH - Division of Health Improvement
DOH - Developmental Disabilities Supports Division
DOH - Office of Internal Audit
HSD - Medical Assistance
Provider Instructions for Completing the QMB Plan of Correction (POC) Process

- After a QMB Quality Review, your Survey Report will be sent to you via certified mail. You may request that it also be sent to you electronically by calling George Perrault, Plan of Correction Coordinator at 505-222-8647.
- Within 10 business days of the date you received your survey report, you must develop and send your Plan of Correction response to the QMB office. (Providers who do not pick up their mail will be referred to the Internal Review Committee [IRC]).
- For each Deficiency in your Survey Report, include specific information about HOW you will correct each Deficiency, WHO will fix each Deficiency (“Responsible Party”), and by WHEN (“Date Due”).
- Your POC must not only address HOW, WHO and WHEN each Deficiency will be corrected, but must also address overall systemic issues to prevent the Deficiency from reoccurring, i.e., Quality Assurance (QA). Your description of your QA must include specifics about your self-auditing processes, such as HOW OFTEN you will self-audit, WHO will do it, and WHAT FORMS will be used.
- Corrective actions should be incorporated into your agency’s Quality Assurance/Quality Improvement policies and procedures.
- You may send your POC response electronically to George.Perrault@state.nm.us, by fax (505-222-8661), or by postal mail.
- Do not send supporting documentation to QMB until after your POC has been approved by QMB.
- QMB will notify you if your POC has been “Approved” or “Denied”.
- Whether your POC is “Approved” or “Denied”, you have a maximum of 45 business days to correct all survey Deficiencies from the date of receipt of your Survey Report. If your POC is “Denied” it must be revised and resubmitted ASAP, as the 45 working day limit is in effect. Providers whose revised POC is denied will be referred to the IRC.
- The POC must be completed on the official QMB Survey Report and Plan of Correction Form, unless approved in advance by the POC Coordinator.
- The following Deficiencies must be corrected within the deadlines below (after receipt of your Survey Report):
  - CCHS and EAR: 10 working days
  - Medication errors: 10 working days
  - IMS system/training: 20 working days
  - ISP related documentation: 30 working days
  - DDSD Training 45 working days
- If you have questions about the POC process, call the QMB POC Coordinator, George Perrault at 505-222-8647 for assistance.
- For Technical Assistance (TA) in developing or implementing your POC, contact your local DDSD Regional Office.
- Once your POC has been approved by QMB, the POC may not be altered or the dates changed.
- Requests for an extension or modification of your POC (post approval) must be made in writing and submitted to the POC Coordinator at QMB, and are approved on a case-by-case basis.
- When submitting supporting documentation, organize your documents by Tag #s, and annotate or label each document using Individual #s.
- Do not submit original documents, hard copies or scanned and electronically submitted copies are fine. Originals must be maintained in the agency/client file(s) as per DDSD Standards.
- Failure to submit, complete or implement your POC within the required timeframes will result in a referral to the IRC and the possible imposition of a $200 per day Civil Monetary Penalty until it is received, completed and/or implemented.
QMB Scope and Severity Matrix of survey results

Each deficiency in your Report of Findings is scored on a Scope and Severity Scale. The culmination of each deficiency’s Scope and Severity is used to determine degree of compliance to standards and regulations and level of QMB Certification.

<table>
<thead>
<tr>
<th>Scope</th>
<th>Isolated 01% - 15%</th>
<th>Pattern 16% - 79%</th>
<th>Widespread 80% - 100%</th>
</tr>
</thead>
<tbody>
<tr>
<td>High Impact</td>
<td>Immediate Jeopardy to individual health and or safety</td>
<td>J.</td>
<td>K.</td>
</tr>
<tr>
<td>Medium Impact</td>
<td>No Actual Harm Potential for more than minimal harm</td>
<td>D.</td>
<td>E.</td>
</tr>
<tr>
<td>Low Impact</td>
<td>No Actual Harm Minimal potential for harm.</td>
<td>A.</td>
<td>B.</td>
</tr>
</tbody>
</table>

Scope and Severity Definitions:

**Key to Scope scale:**

Isolated:
A deficiency that is limited to 1% to 15% of the sample, usually impacting no more than one or two individuals in the sample.

Pattern:
A deficiency that impacts a number or group of individuals from 16% to 79% of the sample is defined as a pattern finding. Pattern findings suggest the need for system wide corrective actions.

Widespread:
A deficiency that impacts most or all (80% to 100%) of the individuals in the sample is defined as widespread or pervasive. Widespread findings suggest the need for system wide corrective actions as well as the need to implement a Continuous Quality Improvement process to improve or build infrastructure. Widespread findings must be referred to the Internal Review Committee for review and possible actions or sanctions.

**Key to Severity scale:**

Low Impact Severity: (Blue)
Low level findings have no or minimal potential for harm to an individual. Providers that have no findings above a “C” level may receive a “Quality” Certification approval rating from QMB.

Medium Impact Severity: (Tan)
Medium level findings have a potential for harm to an individual. Providers that have no findings above a “F” level and/or no more than two F level findings and no F level Conditions of Participation may receive a “Merit” Certification approval rating from QMB.

High Impact Severity: (Green or Yellow)
High level findings are when harm to an individual has occurred. Providers that have no findings above “I” level may only receive a “Standard” Approval rating from QMB and will be referred to the IRC.

High Impact Severity: (Yellow)
“J, K, and L” Level findings:
This is a finding of Immediate Jeopardy. If a provider is found to have “I” level findings or higher, with an outcome of Immediate Jeopardy, including repeat findings or Conditions of Participation they will be referred to the Internal Review Committee.

### The QMB Approval Rating

The QMB approval rating is the provider incentive to encourage quality service and correlates the review outcome with the QMB review frequency and its recommendation to DDSD to determine the length of the provider agreement. The “Approval rating” is based on the Scope and Severity of the review findings. There are five levels of “Approval” that a provider may receive. They are:

**“Quality” Approval Rating:**
The QMB DD Manager will review the Report of Findings and determine if the provider qualifies for a “Quality” Rating. To qualify for a QMB “Quality” rating of approval and a three (3) year QMB review cycle and provider agreement recommendation, the provider must not have any findings that are a condition of participation and no findings of “F” level or higher on the Scope and Severity Matrix with no more than three (3) D or E level findings.

**“Merit” Approval Rating:**
The QMB DD Manager will review the Report of Findings and determine if the provider qualifies for a “Merit” Rating. To qualify for a QMB “Merit” rating of approval and a two (2) year QMB review cycle and provider agreement recommendation, the provider must not have more than three (3) findings that are a condition of participation and no more than three (3) “F” level findings with no findings of a “G” level or higher on the Scope and Severity Matrix and no more that six (6) D or E level findings.

**“Standard” Approval Rating:**
The QMB DD Manager will review the Report of Findings and determine if the provider qualifies for a “Standard” Rating. To qualify for a QMB “Standard” rating of approval and a one (1) year QMB review cycle and provider agreement recommendation, the provider must not have more than six (6) findings that are a condition of participation and no more than six (6) “F” level findings with no findings of a “G” level or higher on the Scope and Severity Matrix and no more that six (6) D or E level findings.

**“Sub-Standard” Approval Rating:**
The QMB DD Manager will review the Report of Findings and determine if the provider has “Sub-standard” performance. To qualify for a QMB “Sub-Standard” rating of approval and a three to six month QMB review cycle, with a referral to the Internal Review Committee and provider agreement recommendation, the provider may have any of the following findings:

- seven (7) or more findings that are a condition of participation
- seven (7) or more “F” level findings
- any findings of a “G” level or higher
- nine (9) or more D or E level findings

A referral to the IRC is required for any “Sub-standard” rating. Depending upon the egregious nature of the findings the IRC shall take appropriate sanction actions up to and including contract termination.

**“Provisional” Approval Rating:**
New DD service providers may qualify for a QMB “Provisional” Approval Rating upon successfully completing their initial QMB Quality Survey. The QMB DD Manager will review the Report of Findings and determine if the provider has achieved at least a standard rating of approval. If successful, the provider may receive a one (1) year contract extension. QMB will notify the DDSD Contract unit of the “Provisional” approval rating.
Attachment C

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

Introduction:
Throughout the process, surveyors are openly communicating with providers. Open communication means that surveyors have clarified issues and/or requested missing information before completing the review. Regardless, there may still be instances where the provider disagrees with a specific finding.

To informally dispute a finding the provider must request in writing an Informal Reconsideration of the Finding (IRF) to the QMB Deputy Bureau Chief within 10 working days of receipt of the final report.

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) and must specify in detail the request for reconsideration and why the finding is inaccurate. The IRF request must include all supporting documentation or evidence that was not previously reviewed during the survey process.

The following limitations apply to the IRF process:

- The request for an IRF and all supporting evidence must be received in 10 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 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 Scope and Severity of a finding.
- 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 QMB Quality Approval Rating and the length of their DDSD provider contract.

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

QMB has 30 working days to complete the review and notify the provider of the decision. The request will be reviewed by the IRF committee. 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 successfully reconsidered, it will be noted and will be removed or modified from the report. 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.

Administrative Review Process:
If a Provider desires to challenge the decision of the IRF committee they may request an Administrative Review by the DHI and DDSD Director. The Request must be made in writing to the QMB Bureau Chief and received within 5 days of notification from the IRF decision.

Regarding IRC Sanctions:
The Informal Reconsideration of the Finding process is a separate process specific to QMB Survey Findings and should not be confused with any process associated with IRC Sanctions.

If a Provider desires to Dispute or Appeal an IRC Sanction that is a separate and different process. Providers may choose the Informal Dispute Resolution Process or the Formal Medicaid Fair Hearing Process to dispute or appeal IRC sanctions, please refer to the DOH Sanction policy and section 39 of the provider contract agreement.
Tag # 1A08 Agency Case File

Scope and Severity Rating: B

Based on record review, the Agency failed to maintain at the administrative office a confidential case file for 2 of 3 individuals.

Review of the Agency individual case files revealed the following items were not found, incomplete, and/or not current:

- ISP Signature Page (#2)
- Addendum A (#2)
- Positive Behavioral Crisis Plan (#2)
- Occupational Therapy Plan (#3)


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.

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:

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.
<table>
<thead>
<tr>
<th>Tag # 1A09</th>
<th>Medication Delivery (MAR) - Routine Medication</th>
<th>Scope and Severity Rating: F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</td>
<td>Medication Administration Records (MAR) were reviewed for the months of September, October &amp; November 2009.</td>
<td></td>
</tr>
</tbody>
</table>

**CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS:** The objective of these standards is to establish Provider Agency policy, procedure and reporting requirements for DD Medicaid Waiver program. These requirements apply to all such Provider Agency staff, whether directly employed or subcontracting with the Provider Agency. Additional Provider Agency requirements and personnel qualifications may be applicable for specific service standards.

**E. Medication Delivery:** Provider Agencies that provide Community Living, Community Inclusion or Private Duty Nursing services shall have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in accordance with DDSD Medication Assessment and Delivery Policy and Procedures, the Board of Nursing Rules and Board of Pharmacy standards and regulations.

(2) When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) shall be maintained and include:

- (a) The name of the individual, a transcription of the physician’s written or licensed health care provider’s prescription including the brand and generic name of the medication, diagnosis for which the medication is prescribed;
- (b) Prescribed dosage, frequency and method/route of administration, times and dates of administration;
- (c) Initials of the individual administering or assisting with the medication;
- (d) Explanation of any medication irregularity;
- (e) Documentation of any allergic reaction or adverse medication effect; and

Medication Administration Records contain the following medications. No Physician's Orders were found for the following medications:

- Nitrofurantion Monomac 100mg (2 times daily)
- Sertraline HCL 50mg (1 time daily)

As indicated by the Medication Administration Records the individual is taking Chlorhexidine Oral Rinse 473 (2 times weekly). According to the Physician's Orders, Chlorhexidine Oral Rinse 0.12% is to be taken 3 times weekly. Medication Administration Record & Physician’s Orders do not match.

Medication Administration Records contain the following medications. No Physician’s Orders were found for the following medications:

- Nitrofurantion Monomac 100mg (2 times daily)

October 2009

As indicated by the Medication Administration Records the individual is taking Chlorhexidine Oral Rinse 473 (1 time weekly). According to the Physician’s Orders, Chlorhexidine Oral Rinse 0.12% is to be taken 3 times weekly. Medication Administration Record & Physician’s Orders do not match.

Medication Administration Records contain the following medications. No Physician’s Orders were found for the following medications:

- Nitrofurantion Monomac 100mg (2 times daily)

- Sertraline HCL 50mg (1 time daily)
(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 administering 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

Individual #2
November 2009
Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries:

- Venlafaxine (2 times daily) – Blank 11/20 (8:30PM)

Individual #3
October 2009
During on-site survey Medication Administration Records were requested for the month of October 2009. As of December 30, 2009, Medication Administration Records had not been provided. When Service Coordinator, #51, was asked about the missing MAR she stated, “the medication book was lost, along with the MAR for October.”

November 2009
As indicated by the Medication Administration Records the individual is to take Vitamin C 500mg (1 time daily). According to the Physician’s Orders, Vitamin C 1000mg is to be taken 1 time daily. Medication Administration Record & Physician’s Orders do not match.
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.
### Tag # 1A09 Medication Delivery - PRN Medication

|---------------------------------------------------------------------------|

**CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS:** The objective of these standards is to establish Provider Agency policy, procedure and reporting requirements for DD Medicaid Waiver program. These requirements apply to all such Provider Agency staff, whether directly employed or subcontracting with the Provider Agency. Additional Provider Agency requirements and personnel qualifications may be applicable for specific service standards.

**E. Medication Delivery:** Provider Agencies that provide Community Living, Community Inclusion or Private Duty Nursing services shall have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in accordance with DDSD Medication Assessment and Delivery Policy and Procedures, the Board of Nursing Rules and Board of Pharmacy standards and regulations.

(2) When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) shall be maintained and include:

<table>
<thead>
<tr>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>(a) The name of the individual, a transcription of the physician’s written or licensed health care provider’s prescription including the brand and generic name of the medication, diagnosis for which the medication is prescribed;</td>
</tr>
<tr>
<td>(b) Prescribed dosage, frequency and method/route of administration, times and dates of administration;</td>
</tr>
<tr>
<td>(c) Initials of the individual administering or assisting with the medication;</td>
</tr>
<tr>
<td>(d) Explanation of any medication irregularity;</td>
</tr>
<tr>
<td>(e) Documentation of any allergic reaction or adverse medication effect;</td>
</tr>
</tbody>
</table>

Based on record review, the Agency failed to maintain PRN Medication Administration Records which contained all elements required by standard for 2 of 3 Individuals.

**Individual #1**

**October 2009**

- No Signs/Symptoms were noted on the Medication Administration Record for the following PRN medication:
  - Stool Softener (Colace) 100mg – PRN – 10/23 (given 1 time daily)

- No Effectiveness was noted on the Medication Administration Record for the following PRN medication:
  - Stool Softener (Colace) 100mg – PRN – 10/23 (given 1 time daily)

**November 2009**

- No Signs/Symptoms were noted on the Medication Administration Record for the following PRN medication:
  - Seroquel 25mg – PRN – 11/20 (given 1 time daily)
  - Stool Softener (Colace) 100mg – PRN – 11/23 (given 2 times daily)

- No Effectiveness was noted on the Medication Administration Record for the following PRN medication:
  - Seroquel 25mg – PRN – 11/20 & 23 (given 1 time daily)
  - Stool Softener (Colace) 100mg – PRN – 11/22 (given 1 time daily) & 11/23 (given 2 times daily)

**Individual #2**

**September 2009**

- No Signs/Symptoms were noted on the Medication Administration Record for the following PRN medication:
  - Seroquel 25mg – PRN – 11/20 & 23 (given 1 time daily)
  - Stool Softener (Colace) 100mg – PRN – 11/22 (given 1 time daily) & 11/23 (given 2 times daily)
(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 administering 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

No Signs/Symptoms were noted on the Medication Administration Record for the following PRN medication:

- Peptobismol – PRN – 09/19 (given 1 time daily)

No Effectiveness was noted on the Medication Administration Record for the following PRN medication:

- Peptobismol – PRN – 09/19 (given 1 time daily)

**October 2009**

No Effectiveness was noted on the Medication Administration Record for the following PRN medication:

- Peptobismol – PRN – 10/11 & 14 (given 1 time daily)

**November 2009**

No Effectiveness was noted on the Medication Administration Record for the following PRN medication:

- Acetaminophen 500mgl – PRN – 11/15 (given 1 time daily)
(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.

Department of Health

Developmental Disabilities Supports Division (DDSD) Medication Assessment and Delivery Policy - Eff. November 1, 2006

F. PRN Medication

3. Prior to self-administration, self-administration with physical assist or assisting with delivery of PRN medications, the direct support staff must contact the agency nurse to describe observed symptoms and thus assure that the PRN medication is being used according to instructions given by the ordering PCP. In cases of fever, respiratory distress (including coughing), severe pain, vomiting, diarrhea, change in responsiveness/level of consciousness, the nurse must strongly consider the need to conduct a face-to-face assessment to assure that the PRN does not mask a condition better treated by seeking medical attention. This does not apply to home based/family living settings where the provider is related by affinity or by consanguinity to the individual.
4. The agency nurse shall review the utilization of PRN medications routinely. Frequent or escalating use of PRN medications must be reported to the PCP and discussed by the Interdisciplinary for changes to the overall support plan (see Section H of this policy).

H. Agency Nurse Monitoring
1. Regardless of the level of assistance with medication delivery that is required by the individual or the route through which the medication is delivered, the agency nurses must monitor the individual's response to the effects of their routine and PRN medications. The frequency and type of monitoring must be based on the nurse's assessment of the individual and consideration of the individual's diagnoses, health status, stability, utilization of PRN medications and level of support required by the individual's condition and the skill level and needs of the direct care staff. Nursing monitoring should be based on prudent nursing practice and should support the safety and independence of the individual in the community setting. The health care plan shall reflect the planned monitoring of the individual's response to medication.

Department of Health Developmental Disabilities Supports Division (DDSD) - Procedure Title: Medication Assessment and Delivery Procedure Eff Date: November 1, 2006
C. 3. Prior to delivery of the PRN, direct support staff must contact the agency nurse to describe observed symptoms and thus assure that the PRN is being used according to instructions given by the ordering PCP. In cases of fever, respiratory distress (including coughing), severe pain, vomiting, diarrhea, change in responsiveness/level of consciousness, the nurse must strongly consider the need to conduct a face-to-face assessment to assure that the PRN does not mask a condition better treated by seeking medical attention.
(References: Psychotropic Medication Use Policy, Section D, page 5 Use of PRN Psychotropic Medications; and, Human Rights Committee Requirements Policy, Section B, page 4 Interventions Requiring Review and Approval – Use of PRN Medications).

a. Document conversation with nurse including all reported signs and symptoms, advice given and action taken by staff.

4. Document on the MAR each time a PRN medication is used and describe its effect on the individual (e.g., temperature down, vomiting lessened, anxiety increased, the condition is the same, improved, or worsened, etc.).
Tag # 1A11 (CoP)  Transportation P&P


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.

G. Transportation: Provider agencies that provide Community Living, Community Inclusion or Non-Medical Transportation services shall have a written policy and procedures regarding the safe transportation of individuals in the community, which comply with New Mexico regulations governing the operation of motor vehicles to transport individuals, and which are consistent with DDSD guidelines issued July 1, 1999 titled “Client Transportation Safety”. The policy and procedures must address at least the following topics:

1. Drivers' requirements,
2. Individual safety, including safe locations for boarding and disembarking passengers, appropriate responses to hazardous weather and other adverse driving conditions,
3. Vehicle maintenance and safety inspections,
4. Staff training regarding the safe operation of the vehicle, assisting passengers and safe lifting procedures,
5. Emergency Plans, including vehicle evacuation techniques,
6. Documentation, and
7. Accident Procedures

Scope and Severity Rating: F

Based on record review, the Agency failed to have a written policies and procedures regarding the safe transportation of individuals in the community, which comply with New Mexico regulations governing the operation of motor vehicles to transport individuals.

Review of Agency's policies and procedures indicated the following elements were not found:

2. Individual safety, including safe locations for boarding and disembarking passengers, appropriate responses to hazardous weather and other adverse driving conditions,
3. Vehicle maintenance and safety inspections,
5. Emergency Plans, including vehicle evacuation techniques,
6. Documentation, and
7. Accident Procedures
Staff Policy  
**Eff Date:** March 1, 2007  

**II. POLICY STATEMENTS:**  

1. Staff providing direct services shall complete safety training within the first thirty (30) days of employment and before working alone with an individual receiving services. The training shall address at least the following:  

   1. Operating a fire extinguisher  
   2. Proper lifting procedures  
   3. General vehicle safety precautions (e.g., pre-trip inspection, removing keys from the ignition when not in the driver’s seat)  
   4. Assisting passengers with cognitive and/or physical impairments (e.g., general guidelines for supporting individuals who may be unaware of safety issues involving traffic or those who require physical assistance to enter/exit a vehicle)  
   5. Operating wheelchair lifts (if applicable to the staff’s role)  
   6. Wheelchair tie-down procedures (if applicable to the staff’s role)  
   7. Emergency and evacuation procedures (e.g., roadside emergency, fire emergency)
Tag # 1A11 (CoP)  Transportation Training | Scope and Severity Rating:  E
---|---

**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.

**G. Transportation:** Provider agencies that provide Community Living, Community Inclusion or Non-Medical Transportation services shall have a written policy and procedures regarding the safe transportation of individuals in the community, which comply with New Mexico regulations governing the operation of motor vehicles...

---

**Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy**

- **Training Requirements for Direct Service Agency Staff Policy**
  - **Eff Date:** March 1, 2007

**II. POLICY STATEMENTS:**

1. Staff providing direct services shall complete safety training within the first thirty (30) days of employment and before working alone with an individual receiving services. The training shall address at least the following:
   1. Operating a fire extinguisher
   2. Proper lifting procedures
   3. General vehicle safety precautions (e.g., pre-trip inspection, removing keys from the ignition when not in the driver’s seat)
   4. Assisting passengers with cognitive and/or physical impairments (e.g., general guidelines for supporting individuals who may be unaware of safety issues involving traffic or those who require physical assistance to enter/exit a vehicle)

---

Based on record review, the Agency failed to provide staff training regarding the safe operation of the vehicle, assisting passengers and safe lifting procedures for 5 of 11 Direct Service Personnel.

No documented evidence was found of the following required training:

- Transportation (DSP #43, 44, 45, 47 & 50)
5. Operating wheelchair lifts (if applicable to the staff's role)
6. Wheelchair tie-down procedures (if applicable to the staff's role)
7. Emergency and evacuation procedures (e.g., roadside emergency, fire emergency)
<table>
<thead>
<tr>
<th>Tag # 1A15 Healthcare Documentation</th>
<th>Scope and Severity Rating: F</th>
</tr>
</thead>
</table>

**CHAPTER 1. III. PROVIDER AGENCY DOCUMENTATION OF SERVICE DELIVERY AND LOCATION - Healthcare Documentation by Nurses For Community Living Services, Community Inclusion Services and Private Duty Nursing Services:** Nursing services must be available as needed and documented for Provider Agencies delivering Community Living Services, Community Inclusion Services and Private Duty Nursing Services.

**Chapter 1. III. E. (1 - 4) (1) Documentation of nursing assessment activities**

(a) The following hierarchy shall be used to determine which provider agency is responsible for completion of the HAT and MAAT and related subsequent planning and training:

(i) Community living services provider agency;
(ii) Private duty nursing provider agency;
(iii) Adult habilitation provider agency;
(iv) Community access provider agency; and
(v) Supported employment provider agency.

(b) The provider agency must arrange for their nurse to complete the Health Assessment Tool (HAT) and the Medication Administration Assessment Tool (MAAT) on at least an annual basis for each individual receiving community living, community inclusion or private duty nursing services, unless the provider agency arranges for the individual’s Primary Care Practitioner (PCP) to voluntarily complete these assessments in lieu of the agency nurse. 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

Based on record review, the Agency failed to maintain the required documentation in the Individuals Agency Record as required per standard for 3 of 3 individual

The following were not found, incomplete and/or not current:

- **Quarterly Nursing Review of HCP/Crisis Plans:**
  - None found for 02/20/2009 – 07/19/2009 (#2)

- **Special Health Care Needs:**
  - Nutritional Plan
    - Individual #3 - As indicated by the IST section of ISP the individual is required to have a plan.

- **Crisis Plans**
  - Gastrointestinal
    - Individual #1 - As indicated by the IST section of ISP the individual is required to have a plan.
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 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.
<table>
<thead>
<tr>
<th>Tag #</th>
<th>DSP Training Documents</th>
<th>Scope and Severity Rating: E</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A20</td>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</td>
<td>Based on record review, the Agency failed to ensure that Orientation and Training requirements were met for 5 of 11 Direct Service Personnel.</td>
</tr>
<tr>
<td></td>
<td><strong>CHAPTER 1 IV. GENERAL REQUIREMENTS FOR PROVIDER AGENCY SERVICE</strong></td>
<td>Review of Direct Service Personnel training records found no evidence of the following required DOH/DDSD trainings and certification being completed:</td>
</tr>
<tr>
<td></td>
<td><strong>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.</td>
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<td></td>
<td><strong>C. Orientation and Training Requirements:</strong> Orientation and training for direct support staff and his or her supervisors shall comply with the DDSD/DOH Policy Governing the Training Requirements for Direct Support Staff and Internal Service Coordinators Serving Individuals with Developmental Disabilities to include the following:</td>
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<td>(1) Each new employee shall receive appropriate orientation, including but not limited to, all policies relating to fire prevention, accident prevention, incident management and reporting, and emergency procedures; and</td>
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<td>(2) Individual-specific training for each individual under his or her direct care, as described in the individual service plan, prior to working alone with the individual.</td>
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<tr>
<td></td>
<td>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:</td>
<td></td>
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<tr>
<td></td>
<td>A. Individuals shall receive services from competent and qualified staff…</td>
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<tr>
<td>Tag # 1A22</td>
<td>Staff Competence</td>
<td>Scope and Severity Rating: F</td>
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<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</td>
<td>Based on interview, the Agency failed to ensure that training competencies were met for 3 of 3 Direct Service Personnel.</td>
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</tbody>
</table>

**CHAPTER 1 IV. GENERAL REQUIREMENTS FOR PROVIDER AGENCY SERVICE PERSONNEL:**

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.

**F. Qualifications for Direct Service Personnel:**

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 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.

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[Table rows with specific details about training and competency requirements, as well as examples of direct service personnel's responses to questions about training and individual plans.]

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Report #: Q10.02.32408382.METRO.001.INT.01
(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.
<table>
<thead>
<tr>
<th>Tag # 1A25 (CoP) CCHS</th>
<th>Scope and Severity Rating: D</th>
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<tbody>
<tr>
<td><strong>NMAC 7.1.9.8 CAREGIVER AND HOSPITAL CAREGIVER EMPLOYMENT REQUIREMENTS:</strong></td>
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<tr>
<td><strong>F. Timely Submission:</strong> Care providers shall submit all fees and pertinent application information for all individuals who meet the definition of an applicant, caregiver or hospital caregiver as described in Subsections B, D and K of 7.1.9.7 NMAC, no later than twenty (20) calendar days from the first day of employment or effective date of a contractual relationship with the care provider.</td>
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</table>

Based on record review, the Agency failed to maintain documentation indicating no “disqualifying convictions” or documentation of the timely submission of pertinent application information to the Caregiver Criminal History Screening Program was on file for 2 of 12 Agency Personnel.

**The following Agency Personnel Files contained no evidence of Caregiver Criminal History Screenings:**

- #44 – Date of hire 10/30/2009

**The following Agency Personnel Files contained Caregiver Criminal History Screenings, which were completed after the twenty (20) calendar days from the first day of employment:**

- #47 – Date of hire 11/02/2009 (date fingerprints taken – 12/17/2009)

**NMAC 7.1.9.9 CAREGIVERS OR HOSPITAL CAREGIVERS AND APPLICANTS WITH DISQUALIFYING CONVICTIONS:**

**A. Prohibition on Employment:** A care provider shall not hire or continue the employment or contractual services of any applicant, caregiver or hospital caregiver for whom the care provider has received notice of a disqualifying conviction, except as provided in Subsection B of this section.

**NMAC 7.1.9.11 DISQUALIFYING CONVICTIONS.**

The following felony convictions disqualify an applicant, caregiver or hospital caregiver from employment or contractual services with a care provider:

- A. homicide;
- B. trafficking, or trafficking in controlled substances;
- C. kidnapping, false imprisonment, aggravated assault or aggravated battery;
- D. rape, criminal sexual penetration, criminal sexual contact, incest, indecent exposure, or other related felony sexual offenses;
- E. crimes involving adult abuse, neglect or financial exploitation;
- F. crimes involving child abuse or neglect;
- G. crimes involving robbery, larceny, extortion, burglary, fraud, forgery, embezzlement, credit card fraud, or receiving stolen property; or
- H. an attempt, solicitation, or conspiracy involving any of the felonies in this subsection.
<table>
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<tr>
<th>Tag # 1A26 (CoP) COR / EAR</th>
<th>Scope and Severity Rating: E</th>
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<tbody>
<tr>
<td>NMAC 7.1.12.8 REGISTRY ESTABLISHED; PROVIDER INQUIRY REQUIRED: Upon the effective date of this rule, the department has established and maintains an accurate and complete electronic registry that contains the name, date of birth, address, social security number, and other appropriate identifying information of all persons who, while employed by a provider, have been determined by the department, as a result of an investigation of a complaint, to have engaged in a substantiated registry-referred incident of abuse, neglect or exploitation of a person receiving care or services from a provider. Additions and updates to the registry shall be posted no later than two (2) business days following receipt. Only department staff designated by the custodian may access, maintain and update the data in the registry. A. Provider requirement to inquire of registry. A provider, prior to employing or contracting with an employee, shall inquire of the registry whether the individual under consideration for employment or contracting is listed on the registry. B. Prohibited employment. A provider may not employ or contract with an individual to be an employee if the individual is listed on the registry as having a substantiated registry-referred incident of abuse, neglect or exploitation of a person receiving care or services from a provider. D. Documentation of inquiry to registry. The provider shall maintain documentation in the employee’s personnel or employment records that evidences the fact that the provider made an inquiry to the registry concerning that employee prior to employment. Such documentation must include evidence, based on the response to such inquiry received from the custodian by the provider, that the employee was not listed on the registry as having a substantiated registry-referred incident of abuse, neglect or exploitation.</td>
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<tr>
<td>Based on record review, the Agency failed to maintain documentation in the employee’s personnel records that evidenced inquiry to the Employee Abuse Registry prior to employment for 6 of 12 Agency Personnel. The following Agency Personnel records contained evidence that indicated the Employee Abuse Registry was completed after hire:</td>
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<tr>
<td>• #43 – Date of hire 12/23/2008</td>
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<td>• #44 – Date of hire 10/30/2009</td>
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<tr>
<td>• #45 – Date of hire 06/29/2009</td>
<td></td>
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<tr>
<td>• #47 – Date of hire 11/02/2009</td>
<td></td>
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<tr>
<td>• #48 – Date of hire 09/04/2009</td>
<td></td>
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<tr>
<td>• #51 – Date of hire 08/28/2007</td>
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</tbody>
</table>
E. **Documentation for other staff.** With respect to all employed or contracted individuals providing direct care who are licensed health care professionals or certified nurse aides, the provider shall maintain documentation reflecting the individual's current licensure as a health care professional or current certification as a nurse aide.

F. **Consequences of noncompliance.** The department or other governmental agency having regulatory enforcement authority over a provider may sanction a provider in accordance with applicable law if the provider fails to make an appropriate and timely inquiry of the registry, or fails to maintain evidence of such inquiry, in connection with the hiring or contracting of an employee; or for employing or contracting any person to work as an employee who is listed on the registry. Such sanctions may include a directed plan of correction, civil monetary penalty not to exceed five thousand dollars ($5000) per instance, or termination or non-renewal of any contract with the department or other governmental agency.


**Chapter 1.IV. General Provider Requirements.**

**D. Criminal History Screening:** All personnel shall be screened by the Provider Agency in regard to the employee’s qualifications, references, and employment history, prior to employment. All Provider Agencies shall comply with the Criminal Records Screening for Caregivers 7.1.12 NMAC and Employee Abuse Registry 7.1.12 NMAC as required by the Department of Health, Division of Health Improvement.
### Tag # 1A28 (CoP) Incident Mgt. System - Policy & Procedure

<table>
<thead>
<tr>
<th><strong>Scope &amp; Severity Rating:</strong> F</th>
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<tbody>
<tr>
<td><strong>7.1.13.9 INCIDENT MANAGEMENT SYSTEM REPORTING REQUIREMENTS FOR COMMUNITY BASED SERVICE PROVIDERS:</strong></td>
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<tr>
<td><strong>C. Incident Policies:</strong> All community based service providers shall maintain policies and procedures, which describe the community based service provider’s immediate response to all reported allegations of incidents involving abuse, neglect, or misappropriation of property; all unexpected deaths or natural/expected deaths, and other reportable incidents required as required in Paragraph (2) of Subsection A of 7.1.13.9 NMAC.</td>
<td>Based on interview, the Agency failed to establish and maintain an incident management system, which emphasizes the principles of prevention and staff involvement. When Executive Director, #52, was asked if the Agency conducted internal investigations the following was reported: #52 stated, “Informal investigations are supposed to be done internally but there hasn’t been anything that I feel requires an investigation, per say.” When Service Coordinator/Incident Management Coordinator, #51, was asked if the Agency conducted internal investigations the following was reported: #51 stated, “There is no formal process for the investigation of external Incident Reports.” When #52 was asked if the Agency did any type of tracking &amp; trending of incidents as required by the Incident Management Quality Improvement System for Community Based Service Providers, the following was reported: #52 stated, “No, we do not do any type of tracking or trending. We also do not have an Incident Management Committee at this time.”</td>
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<tr>
<td><strong>NMAC 7.1.13.10 INCIDENT MANAGEMENT SYSTEM REQUIREMENTS:</strong></td>
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<tr>
<td><strong>A. General:</strong> All licensed health care facilities and community based service providers shall establish and maintain an incident management system, which emphasizes the principles of prevention and staff involvement. The licensed health care facility or community based service provider shall ensure that the incident management system policies and procedures requires all employees to be competently trained to respond to, report, and document incidents in a timely and accurate manner.</td>
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<td><strong>B. Training Curriculum:</strong> The licensed health care facility and community based service provider shall provide all employees and volunteers with a written training curriculum on incident policies and procedures for identification, and timely reporting of abuse, neglect, misappropriation of consumers’ property, and where applicable to community based service providers, unexpected deaths or other reportable incidents, within thirty (30) days of the employees’ initial employment, and by annual review not to exceed twelve (12) month intervals. The training curriculum may include computer-</td>
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based training. Periodic reviews shall include, at a minimum, review of the written training curriculum and site-specific issues pertaining to the licensed health care facilities or community based service provider’s facility. Training shall be conducted in a language that is understood by the employee and volunteer.

C. Incident Management System Training
Curriculum Requirements:
(1) The licensed health care facility and community based service provider shall conduct training, or designate a knowledgeable representative to conduct training, in accordance with the written training curriculum that includes but is not limited to:
   (a) an overview of the potential risk of abuse, neglect, misappropriation of consumers’ property;
   (b) informational procedures for properly filing the division's incident management report form;
   (c) specific instructions of the employees' legal responsibility to report an incident of abuse, neglect and misappropriation of consumers’ property.
   (d) specific instructions on how to respond to abuse, neglect, misappropriation of consumers’ property;
   (e) emergency action procedures to be followed in the event of an alleged incident or knowledge of abuse, neglect, misappropriation of consumers’ property; and
   (f) where applicable to employees of community based service providers, informational procedures for properly filing the division's incident management report form for unexpected deaths or other reportable incidents.
<table>
<thead>
<tr>
<th>Tag # 1A28 (CoP)</th>
<th>Incident Mgt. System - Personnel Training</th>
<th>Scope &amp; Severity Rating: D</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>NMAC 7.1.13.10 INCIDENT MANAGEMENT SYSTEM REQUIREMENTS:</strong></td>
<td>Based on record review and interview, the Agency failed to provide documentation verifying completion of Incident Management Training for 2 of 12 Agency Personnel.</td>
<td></td>
</tr>
<tr>
<td><strong>A. General:</strong> All licensed health care facilities and community based service providers shall establish and maintain an incident management system, which emphasizes the principles of prevention and staff involvement. The licensed health care facility or community based service provider shall ensure that the incident management system policies and procedures requires all employees to be competently trained to respond to, report, and document incidents in a timely and accurate manner.</td>
<td>- Incident Management Training (Abuse, Neglect &amp; Misappropriation of Consumers' Property) (#49)</td>
<td></td>
</tr>
<tr>
<td><strong>D. Training Documentation:</strong> All licensed health care facilities and community based service providers shall prepare training documentation for each employee to include a signed statement indicating the date, time, and place they received their incident management reporting instruction. The licensed health care facility and community based service provider shall maintain documentation of an employee's training for a period of at least twelve (12) months, or six (6) months after termination of an employee's employment. Training curricula shall be kept on the provider premises and made available on request by the department. Training documentation shall be made available immediately upon a division representative's request. Failure to provide employee training documentation shall subject the licensed health care facility or community based service provider to the penalties provided for in this rule.</td>
<td><strong>When DSP were asked what two State Agencies must be contacted when there is suspected Abuse, Neglect &amp; Misappropriation of Consumers' Property, the following was reported:</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007</strong></td>
<td>- DSP #40 stated, “DOH.” DSP failed to mention Adult Protective Services.</td>
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<tr>
<td><strong>II. POLICY STATEMENTS:</strong></td>
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<tr>
<td>Tag # 1A28 (CoP) Incident Mgt. System - Parent/Guardian Training</td>
<td>Scope &amp; Severity Rating: F</td>
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<tr>
<td><strong>NMAC 7.1.13.10 INCIDENT MANAGEMENT SYSTEM REQUIREMENTS:</strong></td>
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<td><strong>A. General:</strong> All licensed health care facilities and community based service providers shall establish and maintain an incident management system, which emphasizes the principles of prevention and staff involvement. The licensed health care facility or community based service provider shall ensure that the incident management system policies and procedures requires all employees to be competently trained to respond to, report, and document incidents in a timely and accurate manner.</td>
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<tr>
<td>Based on record review, the Agency failed to provide documentation indicating consumer, family members, or legal guardians had received an orientation packet including incident management system policies and procedural information concerning the reporting of Abuse, Neglect and Misappropriation of Consumers’ Property, for 3 of 3 individuals.</td>
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<tr>
<td><strong>E. Consumer and Guardian Orientation Packet:</strong> Consumers, family members and legal guardians shall be made aware of and have available immediate accessibility to the licensed health care facility and community based service provider incident reporting processes. The licensed health care facility and community based service provider shall provide consumers, family members or legal guardians an orientation packet to include incident management systems policies and procedural information concerning the reporting of abuse, neglect or misappropriation. The licensed health care facility and community based service provider shall include a signed statement indicating the date, time, and place they received their orientation packet to be contained in the consumer’s file. The appropriate consumer, family member or legal guardian shall sign this at the time of orientation.</td>
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<tr>
<td>• Parent/Guardian Incident Management Training (Abuse, Neglect &amp; Misappropriation of Consumers’ Property) (#1, 2 &amp; 3)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tag # 1A31 (CoP) Client Rights/Human Rights</td>
<td>Scope and Severity Rating: E</td>
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<td>---------------------------------------------</td>
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<tr>
<td><strong>7.26.3.11 RESTRICTIONS OR LIMITATION OF CLIENT’S RIGHTS:</strong></td>
<td>Based on record review and interview, the Agency failed to ensure the rights of Individuals were not restricted or limited for 2 of 3 Individuals.</td>
<td></td>
</tr>
<tr>
<td>A. A service provider shall not restrict or limit a client's rights except: (1) where the restriction or limitation is allowed in an emergency and is necessary to prevent imminent risk of physical harm to the client or another person; or (2) where the interdisciplinary team has determined that the client's limited capacity to exercise the right threatens his or her physical safety; or (3) as provided for in Section 10.1.14 [now Subsection N of 7.26.3.10 NMAC].</td>
<td>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. (#1 &amp; 2)</td>
<td></td>
</tr>
<tr>
<td>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.</td>
<td>When #52 was asked if the Agency had documentation of Human Rights approval, the following was reported,</td>
<td></td>
</tr>
<tr>
<td>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]</td>
<td>#52 stated, “Yes we do but we only meet once a year.”</td>
<td></td>
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</tbody>
</table>

**Long Term Services Division**  
**Policy Title: Human Rights Committee**  
**Requirements Eff Date: March 1, 2003**  
**IV. POLICY STATEMENT - Human Rights**  
Committees are required for residential service provider agencies. The purpose of these
committees with respect to the provision of Behavior Supports is to review and monitor the implementation of certain Behavior Support Plans.

Human Rights Committees may not approve any of the interventions specifically prohibited in the following policies:
- Aversive Intervention Prohibitions
- Psychotropic Medications Use
- Behavioral Support Service Provision.

A Human Rights Committee may also serve other agency functions as appropriate, such as the review of internal policies on sexuality and incident management follow-up.

A. HUMAN RIGHTS COMMITTEE ROLE IN BEHAVIOR SUPPORTS

Only those Behavior Support Plans with an aversive intervention included as part of the plan or associated Crisis Intervention Plan need to be reviewed prior to implementation. Plans not containing aversive interventions do not require Human Rights Committee review or approval.

2. The Human Rights Committee will determine and adopt a written policy stating the frequency and purpose of meetings. Behavior Support Plans approved by the Human Rights Committee will be reviewed at least quarterly.

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.

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
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 # 1A36 SC Training

Scope and Severity Rating: C

Based on record review, the Agency failed to ensure that Orientation and Training requirements were met for 1 of 1 Service Coordinators.

Review of Service Coordinators training records found no evidence of the following required DOH/DDSD trainings being completed:

- Pre-Service Manual (SC #51)

**Tag # 1A36 SC Training**


**CHAPTER 1 IV. GENERAL REQUIREMENTS FOR PROVIDER AGENCY SERVICE PERSONNEL:** 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.

**C. Orientation and Training Requirements:**

Orientation and training for direct support staff and his or her supervisors shall comply with the DDSD/DOH Policy Governing the Training Requirements for Direct Support Staff and Internal Service Coordinators Serving Individuals with Developmental Disabilities to include the following:

1. Each new employee shall receive appropriate orientation, including but not limited to, all policies relating to fire prevention, accident prevention, incident management and reporting, and emergency procedures; and
<table>
<thead>
<tr>
<th>Tag # 1A37 Individual Specific Training</th>
<th>Scope and Severity Rating: D</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 ensure that Individual Specific Training requirements were met for 1 of 12 Agency Personnel.</td>
</tr>
<tr>
<td><strong>CHAPTER 1 IV. GENERAL REQUIREMENTS FOR PROVIDER AGENCY SERVICE</strong></td>
<td>Review of personnel records found no evidence of the following:</td>
</tr>
<tr>
<td><strong>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.</td>
<td>• Individual Specific Training (#44)</td>
</tr>
<tr>
<td><strong>C. Orientation and Training Requirements:</strong> Orientation and training for direct support staff and his or her supervisors shall comply with the DDSD/DOH Policy Governing the Training Requirements for Direct Support Staff and Internal Service Coordinators Serving Individuals with Developmental Disabilities to include the following:</td>
<td></td>
</tr>
<tr>
<td>(2) <strong>Individual-specific training</strong> for each individual under his or her direct care, as described in the individual service plan, prior to working alone with the individual.</td>
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</tbody>
</table>

Department of Health (DOH) Developmental Disabilities Supports Division (DDSD) Policy - Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007 - II. POLICY STATEMENTS:

A. Individuals shall receive services from competent and qualified staff.
B. Staff shall complete individual-specific (formerly known as “Addendum B”) training requirements in accordance with the specifications described in the individual service plan (ISP) of each individual served.
### Tag # 6L13 (CoP) - CL Healthcare Reqts.

<table>
<thead>
<tr>
<th>Tag # 6L13 (CoP) - CL Healthcare Reqts.</th>
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</thead>
<tbody>
<tr>
<td><strong>CHAPTER 6. VI. GENERAL REQUIREMENTS FOR COMMUNITY LIVING</strong></td>
</tr>
<tr>
<td><strong>G. Health Care Requirements for Community Living Services.</strong></td>
</tr>
<tr>
<td>(1) The Community Living Service providers shall ensure completion of a HAT for each individual receiving this service. The HAT shall be completed 2 weeks prior to the annual ISP meeting and submitted to the Case Manager and all other IDT Members. A revised HAT is required to also be submitted whenever the individual’s health status changes significantly. For individuals who are newly allocated to the DD Waiver program, the HAT may be completed within 2 weeks following the initial ISP meeting and submitted with any strategies and support plans indicated in the ISP, or within 72 hours following admission into direct services, which ever comes first.</td>
</tr>
<tr>
<td>(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.</td>
</tr>
<tr>
<td>(3) For each individual receiving Community Living Services, the provider agency shall ensure and document the following:</td>
</tr>
<tr>
<td>(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.</td>
</tr>
<tr>
<td>b) That each individual with a score of 4, 5, or 6</td>
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</table>

### Scope and Severity Rating: D

Based on record review, the Agency failed to provide documentation of annual physical examinations and/or other examinations as specified by a licensed physician for 1 of 3 individuals receiving Community Living Services.

- **Auditory Exam**
  ° Individual #2 - As indicated by the documentation reviewed, exam was completed in 09/2008. Follow-up was to be completed annually. No evidence of follow-up found.

- **Abnormal Involuntary Movement Screening**
  ° None found 10/2009 for Seroquel (#2)
on the HAT, has a Health Care Plan developed by a licensed nurse.
(c) That an individual with chronic condition(s)
with the potential to exacerbate into a life
threatening condition, has Crisis Prevention/
Intervention Plan(s) developed by a licensed
nurse or other appropriate professional for
each such condition.
(4) That an average of 3 hours of documented
nutritional counseling is available annually, if
recommended by the IDT.
(5) That the physical property and grounds are free
of hazards to the individual’s health and safety.
(6) In addition, for each individual receiving
Supported Living or Family Living Services, the
provider shall verify and document the following:
(a) The individual has a primary licensed
physician;
(b) The individual receives an annual physical
examination and other examinations as
specified by a licensed physician;
(c) The individual receives annual dental check-
ups and other check-ups as specified by a
licensed dentist;
(d) The individual receives eye examinations as
specified by a licensed optometrist or
ophthalmologist; and
(e) Agency activities that occur as follow-up to
medical appointments (e.g. treatment, visits to
specialists, changes in medication or daily
routine).
<table>
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<tr>
<th>Tag # 6L14 Residential Case File</th>
<th>Scope and Severity Rating: F</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 3 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:</td>
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<tr>
<td>(1) Complete and current ISP and all supplemental plans specific to the individual;</td>
<td>• ISP Signature Page (#2)</td>
</tr>
<tr>
<td>(2) Complete and current Health Assessment Tool;</td>
<td>• Addendum A (#2)</td>
</tr>
<tr>
<td>(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;</td>
<td>• Positive Behavioral Crisis Plan (#2)</td>
</tr>
<tr>
<td>(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);</td>
<td>• Occupational Therapy Plan (#3)</td>
</tr>
<tr>
<td>(5) Data collected to document ISP Action Plan implementation</td>
<td>• Health Assessment Tool (#1)</td>
</tr>
<tr>
<td>(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;</td>
<td>• Special Health Care Needs</td>
</tr>
<tr>
<td>(7) Physician’s or qualified health care providers written orders;</td>
<td>° Nutritional Plan (#3)</td>
</tr>
<tr>
<td>(8) Progress notes documenting implementation of</td>
<td>• Crisis Plan</td>
</tr>
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<td></td>
<td>° Gastrointestinal (#1) (per IST)</td>
</tr>
<tr>
<td></td>
<td>• Health Care Providers Written Orders (#3)</td>
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</table>
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.
<table>
<thead>
<tr>
<th><strong>Tag # 6L26  SL Reimbursement</strong></th>
<th><strong>Scope and Severity Rating: B</strong></th>
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<tbody>
<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</td>
<td>Based on record review, the Agency failed to provide written or electronic documentation as evidence for each unit billed for Supported Living Services for 2 of 3 individuals.</td>
</tr>
<tr>
<td><strong>CHAPTER 6. IX. REIMBURSEMENT FOR COMMUNITY LIVING SERVICES</strong></td>
<td>Individual #2</td>
</tr>
<tr>
<td>A. Reimbursement for Supported Living Services</td>
<td>November 2009</td>
</tr>
<tr>
<td>(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.</td>
<td>• The Agency billed 6 units of Supported Living from 11/20/2009 through 11/25/2009. Documentation received accounted for 5 units.</td>
</tr>
<tr>
<td>(2) <strong>Billable Activities</strong></td>
<td>Individual #3</td>
</tr>
<tr>
<td>(a) Direct care provided to an individual in the residence any portion of the day.</td>
<td>October 2009</td>
</tr>
<tr>
<td>(b) Direct support provided to an individual by community living direct service staff away from the residence, e.g., in the community.</td>
<td>• The Agency billed 7 units of Supported Living from 10/23/2009 through 10/29/2009. Documentation received accounted for 6 units.</td>
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<tr>
<td>(c) Any activities in which direct support staff provides in accordance with the Scope of Services.</td>
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<tr>
<td>(3) <strong>Non-Billable Activities</strong></td>
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<tr>
<td>(a) The Supported Living Services provider shall not bill DD Waiver for Room and Board.</td>
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<tr>
<td>(b) Personal care, respite, nutritional counseling and nursing supports shall not be billed as separate services for an individual receiving Supported Living Services.</td>
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<td>(c) The provider shall not bill when an individual is hospitalized or in an institutional care setting.</td>
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