Dear Ms. Kenny and Ms. Schulz:

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.

   “Assuring safety and quality of care in New Mexico’s health facilities and community-based programs.”

   **David Rodriguez, Division Director • Division of Health Improvement**
   Division of Health Improvement • Quality Management Bureau • 5301 Central Ave NE • Suite 400 • Albuquerque, New Mexico 87108  
   (505) 222-8633 • FAX: (505) 222-8661


Survey Report #: Q10.02.43471889.METRO.001.INT.01
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 505-231-0088, if you have questions about the survey or the report. Thank you for your cooperation and for the work you perform.

Sincerely,

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

Entrance Conference Date: November 30, 2009

Present:

Alianza Family Services
Debbie Kenny, Managing Member
Susan Schulz, Managing Member

DOH/DHI/QMB
Marti Madrid, LBSW, Team Lead/Healthcare Surveyor
Cynthia Nielsen, MSN, Healthcare Surveyor
Barbara Czinger, MSW, LISW, Healthcare Surveyor

DDSD – Metro Regional Office
Carol Sena, Social and Community Services Coordinator

Exit Conference Date: December 3, 2009

Present:

Alianza Family Services
Debbie Kenny, Managing Member
Susan Schulz, Managing Member

DOH/DHI/QMB
Marti Madrid, LBSW, Team Lead/Healthcare Surveyor
Cynthia Nielsen, MSN, Healthcare Surveyor

DDSD - Metro Regional Office
Carol Sena, Social and Community Supports Coordinator

Homes Visited Number: 7

Administrative Locations Visited Number: 1

Total Sample Size Number: 9
0 - Jackson Class Members
9 - Non-Jackson Class Members
8 - Family Living
1 - Community Access

Persons Served Interviewed Number: 4

Records Reviewed (Persons Served) Number: 9

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 Division
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>SEVERITY</th>
<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>
<td>L.</td>
</tr>
<tr>
<td>Medium Impact</td>
<td>Actual harm</td>
<td>G.</td>
<td>H.</td>
<td>I.</td>
</tr>
<tr>
<td>Low Impact</td>
<td>No Actual Harm Potential for more than minimal harm</td>
<td>D. (2 or less)</td>
<td>E.</td>
<td>F. (3 or more)</td>
</tr>
<tr>
<td>Low Impact</td>
<td>No Actual Harm Minimal potential for harm.</td>
<td>A.</td>
<td>B.</td>
<td>C.</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 “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.
Agency: Alianza Family Services – Metro Region
Program: Developmental Disabilities Waiver
Service: Community Living (Family Living) & Community Inclusion (Community Access)
Monitoring Type: Initial Survey
Date of Survey: November 30 – December 3, 2009

<table>
<thead>
<tr>
<th>Statute</th>
<th>Deficiency</th>
<th>Agency Plan of Correction and Responsible Party</th>
<th>Date Due</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tag # 1A08  Agency Case File</td>
<td>Scope and Severity Rating: C</td>
<td>Based on record review, the Agency failed to maintain at the administrative office a confidential case file for 6 of 9 individuals. Review of the Agency individual case files revealed the following items were not found, incomplete, and/or not current:</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></td>
<td></td>
<td></td>
</tr>
<tr>
<td>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:</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(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;</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(2) The individual’s complete and current ISP, with all supplemental plans specific to the individual,</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Speech Therapy Plan (#1)</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Occupational Therapy Plan (#6)</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Physical Therapy Plan (#6)</td>
<td></td>
</tr>
</tbody>
</table>
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>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Scope and Severity Rating: E</strong></td>
<td></td>
</tr>
<tr>
<td>Medication Administration Records (MAR) were reviewed for the months of August, September and October, 2009. Based on record review, 3 of 9 individuals had Medication Administration Records, which contained missing medications entries and/or other errors: Individual #2 September 2009 Medication Administration Records did not contain the diagnosis for which the medication is prescribed: • Calcium 500 + D (One time daily) Individual #6 Medication Administration Record did not contain the specific time the medication should be given, for the following medications: • Omeprazole 20 mg/Tablet (One time daily) MAR indicated time to be given as 9:00am and 11:00am and did not indicate the exact time. Individual #8 Medication Administration Records did not contain the route of administration for the following medications: • Valproic Acid 250mg (Two times daily)</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
(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
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 # 1A09 Medication Delivery - PRN Medication</th>
<th>Scope and Severity Rating: E</th>
</tr>
</thead>
<tbody>
<tr>
<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>
<td></td>
</tr>
<tr>
<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>
<td></td>
</tr>
<tr>
<td>(2) When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) shall be maintained and include:</td>
<td></td>
</tr>
<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>
<td></td>
</tr>
<tr>
<td>(b) Prescribed dosage, frequency and method/route of administration, times and dates of administration;</td>
<td></td>
</tr>
<tr>
<td>(c) Initials of the individual administering or assisting with the medication;</td>
<td></td>
</tr>
<tr>
<td>(d) Explanation of any medication irregularity;</td>
<td></td>
</tr>
<tr>
<td>(e) Documentation of any allergic reaction or adverse medication effect; and</td>
<td></td>
</tr>
<tr>
<td>Based on record review, the Agency failed to maintain PRN Medication Administration Records which contained all elements required by standard for 2 of 9 Individuals.</td>
<td></td>
</tr>
<tr>
<td>Individual #7</td>
<td></td>
</tr>
<tr>
<td>September 2009</td>
<td></td>
</tr>
<tr>
<td>No Signs/Symptoms were noted on the Medication Administration Record for the following PRN medication:</td>
<td></td>
</tr>
<tr>
<td>• Ibuprofen 300mg - PRN - September 7, 8, 27 &amp; 28 (given 1 time daily)</td>
<td></td>
</tr>
<tr>
<td>• Enema 4.5 fluid oz - PRN - September 3, 6, 10, 13, 16, 20, 23, 26 &amp; 30 (given 1 time daily)</td>
<td></td>
</tr>
<tr>
<td>• Phisohex 148ml - PRN - September 4 (given 1 time daily)</td>
<td></td>
</tr>
<tr>
<td>• Acidophilus 100mg - PRN - September 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27 &amp; 29 (given 1 time per day)</td>
<td></td>
</tr>
<tr>
<td>• Albuterol Sulfate 0.083% - PRN - September 9, 11 &amp; 12 (given 1 time) &amp; September 10 (given 3 times)</td>
<td></td>
</tr>
<tr>
<td>• Flovent HFA 44mcg - PRN - September 9, 19, 20, 21 &amp; 22 (given 1 time) &amp; September 10, 11, 12, 13, 14, 15, 16, 17, 18 (given 2 times per day)</td>
<td></td>
</tr>
<tr>
<td>No Effectiveness was noted on the Medication Administration Record for the following PRN medication:</td>
<td></td>
</tr>
<tr>
<td>• Ibuprofen 300mg - PRN - September 7, 8, 27 &amp; 28 (given 1 time daily)</td>
<td></td>
</tr>
<tr>
<td>• Enema 4.5 fluid oz - PRN - September 3, 6, 10, 13, 16, 20, 23, 26 &amp; 30 (given 1 time daily)</td>
<td></td>
</tr>
</tbody>
</table>
(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

<table>
<thead>
<tr>
<th>PRN Medication</th>
<th>Date and Administration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phisohex 148ml - PRN</td>
<td>September 4</td>
</tr>
<tr>
<td>Acidophilus 100mg - PRN</td>
<td>September 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27 &amp; 29</td>
</tr>
<tr>
<td>Albuterol Sulfate 0.083% - PRN</td>
<td>September 9, 11 &amp; 12 (given 1 time) &amp; September 10 (given 3 times)</td>
</tr>
<tr>
<td>Flovent HFA 44mcg - PRN</td>
<td>September 9, 19, 20, 21 &amp; 22 (given 1 time) &amp; September 10, 11, 12, 13, 14, 15, 16, 17, 18 (given 2 times per day)</td>
</tr>
</tbody>
</table>

October 2009

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

- Enema 4.5 fluid oz – October 2, 6, 9, 13, 17, 20, 24, 28 & 31 (given 1 time daily).
- Acidophilus 100mg – October 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29 & 31 (given 1 time daily)

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

- Enema 4.5 fluid oz – October 2, 6, 9, 13, 17, 20, 24, 28 & 31 (given 1 time daily)
- Acidophilus 100mg – October 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27, 29 & 31 (given 1 time daily)

Individual #9 September 2009

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

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.

- Erythromycin 0.5% - PRN - September 3, 5, 8, 12, 21, 25 & 30. (given 1 time daily)

No Effectiveness was noted on the Medication Administration Record for the following PRN medication:
- Erythromycin 0.5% - PRN - September 3, 5, 8, 12, 21, 25 & 30. (given 1 time daily)

October 2009
No Symptoms were noted on the Medication Administration Record for the following PRN medication
- Erythromycin 0.5% - October 1, 4, 8, 11, 14, 18, 22, 27, 30 (given 1 time daily)

No Effectiveness was noted on the Medication Administration Record for the following PRN medication:
- Erythromycin 0.5% - October 1, 4, 8, 11, 14, 18, 22, 27, 30 (given 1 time daily)
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**  

<table>
<thead>
<tr>
<th>Scope and Severity Rating: F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Based on record review the Agency failed to have 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.</td>
</tr>
<tr>
<td>Review of Agency’s policies and procedures found no evidence of the Agency’s transportation policy &amp; procedure.</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.

**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.

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

Training Requirements for Direct Service Agency

- Transportation (DSP #40, 41, 42, 43, 44, 45, 46, 47, 48, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86 & 87)

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 43 of 43 Direct Service Personnel.

No documented evidence was found of the following required training:

- Transportation (DSP #40, 41, 42, 43, 44, 45, 46, 47, 48, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86 & 87)
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)
<table>
<thead>
<tr>
<th>Tag # 1A12 Reimbursement/Billable Units</th>
<th>Scope and Severity Rating: A</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 provide written or electronic documentation as evidence for each unit billed, which contained the required information for 1 of 9 individuals.</td>
</tr>
<tr>
<td>CHAPTER 1 III. PROVIDER AGENCY DOCUMENTATION OF SERVICE DELIVERY AND LOCATION</td>
<td></td>
</tr>
<tr>
<td>A. General: All Provider Agencies shall maintain all records necessary to fully disclose the service, quality, quantity and clinical necessity furnished to individuals who are currently receiving services. The Provider Agency records shall be sufficiently detailed to substantiate the date, time, individual name, servicing Provider Agency, level of services, and length of a session of service billed.</td>
<td>Individual #4 August 2009 • The Agency billed a total of 29 units of Family Living on August 2009. Documentation did not contain a description of what occurred during the encounter or service interval to justify billing.</td>
</tr>
<tr>
<td>B. Billable Units: The documentation of the billable time spent with an individual shall be kept on the written or electronic record that is prepared prior to a request for reimbursement from the HSD. For each unit billed, the record shall contain the following:</td>
<td>September 2009 • The agency billed a total of 30 units of Family Living in September 2009. Documentation did not contain a description of what occurred during the encounter or service interval to justify billing.</td>
</tr>
<tr>
<td>(1) Date, start and end time of each service encounter or other billable service interval;</td>
<td>October 2009 • The agency billed a total of 31 units of Family Living in October 2009. Documentation did not contain a description of what occurred during the encounter or service interval to justify billing.</td>
</tr>
<tr>
<td>(2) A description of what occurred during the encounter or service interval; and</td>
<td></td>
</tr>
<tr>
<td>(3) The signature or authenticated name of staff providing the service.</td>
<td></td>
</tr>
</tbody>
</table>

MAD-MR: 03-59 Eff 1/1/2004
8.314.1 BI RECORD KEEPING AND 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.
Tag # 1A15  Healthcare Documentation


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

Scope and Severity Rating: E

Based on record review, the Agency failed to maintain the required documentation in the Individuals Agency Record as required per standard for 5 of 9 individuals:

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

- Medication Administration Assessment Tool (#5)
- Quarterly Nursing Review of HCP/Crisis Plans:
  - None found for 9/2009 - 12/2009 (#7)
  - None found for 9/2009 - 12/2009 (#8)
- Special Health Care Needs:
  - Tube Feeding Protocol
    - Individual #7 - As indicated by the IST section of ISP the individual is required to have a tube feeding plan
  - Nutritional Plan
    - Individual #7 - As indicated by the IST section of ISP the individual is required to have a nutritional plan.
    - Individual #9 - As indicated by the IST section of ISP the individual is required to have a nutritional plan.
- Health Care Plans
  - No Healthcare Plan found.
  - Individual #5 - According to documents reviewed the individual is required to have a plan, as they have a HAT score of 5.
- Diabetes
  - Individual #2 - As indicated by the IST section of ISP the individual is required to have a diabetic plan
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.

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).

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.

<table>
<thead>
<tr>
<th>Crisis Plans</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Diabetes Crisis Plan</strong></td>
</tr>
<tr>
<td>° Individual #2 - As indicated by the IST section of ISP the individual is required to have a diabetic crisis plan.</td>
</tr>
</tbody>
</table>

| Cardiac Condition |
| ° Individual #2 - As indicated by the IST section of ISP the individual is required to have a cardiac condition plan. |

| Allergies Crisis Plan |
| ° Individual #2 - As indicated by the IST section of ISP the individual is required to have a plan. |
(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 # 1A20</th>
<th>DSP Training Documents</th>
<th>Scope and Severity Rating:  E</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 Orientation and Training requirements were met for 10 of 43 Direct Service Personnel.</td>
<td></td>
</tr>
<tr>
<td>CHAPTER 1 IV. GENERAL REQUIREMENTS FOR PROVIDER AGENCY SERVICE</td>
<td></td>
<td></td>
</tr>
<tr>
<td>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.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>C. Orientation and Training Requirements:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>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>
<td></td>
</tr>
<tr>
<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>
<td></td>
<td></td>
</tr>
<tr>
<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>
<td></td>
<td></td>
</tr>
<tr>
<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>
<td></td>
</tr>
<tr>
<td>A. Individuals shall receive services from competent and qualified staff.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

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

Review of Direct Service Personnel training records found no evidence of the following required DOH/DDSD trainings and certification being completed:

- First Aid (DSP #63, 66, 69, 71, 74 & 76)
- CPR (DSP #63, 66, 69, 71, 74 & 76)
- Assisting With Medication Delivery (DSP #57, 71 & 74)
- Rights & Advocacy (DSP #54)
- Level 1 Health (DSP #54)
- Positive Behavior Supports Strategies (DSP #54)
- PCP (DSP #69 & 78)
- Pre – Service (DSP# 69 & 81)
- Basic Health (DSP #69 & 81)
<table>
<thead>
<tr>
<th>Tag # 1A28 (CoP) Incident Mgt. System - Personnel Training</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Scope and Severity Rating: E</strong></td>
<td>Based on record review the Agency failed to provide documentation verifying completion of Incident Management Training for 25 of 47 Agency Personnel.</td>
</tr>
<tr>
<td><strong>NMAC 7.1.13.10 INCIDENT MANAGEMENT SYSTEM REQUIREMENTS:</strong></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></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></td>
</tr>
<tr>
<td><strong>Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007</strong></td>
<td></td>
</tr>
<tr>
<td><strong>II. POLICY STATEMENTS:</strong></td>
<td></td>
</tr>
<tr>
<td><strong>A.</strong> Individuals shall receive services from competent and qualified staff.</td>
<td></td>
</tr>
<tr>
<td><strong>C.</strong> Staff shall complete training on DOH-approved incident reporting procedures in accordance with 7 NMAC 1.13.</td>
<td></td>
</tr>
<tr>
<td>Tag # 1A29 Complaints / Grievances - Acknowledgement</td>
<td>Scope and Severity Rating: A</td>
</tr>
<tr>
<td>--------------------------------------------------</td>
<td>----------------------------</td>
</tr>
<tr>
<td>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].</td>
<td>Based on record review, the Agency failed to provide documentation, the complaint procedure had been made available to individuals or their legal guardians for 1 of 9 individuals.</td>
</tr>
</tbody>
</table>

- **Grievance/Complaint Procedure Acknowledgement (#1)**

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.
Tag # 1A31 (CoP) Client Rights/Human Rights

Scope and Severity Rating: D

<table>
<thead>
<tr>
<th>Rule</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>7.26.3.11 RESTRICTIONS OR LIMITATION OF CLIENT'S RIGHTS:</td>
<td>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]. 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. 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>
</tr>
</tbody>
</table>

| Based on record review, the Agency failed to ensure the rights of Individuals was not restricted or limited for 1 of 9 Individuals. (Individual #1) A review of Agency Individual files indicated 1 of 9 Individuals required Human Rights Committee Approval for restrictions. No documentation was found regarding Human Rights Approval for the following: • Psychotropic Medications to control behaviors. No evidence found of Human Rights Committee approval. (Individual #1) |

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 response to psychiatric and/or behavioral symptoms in addition to the above requirements, obtain current written consent from the individual, guardian or surrogate health decision maker and submit for review by the agency’s Human Rights Committee (References: Psychotropic Medication Use Policy, Section D, page 5 Use of PRN Psychotropic
Medications; and Human Rights Committee Requirements Policy, Section B, page 4 - Interventions Requiring Review and Approval – Use of PRN Medications).
<table>
<thead>
<tr>
<th>Tag # 1A32 (CoP)</th>
<th>ISP Implementation</th>
</tr>
</thead>
<tbody>
<tr>
<td>NMAC 7.26.5.16.C and D Development of the ISP. Implementation of the ISP. The ISP shall be implemented according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan.</td>
<td></td>
</tr>
<tr>
<td>C. The IDT shall review and discuss information and recommendations with the individual, with the goal of supporting the individual in attaining desired outcomes. The IDT develops an ISP based upon the individual's personal vision statement, strengths, needs, interests and preferences. The ISP is a dynamic document, revised periodically, as needed, and amended to reflect progress towards personal goals and achievements consistent with the individual's future vision. This regulation is consistent with standards established for individual plan development as set forth by the commission on the accreditation of rehabilitation facilities (CARF) and/or other program accreditation approved and adopted by the developmental disabilities division and the department of health. It is the policy of the developmental disabilities division (DDD), that to the extent permitted by funding, each individual receive supports and services that will assist and encourage independence and productivity in the community and attempt to prevent regression or loss of current capabilities. Services and supports include specialized and/or generic services, training, education and/or treatment as determined by the IDT and documented in the ISP.</td>
<td></td>
</tr>
<tr>
<td>D. The intent is to provide choice and obtain opportunities for individuals to live, work and play with full participation in their communities. The following principles provide direction and purpose in planning for individuals with developmental disabilities. [05/03/94; 01/15/97; Recompiled 10/31/01]</td>
<td></td>
</tr>
<tr>
<td>Scope and Severity Rating: E</td>
<td></td>
</tr>
<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 4 of 8 individuals.</td>
<td></td>
</tr>
<tr>
<td>Per Individuals ISP the following was found with regards to the implementation of ISP Outcomes:</td>
<td></td>
</tr>
<tr>
<td>Family Living Data Collection/Data Tracking/Progress with regards to ISP Outcomes:</td>
<td></td>
</tr>
<tr>
<td>Individual #1</td>
<td></td>
</tr>
<tr>
<td>• None found for September 2009 - October 2009.</td>
<td></td>
</tr>
<tr>
<td>Individual #2</td>
<td></td>
</tr>
<tr>
<td>• None found for September 2009.</td>
<td></td>
</tr>
<tr>
<td>Individual #8</td>
<td></td>
</tr>
<tr>
<td>• None found for September 2009.</td>
<td></td>
</tr>
<tr>
<td>Individual #9</td>
<td></td>
</tr>
<tr>
<td>• None found for September 2009 - October 2009.</td>
<td></td>
</tr>
<tr>
<td>Tag # 1A37 Individual Specific Training</td>
<td>Scope and Severity Rating: F</td>
</tr>
<tr>
<td>---------------------------------------</td>
<td>-----------------------------</td>
</tr>
<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 38 of 47 Agency Personnel.</td>
</tr>
<tr>
<td><strong>CHAPTER 1 IV. GENERAL REQUIREMENTS FOR PROVIDER AGENCY SERVICE</strong></td>
<td><strong>Review of personnel records found no evidence of the following:</strong></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 (#40, 41, 42, 43, 44, 46, 48, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 70, 71, 72, 73, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86 &amp; 87)</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: (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>
<td></td>
</tr>
<tr>
<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>
</tr>
<tr>
<td>A. Individuals shall receive services from competent and qualified staff.</td>
<td></td>
</tr>
<tr>
<td>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.</td>
<td></td>
</tr>
</tbody>
</table>
## Tag # 6L14 Residential Case File

### Scope and Severity Rating: E

Based on record review, the Agency failed to maintain a complete and confidential case file in the residence for 5 of 8 individuals receiving Family Living Services.

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

- **Annual ISP (#6)**
- **Speech Therapy Plan (#1, 6 & 8)**
- **Occupational Therapy Plan (#6)**
- **Physical Therapy Plan (#6 & 8)**
- **Special Health Care Needs**
  - Nutritional Plan (#8)
- **Health Care Plans**
  - G Tube (#7)
- **Crisis Plan**
  - Seizures (#4)
  - Sleep Apnea (#7)

### Service Provider Agency Requirements

#### Chapter 4: Community Living Service Provider Agency Requirements

##### A. Residence Case File:

- For individuals receiving Supported Living or Family Living, the Agency shall maintain in the individual's home a complete and current confidential case file for each individual. For individuals receiving Independent Living Services, rather than maintaining this file at the individual’s home, the complete and current confidential case file for each individual shall be maintained at the agency’s administrative site. Each file shall include the following:
  1. Complete and current ISP and all supplemental plans specific to the individual;
  2. Complete and current Health Assessment Tool;
  3. Current emergency contact information, which includes the individual’s address, telephone number, names and telephone numbers of residential Community Living Support providers, relatives, or guardian or conservator, primary care physician’s name(s) and telephone number(s), pharmacy name, address and telephone number and dentist name, address and telephone number, and health plan;
  4. Up-to-date progress notes, signed and dated by the person making the note for at least the past month (older notes may be transferred to the agency office);
  5. Data collected to document ISP Action Plan implementation
  6. Progress notes written by direct care staff and by nurses regarding individual health status and physical conditions including action taken in response to identified changes in condition for at least the past month;
  7. Physician’s or qualified health care providers written orders;
  8. Progress notes documenting implementation of...
a physician’s or qualified health care provider’s order(s);  
(9) Medication Administration Record (MAR) for the past three (3) months which includes:  
(a) The name of the individual;  
(b) A transcription of the healthcare 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
<p>| discharge summaries for past twelve (12) months, past medical history including hospitalizations, surgeries, injuries, family history and current physical exam. |   |   |</p>
<table>
<thead>
<tr>
<th>Tag # 6L25 (CoP)</th>
<th>Residential Health &amp; Safety (Family Living)</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></td>
<td>Based on observation, the Agency failed to ensure that each individual’s residence met all requirements within the standard for 7 of 7 Family Living residences.</td>
</tr>
<tr>
<td>CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS</td>
<td></td>
<td>The following items were not found, not functioning or incomplete:</td>
</tr>
<tr>
<td>L. Residence Requirements for Family Living Services and Supported Living Services</td>
<td>(1) Supported Living Services and Family Living Services providers shall assure that each individual’s residence has:</td>
<td>• Accessible written procedures for the safe storage of all medications with dispensing instructions for each individual that are consistent with the Assisting with Medication Administration training or each individual’s ISP (#1, 2, 4, 5, 7, 8 &amp; 9)</td>
</tr>
<tr>
<td>(a) Battery operated or electric smoke detectors, heat sensors, or a sprinkler system installed in the residence;</td>
<td>(b) General-purpose first aid kit;</td>
<td></td>
</tr>
<tr>
<td>(c) When applicable due to an individual’s health status, a blood borne pathogens kit;</td>
<td>(d) Accessible written procedures for emergency evacuation e.g. fire and weather-related threats;</td>
<td></td>
</tr>
<tr>
<td>(e) Accessible telephone numbers of poison control centers located within the line of sight of the telephone;</td>
<td>(f) Accessible written documentation of actual evacuation drills occurring at least three (3) times a year. For Supported Living evacuation drills shall occur at least once a year during each shift;</td>
<td></td>
</tr>
<tr>
<td>(g) Accessible written procedures for the safe storage of all medications with dispensing instructions for each individual that are consistent with the Assisting with Medication Administration training or each individual’s ISP; and</td>
<td>(h) Accessible written procedures for emergency placement and relocation of individuals in the event of an emergency evacuation that makes the residence unsuitable for occupancy. The emergency evacuation procedures shall address, but are not limited to, fire, chemical and/or hazardous waste spills, and flooding.</td>
<td></td>
</tr>
</tbody>
</table>