Date:	January 8, 2010
To:	Matt Poel, Executive Director
Provider:	Great Livin' LLC
Address:	609 Broadway Blvd, NE, Suite 217
State/Zip:	Albuquerque, New Mexico 87102
E-mail Address:	Matt@Great-Livin.com
Region:	Metro
Survey Date:	October 13 - 15, 2009
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	Community Living (Supported Living)
Survey Type:	Routine
Team Leader:	Stephanie R. Martinez de Berenger, M.P.A., GCDF, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau
Team Members:	Nadine Romero, LBSW, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau;

Dear Mr. Poel,

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/Quality Management Bureau is issuing your agency a "SUB-STANDARD" rating for significant non-compliance with DDSD Standards and regulations; additionally your agency is being referred to the Internal Review Committee for consideration of remedies and possible sanctions.

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 900 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 505-690-7280, if you have questions about the survey or the report. Thank you for your cooperation and for the work you perform.

Sincerely,

Stephanie R. Martinez de Berenger, MPA, GCDF

Stephanie R. Martinez de Berenger, M.P.A., GCDF
Team Lead/Healthcare Surveyor
Division of Health Improvement
Quality Management Bureau

Survey Process Employed:

Entrance Conference Date:	October 13, 2009
Present:	**Great Livin', LLC** Matt Poel, Executive Director Steven Nadolny, Administrative Director **DOH/DHI/QMB** Stephanie R. Martinez de Berenger, MPA, GCDG, Team Lead/Healthcare Surveyor Nadine Romero, LBSW, Healthcare Surveyor
Exit Conference Date:	October 15, 2009
Present:	**Great Livin', LLC** Matt Poel, Executive Director Steven Nadolny, Administrative Director Jackie Martinez, Service Coordinator **DOH/DHI/QMB** Stephanie R. Martinez de Berenger, MPA, GCDG, Team Lead/Healthcare Surveyor Nadine Romero, LBSW, Healthcare Surveyor
Homes Visited	Number: 4
Administrative Locations Visited	Number: 1
Total Sample Size (duplicated)	Number: 4 0 - Jackson Class Members 4 - Non-Jackson Class Members 4 - Supported Living
Persons Served Interviewed	Number: 3
Persons Served Observed	Number: 1
Records Reviewed (Persons Served)	Number: 4
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

Attachment A

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.

Attachment B

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.

			SCOPE		
			Isolated 01% - 15%	**Pattern 16% - 79%**	**Widespread 80% - 100%**
SEVERITY	**High Impact**	Immediate Jeopardy to individual health and or safety	**J.**	**K.**	**L.**
		Actual harm	**G.**	**H.**	**I.**
	Medium Impact	No Actual Harm Potential for more than minimal harm	**D.**	**E.**	**F.** (3 or more)
			D. (2 or less)		**F.** (no conditions of participation)
	Low Impact	No Actual Harm Minimal potential for harm.	**A.**	**B.**	**C.**

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 that 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.

Agency: **Great Livin', LLC – Metro Region**
Program: Developmental Disabilities Waiver
Service: Community Living (Supported Living)
Monitoring Type: Routine Survey
Date of Survey: **October 13 – 15, 2009**

Statute	Deficiency	Agency Plan of Correction and Responsible Party	Date Due
Tag # 1A08 Agency Case File	**Scope and Severity Rating: A**		
Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 **CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS:** The objective of these standards is to establish Provider Agency policy, procedure and reporting requirements for DD Medicaid Waiver program… **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…	Based on record review, the Agency failed to maintain at the administrative office a confidential case file for 1 of 4 individuals. Review of the Agency individual case files revealed the following items were not found, incomplete, and/or not current: • Speech Therapy Plan (#3)		

Tag # 1A08 Agency Case File - Progress Notes	Scope & Severity Rating: A		
Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 **CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS:** The objective of these standards is to establish Provider Agency policy, procedure and reporting requirements for DD Medicaid Waiver program. These requirements apply to all such Provider Agency staff, whether directly employed or subcontracting with the Provider Agency. Additional Provider Agency requirements and personnel qualifications may be applicable for specific service standards. **D. Provider Agency Case File for the Individual:** All Provider Agencies shall maintain at the administrative office a confidential case file for each individual. Case records belong to the individual receiving services and copies shall be provided to the receiving agency whenever an individual changes providers. The record must also be made available for review when requested by DOH, HSD or federal government representatives for oversight purposes. The individual's case file shall include the following requirements: (3) Progress notes and other service delivery documentation;	Based on record review, the Agency failed to maintain progress notes and other service delivery documentation for 1 of 4 Individuals. **Supported Living Progress Notes/Daily Contact Logs** • Individual #3 - None found for August 15 ,17 & 24, 2009		

Tag # 1A09 Medication Delivery (MAR) - Routine Medication	Scope and Severity Rating: E		
Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 **CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS:** The objective of these standards is to establish Provider Agency policy, procedure and reporting requirements for DD Medicaid Waiver program. These requirements apply to all such Provider Agency staff, whether directly employed or subcontracting with the Provider Agency. Additional Provider Agency requirements and personnel qualifications may be applicable for specific service standards. **E. Medication Delivery:** Provider Agencies that provide Community Living, Community Inclusion or Private Duty Nursing services shall have written policies and procedures regarding medication(s) delivery and tracking and reporting of medication errors in accordance with DDSD Medication Assessment and Delivery Policy and Procedures, the Board of Nursing Rules and Board of Pharmacy standards and regulations. (2) When required by the DDSD Medication Assessment and Delivery Policy, Medication Administration Records (MAR) shall be maintained and include: (a) The name of the individual, a transcription of the physician's written or licensed health care provider's prescription including the brand and generic name of the medication, diagnosis for which the medication is prescribed; (b) Prescribed dosage, frequency and method/route of administration, times and dates of administration; (c) Initials of the individual administering or assisting with the medication; (d) Explanation of any medication irregularity; (e) Documentation of any allergic reaction or adverse medication effect; and	Medication Administration Records (MAR) were reviewed for the months of June, July, August 2009. Based on record review, 4 of 4 individuals had Medication Administration Records, which contained missing medications entries and/or other errors: Individual #1 June 2009 Medication Administration Records did not contain the diagnosis for which the medication is prescribed: • Trazodone 150MG (1 time daily) July 2009 Medication Administration Records did not contain the diagnosis for which the medication is prescribed: ° Trazodone 50MG (2 times daily) ° Seroquel 400MG (1 time daily) ° Depakote ER 250MG (7 times daily) ° Folic Acid 1MG (1 time daily) August 2009 Medication Administration Records did not contain the diagnosis for which the medication is prescribed: Trazodone 50MG (2 times daily) Seroquel 400MG (1 time daily) Depakote ER 250MG (7 times daily) Folic Acid 1MG (1 time daily) Individual #2 June 2009		

(f) For PRN medication, an explanation for the use of the PRN medication shall include observable signs/symptoms or circumstances in which the medication is to be used, and documentation of effectiveness of PRN medication administered. (3) The Provider Agency shall also maintain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose; (4) MARs are not required for individuals participating in Independent Living who self-administer their own medications; (5) Information from the prescribing pharmacy regarding medications shall be kept in the home and community inclusion service locations and shall include the expected desired outcomes of administrating the medication, signs and symptoms of adverse events and interactions with other medications; **NMAC 16.19.11.8 MINIMUM STANDARDS:** A. MINIMUM STANDARDS FOR THE DISTRIBUTION, STORAGE, HANDLING AND RECORD KEEPING OF DRUGS: (d) The facility shall have a Medication Administration Record (MAR) documenting medication administered to residents, **including over-the-counter medications.** This documentation shall include: (i) Name of resident; (ii) Date given; (iii) Drug product name; (iv) Dosage and form; (v) Strength of drug; (vi) Route of administration; (vii) How often medication is to be taken; (viii) Time taken and staff initials; (ix) Dates when the medication is discontinued or changed; (x) The name and initials of all staff	Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Risperdal 2MG (1 time daily) – Blank June 8, 2009 (8:00 PM) • Phenytoin SO EXT 100MG (4 times daily) – Blank June 8, 2009 (8:00PM) • Lithium Carbonate 300 MG (2 times daily) – Blank June 8, 2009 (8:00 AM & 8:00 PM) • Calcium Citrate + D (2 times daily) Blank – June 8, 2009 (8:00 PM) • Lamictal 150 MG (3 times daily) Blank – June 8, 2009 (8:00 AM, 12:00PM & 6:00PM) Medication Administration Records did not contain the diagnosis for which the medication is prescribed: • Multivitamin (1 time daily) • Calcium Citrate + D (2 times daily) • Cephalexin 500MG (4 times daily) Medication Administration Record document did not contain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose for the following medications: • Risperdal 2 MG (1 time daily) • Phenytoin SO EXT 100MG • Multivitamin (1 time daily) • Lithium Carbonate 300 • Calcium Citrate + D (2 times daily)		

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.	• Lamictal 150 MG (3 times daily) July 2009 During on-site survey Medication Administration Records were requested for months of June, July & August 2009. As of October 15, 2009, Medication Administration Records for the month of July 2009 had not been provided. August 2009 Medication Administration Records did not contain the diagnosis for which the medication is prescribed: • Multivitamin (1 time daily) • Calcium Citrate + D (2 times daily) • Phenytoin SOD EXT 100MG (2 times daily) • Metamucil Packet (1 time daily) Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Metamucil Packet (1 time daily) – Blank August 25, 2009 (12:00 PM) Individual #3 During on-site survey Medication Administration Records were requested for months of August and September, 2009. Individual began services on August 15, 2009. As of October 15, 2009, Medication Administration Records had not been provided.		

Tag # 1A09 Medication Delivery - PRN Medication	Scope and Severity Rating: E		
Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 **CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS:** The objective of these standards is to establish Provider Agency policy, procedure and reporting requirements for DD Medicaid Waiver program… **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… (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;	Based on record review, the Agency failed to maintain PRN Medication Administration Records which contained all elements required by standard for 1 of 4 individuals. Individual #2 June 2009 No Signs/Symptoms was noted on the Medication Administration Record for the following PRN medication: • Hydrocodone – APAP5 – 500MG – PRN – June 10, 12, 14 & 16, 2009 (given 1 time daily) and June 11, 2009 (given 2 times daily). • Antacid Tablets Calcium Carbonate 500MG - PRN - June 1, 2009 (given 2 times) & June 2, 2009 (given 3 times). No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Hydrocodone – APAP5 – 500MG – PRN – June 10, 12, 14 & 16, 2009 (given 1 time daily) and June 11, 2009 (given 2 times daily). • Antacid Tablets Calcium Carbonate 500MG - PRN - June 1, 2009 (given 2 times) & June 2, 2009 (given 3 times). Medication Administration Records did not contain the circumstance for which the medication is to be used: • Hydrocodone – APAP5 – 500MG (PRN) Medication Administration Records did not contain the route of administration for the following medications: • Hydrocodone – APAP5 – 500MG (PRN) Medication Administration Record document did		

(4) MARs are not required for individuals participating in Independent Living who self-administer their own medications; (5) Information from the prescribing pharmacy regarding medications shall be kept in the home and community inclusion service locations and shall include the expected desired outcomes of administrating the medication, signs and symptoms of adverse events and interactions with other medications; **NMAC 16.19.11.8 MINIMUM STANDARDS:** **A. MINIMUM STANDARDS FOR THE DISTRIBUTION, STORAGE, HANDLING AND RECORD KEEPING OF DRUGS:** (d) The facility shall have a Medication Administration Record (MAR) documenting medication administered to residents, **including over-the-counter medications.** This documentation shall include: (i) Name of resident; (ii) Date given; (iii) Drug product name; (iv) Dosage and form; (v) Strength of drug; (vi) Route of administration; (vii) How often medication is to be taken; (viii) Time taken and staff initials; (ix) Dates when the medication is discontinued or changed; (x) The name and initials of all staff administering medications. **Model Custodial Procedure Manual - *D. Administration of Drugs*** Unless otherwise stated by practitioner, patients will not be allowed to administer their own medications. Document the practitioner's order authorizing the self-administration of medications. All PRN (As needed) medications shall have complete detail instructions regarding the administering of the medication. This shall include:	not contain a signature page that designates the full name that corresponds to each initial used to document administered or assisted delivery of each dose for the following medications: • Hydrocodone – APAP5 – 500MG (PRN) • Zolpidem Tartrate 5MG (PRN) • Sudafed 60MG (PRN) • Calcium Carbonate 500MG (PRN) Medication Administration Records did not contain the exact amount to be used in a 24 hour period: • Antacid Tablets 500MG (PRN) August 2009 No Signs/Symptoms was noted on the Medication Administration Record for the following PRN medication: • Docusate Sodium 100MG – PRN – August 5, 2009 (given 1 time daily) and August 1, 2009 (given 2 times daily). • Ibuprofen 600MG - PRN - August 19, 2009 (given 1 time). No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Tylenol 500MG – PRN – August 17, 20, 22, 27and 31, 2009(given 1 time daily) and August 30, 2009 (given 2 times daily) • Docusate Sodium 100MG – PRN – August 5, 2009 (given 1 time daily) and August 1, 2009 (given 2 times daily). • Ibuprofen 600MG - PRN - August 19, 2009 (given 1 time).		

➢ 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 Training	Scope and Severity Rating: F		
Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 - **CHAPTER 1 II. PROVIDER AGENCY REQUIREMENTS:** The objective of these standards is to establish Provider Agency policy, procedure and reporting requirements for DD Medicaid Waiver program… **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… **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:** I. 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)	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 15 of 18 Direct Service Personnel **No documented evidence was found of the following required training:** • Transportation (DSP #40, 41, 42, 43, 44, 47, 48, 49, 50, 51, 53, 54, 56 & 57) **When DSP were asked if they had received transportation training including training on wheelchair tie downs and van lift safety the following was reported:** • DSP #43 stated, "No." • DSP #46 stated, "Not at this agency." • DSP #50 stated, "Took with [other agency name]." • DSP #56 stated, "No."		

Tag # 1A20 DSP Training Documents	Scope and Severity Rating: E		
Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 **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 (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. **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…	Based on record review, the Agency failed to ensure that Orientation and Training requirements were met for 7 of 18 Direct Service Personnel. Review of Direct Service Personnel training records found no evidence of the following required DOH/DDSD trainings and certification being completed: • Pre-Service (DSP #43) • Person-Centered Planning (1-Day) (DSP #46) • First Aid (DSP #47, 51 & 54) • CPR (DSP #47, 51, 53 & 54) • Assisting with Medication Delivery (DSP #42) • Participatory Communication & Choice Making (DSP #53) • Level 1 Health (DSP #53) • Teaching & Support Strategies (DSP #53)		

Tag # 1A25 (CoP) CCHS	Scope and Severity Rating: D		
NMAC 7.1.9.8 CAREGIVER AND HOSPITAL CAREGIVER EMPLOYMENT REQUIREMENTS: **F. Timely Submission:** 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. **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 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.	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 1 of 19 Agency Personnel. The following Agency Personnel Files contained Caregiver Criminal History Screenings, which were not specific to the Agency: • #40 – Date of hire 03/13/2009		

Tag # 1A26 (CoP) COR / EAR	Scope and Severity Rating: E		
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.	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 5 of 19 Agency Personnel. The following Agency Personnel records contained evidence that indicated the Employee Abuse Registry was completed after hire: • #41 – Date of hire 11/21/2008 • #42 – Date of hire 01/05/2009 • #47 – Date of hire 01/03/2009 • #48 – Date of hire 08/14/2009 • #58 – Date of hire 07/02/2009		

Tag # 1A28 (CoP) Incident Mgt. System - Personnel Training	Scope & Severity Rating: E		
NMAC 7.1.13.10 INCIDENT MANAGEMENT SYSTEM REQUIREMENTS: **A. General:** 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. **D. Training Documentation:** 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. **Policy Title: Training Requirements for Direct Service Agency Staff Policy - Eff. March 1, 2007** **II. POLICY STATEMENTS:** A. Individuals shall receive services from competent and qualified staff. C. Staff shall complete training on DOH-approved incident reporting procedures in accordance with 7 NMAC 1.13.	Based on record review, the Agency failed to provide documentation verifying completion of Incident Management Training for 10 of 19 Agency Personnel. • Incident Management Training (Abuse, Neglect & Misappropriation of Consumers' Property) (#45, 46, 47, 48, 49, 55, 56, 57 & 58) When DSP were asked what two State Agencies must be contacted when there is suspected Abuse, Neglect & Misappropriation of Consumers' Property, the following was reported: • DSP #43 stated, “Haven’t taken training, do not know.”		

Tag # 1A28 (CoP) Incident Mgt. System - Parent/Guardian Training	Scope & Severity Rating: E		
NMAC 7.1.13.10 INCIDENT MANAGEMENT SYSTEM REQUIREMENTS: **A. General:** 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. **E. Consumer and Guardian Orientation Packet:** 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.	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 2 of 4 individuals. • Parent/Guardian Incident Management Training (Abuse, Neglect & Misappropriation of Consumers' Property) (#3 & 4)		

Tag # 1A29 Complaints / Grievances - Acknowledgement	Scope and Severity Rating: B		
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]. **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	Based on record review, the Agency failed to provide documentation, the complaint procedure had been made available to individuals or their legal guardians for 2 of 4 individuals. • Grievance/Complaint Procedure Acknowledgement (#3 & 4)		

Tag # 1A31 (CoP) Client Rights/Human Rights	Scope and Severity Rating: F		
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	Based on record review and interview, the Agency failed to follow DDSD Policy regarding Human Rights Committee Requirements. Review of the Agency Policy & Procedure found no evidence of Human Rights Committee procedures which addressed the frequency and purpose of meetings. When surveyors asked #59 for the Human Rights Committee Policy, surveyors received the Agency's "Consumer Rights." **When Surveyors asked, if the Agency had a policy regarding the Human Rights Committee, the following was reported:** #59 and #60 reported that they are part of a local Human Rights Committee. However, the agency did not have a Human Rights Committee Policy.		

completion of each individual's Individual Service Plan. **7.26.3.11 RESTRICTIONS OR LIMITATION OF CLIENT'S RIGHTS:** 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]. 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]			

Tag # 1A33 Board of Pharmacy - Lic	Scope and Severity Rating: B		
New Mexico Board of Pharmacy Model Custodial Drug Procedures Manual **6. Display of License and Inspection Reports** A. The following are required to be publicly displayed: ☐ Current Custodial Drug Permit from the NM Board of Pharmacy ☐ Current registration from the consultant pharmacist ☐ Current NM Board of Pharmacy Inspection Report	Based on observation, the Agency failed to provide the current Custodial Drug Permit from the New Mexico Board of Pharmacy, the current registration from the Consultant Pharmacist, or the current New Mexico Board of Pharmacy Inspection Report for 1 of 2 residences: Individual Residence: • Current Custodial Drug Permit from the NM Board of Pharmacy (#1 & 3) • Current Registration of Consulting Pharmacist (#1 & 3)		

Tag # 1A37 Individual Specific Training	Scope and Severity Rating: F		
Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 **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: (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. **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.	Based on record review, the Agency failed to ensure that Individual Specific Training requirements were met for 18 of 19 Agency Personnel. Review of personnel records found no evidence of the following: • Individual Specific Training (#40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54 , 56, 57 & 58)		

Tag # 6L14 Residential Case File	Scope and Severity Rating: F		
Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 **CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS** **A. Residence Case File:** For individuals receiving Supported Living or Family Living, the Agency shall maintain in the individual's home a complete and current confidential case file for each individual. For individuals receiving Independent Living Services, rather than maintaining this file at the individual's home, the complete and current confidential case file for each individual shall be maintained at the agency's administrative site. Each file shall include the following: (1) Complete and current ISP and all supplemental plans specific to the individual; (2) Complete and current Health Assessment Tool; (3) Current emergency contact information, which includes the individual's address, telephone number, names and telephone numbers of residential Community Living Support providers, relatives, or guardian or conservator, primary care physician's name(s) and telephone number(s), pharmacy name, address and telephone number and dentist name, address and telephone number, and health plan; (4) Up-to-date progress notes, signed and dated by the person making the note for at least the past month (older notes may be transferred to the agency office); (5) Data collected to document ISP Action Plan implementation (6) Progress notes written by direct care staff and by nurses regarding individual health status and physical conditions including action taken in response to identified changes in condition for at least the past month; (7) Physician's or qualified health care providers	Based on record review, the Agency failed to maintain a complete and confidential case file in the residence for 4 of 4 Individuals receiving Supported Living Services. The following was not found, incomplete and/or not current: • Annual ISP (#4) • Teaching & Support Strategies (#1 & 3) • ISP Signature Page (#1, 2, 3 & 4) • Addendum A (#1, 2, 3 & 4) • Individual Specific Training (Addendum B) (#3 & 4) • Speech Therapy Plan (#3) • Health Assessment Tool (#1, 2 & 4) • **Health Care Plans** ° Aspiration (#1) ° Seizures (#2) • **Crisis Plans** ° Aspiration (#1) ° Sketal Integrity (#1) ° Seizures (#2) ° Asthma (#4) • **Progress Notes/Daily Contacts Logs:** ° Individual #3 - September and October 2009 • **Data Collection/Data Tracking:** ° Individual #1 - None found for October 2009		

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… (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…	° Individual #2 - None found for October 2009 • **Progress Notes written by DSP and/or Nurses regarding Health Status:** ° Individual #2 - None found for September and October 2009 • Health Care Providers Written Orders (#2) • Record of visits of healthcare practitioners (#2)		

Tag # 6L25 (CoP) Residential Health & Safety (Supported Living)	Scope and Severity Rating: E		
Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 **CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS** **L. Residence Requirements for Family Living Services and Supported Living Services** (1) Supported Living Services and Family Living Services providers shall assure that each individual's residence has: (a) Battery operated or electric smoke detectors, heat sensors, or a sprinkler system installed in the residence; (b) General-purpose first aid kit; (c) When applicable due to an individual's health status, a blood borne pathogens kit; (d) Accessible written procedures for emergency evacuation e.g. fire and weather-related threats; (e) Accessible telephone numbers of poison control centers located within the line of sight of the telephone; (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; (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 (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.	Based on observation, the Agency failed to ensure that each individual's residence met all requirements within the standard for 1 of 2 Supported Living residences. The following items were not found, not functioning or incomplete: • Battery operated or electric smoke detectors, heat sensors, or a sprinkler system installed in the residence (#1 & 3) • Accessible written procedures for emergency evacuation e.g. fire and weather-related threats (#1 & 3) • 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. (#1 & 3)		

Tag # 6L26 SL Reimbursement	Scope and Severity Rating: A		
Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007 **CHAPTER 6. IX. REIMBURSEMENT FOR COMMUNITY LIVING SERVICES** A. **Reimbursement** for Supported Living Services (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. (2) **Billable Activities** (a) Direct care provided to an individual in the residence any portion of the day. (b) Direct support provided to an individual by community living direct service staff away from the residence, e.g., in the community. (c) Any activities in which direct support staff provides in accordance with the Scope of Services. (3) Non-Billable Activities (a) The Supported Living Services provider shall not bill DD Waiver for Room and Board. (b) Personal care, respite, nutritional counseling and nursing supports shall not be billed as separate services for an individual receiving Supported Living Services. (c) The provider shall not bill when an individual is hospitalized or in an institutional care setting.	Based on record review, the Agency failed to provide written or electronic documentation as evidence for each unit billed for Supported Living Services for 1 of 4 individuals. Individual #3 August 2009 • The Agency billed 3 units of Supported Living on August 15, 17 & 24, 2009. No documentation found to justify billing.		