Dear Mr. Turner,

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 agency a determination of “Non Compliance with Conditions of Participation.”

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

1. Quality Management Bureau, Attention: Plan of Correction Coordinator
   5301 Central Ave. NE Suite 400 Albuquerque, NM 87108

2. Developmental Disabilities Supports Division Regional Office for region of service surveyed.

Upon notification from QMB that your Plan of Correction has been approved, you must implement all remedies and corrective actions within 45 working days. If your plan of correction is denied, you must resubmit a revised plan ASAP for approval. All remedies must still be completed within 45 working days of the original submission.

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

David Rodriguez, Division Director • Division of Health Improvement
Quality Management Bureau • 5301 Central Ave. NE Suite 400 • Albuquerque, New Mexico 87108
(505) 222-8623 • FAX: (505) 222-8661 • http://dhi.health.state.nm.us

DHITe World 2010

Survey Report #: Q11.01.61285854.METRO.001.RTN.01
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.

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*

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

<table>
<thead>
<tr>
<th>Entrance Conference Date:</th>
<th>July 27, 2010</th>
</tr>
</thead>
</table>
| Present:                  | Creative Careers  
David Turner, Director  
DOH/DHI/QMB  
Marti Madrid, LBSW, Team Lead/Healthcare Surveyor  
Barbara Czinger, LISW, Healthcare Surveyor  
DDSD - Metro Regional Office  
Linda Clark, Supported Employment Coordinator  

Exit Conference Date: | July 29, 2010 |
|-----------------------|--------------|
| Present:              | Creative Careers  
David Turner, Director  
DOH/DHI/QMB  
Marti Madrid, LBSW, Team Lead/Healthcare Surveyor  
DDSD - Metro Regional Office  
Linda Clark, Supported Employment Coordinator  

<table>
<thead>
<tr>
<th>Administrative Locations Visited</th>
<th>Number: 1</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total Sample Size Number: 8</td>
<td></td>
</tr>
</tbody>
</table>
0 - Jackson Class Members  
8 - Non-Jackson Class Members  
8 - Supported Employment |
| Persons Served Interviewed Number: 2 |
| Persons Served Observed Number: 6 | Individuals were not available during the on-site survey. |
| Records Reviewed (Persons Served) Number: 8 |

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

Scope and Severity Definitions:

Key to Scope scale:

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

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

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

Key to Findings:

“Substantial Compliance with Conditions of Participation”
The QMB determination of “Substantial Compliance with Conditions of Participation” indicates that a provider is in substantial compliance with all ‘Conditions of Participation’ and other standards and regulations. The agency has obtained a level of compliance such that there is a minimal potential for harm to individuals' health and safety. To qualify for a determination of Substantial Compliance with Conditions of Participation, the provider must not have any findings that meet the thresholds for determining non-compliance with any Condition of Participation.

“Non-Compliance with Conditions of Participation”
The QMB determination of “Non-Compliance with Conditions of Participation” indicates that a provider is out of compliance with one (1) or more ‘Conditions of Participation,’ This non-compliance, if not corrected, is likely to result in a serious negative outcome or the potential for more than minimal harm to individuals’ health and safety.

Providers receiving a repeat determination of Non-Compliance may be referred by QMB to the Internal Review Committee (IRC) for consideration of remedies and possible actions.

“Sub-Standard Compliance with Conditions of Participation”:
The QMB determination of “Sub-Standard Compliance with Conditions of Participation” indicates a provider is significantly out of compliance with Conditions of Participation and/or has:
Multiple findings of widespread non-compliance with any standard or regulation with a significant potential for more than minimal harm. Any finding of actual harm or Immediate Jeopardy.

Providers receiving a repeat determination of ‘Substandard Compliance’ will be referred by QMB to the Internal Review Committee (IRC) for consideration of remedies and possible actions.
Introduction:
The process involves open communication between surveyors and 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 formally 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.
<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: B</td>
<td>Based on record review, the Agency failed to maintain at the administrative office a confidential case file for 2 of 8 individuals.</td>
<td></td>
</tr>
<tr>
<td>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.</td>
<td>Review of the Agency individual case files found the following items were not found, incomplete, and/or not current:</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: (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,</td>
<td>• Positive Behavioral Plan (#2)</td>
<td></td>
<td></td>
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<tr>
<td></td>
<td></td>
<td>• Positive Behavioral Crisis Plan (#1 &amp; 2)</td>
<td></td>
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<tr>
<td></td>
<td></td>
<td>• Speech Therapy Plan (#2)</td>
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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.
Tag # 1A20   DSP Training Documents

<table>
<thead>
<tr>
<th>Scope and Severity Rating:</th>
<th>E</th>
</tr>
</thead>
<tbody>
<tr>
<td>Based on record review, the Agency failed to ensure that Orientation and Training requirements were met for 3 of 6 Direct Service Personnel.</td>
<td></td>
</tr>
<tr>
<td>Review of Direct Service Personnel training records found no evidence of the following required DOH/DDSD trainings and certification being completed:</td>
<td></td>
</tr>
<tr>
<td>- Basic Health/Orientation (DSP #43)</td>
<td></td>
</tr>
<tr>
<td>- Person-Centered Planning (1-Day) (DSP #43)</td>
<td></td>
</tr>
<tr>
<td>- First Aid (DSP #42, 43 &amp; 44)</td>
<td></td>
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<tr>
<td>- CPR (DSP #42 &amp; 43)</td>
<td></td>
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</tbody>
</table>


CHAPTER 1 IV. GENERAL REQUIREMENTS FOR PROVIDER AGENCY SERVICE

PERSONNEL: The objective of this section is to establish personnel standards for DD Medicaid Waiver Provider Agencies for the following services: Community Living Supports, Community Inclusion Services, Respite, Substitute Care and Personal Support Companion Services. These standards apply to all personnel who provide services, whether directly employed or subcontracting with the Provider Agency. Additional personnel requirements and qualifications may be applicable for specific service standards.

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.
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.

C. Staff shall complete training on DOH-approved incident reporting procedures in accordance with 7 NMAC 1.13.
D. Staff providing direct services shall complete training in universal precautions on an annual basis. The training materials shall meet Occupational Safety and Health Administration (OSHA) requirements.
E. Staff providing direct services shall maintain certification in first aid and CPR. The training materials shall meet OSHA requirements/guidelines.
F. Staff who may be exposed to hazardous chemicals shall complete relevant training in accordance with OSHA requirements.
G. Staff shall be certified in a DDSD-approved behavioral intervention system (e.g., Mandt, CPI) before using physical restraint techniques. Staff members providing direct services shall maintain certification in a DDSD-approved behavioral intervention system if an individual they support has a behavioral crisis plan that includes the use of physical restraint techniques.
H. Staff shall complete and maintain certification in a DDSD-approved medication course in accordance with the DDSD Medication Delivery Policy M-001.
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.
<table>
<thead>
<tr>
<th>Tag # 1A26 (CoP) COR / EAR</th>
<th>Scope and Severity Rating: D</th>
</tr>
</thead>
</table>
| **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 2 of 7 Agency Personnel.  
**The following Agency Personnel records contained evidence that indicated the Employee Abuse Registry was completed after hire:**  
E. **Documentation for other staff.** With respect to all employed or contracted individuals providing direct care who are licensed health care professionals or certified nurse aides, the provider shall maintain documentation reflecting the individual's current licensure as a health care professional or current certification as a nurse aide.

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


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

**D. Criminal History Screening:** All personnel shall be screened by the Provider Agency in regard to the employee's qualifications, references, and employment history, prior to employment. All Provider Agencies shall comply with the Criminal Records Screening for Caregivers 7.1.12 NMAC and Employee Abuse Registry 7.1.12 NMAC as required by the Department of Health, Division of Health Improvement.
<table>
<thead>
<tr>
<th>Tag # 1A29  Complaints / Grievances - Acknowledgement</th>
<th>Scope and Severity Rating: A</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>NMAC 7.26.3.6</strong>&lt;br&gt;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 8 individuals.</td>
</tr>
</tbody>
</table>

- **Grievance/Complaint Procedure Acknowledgement (#5)**

**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:**<br>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
**ADDITIONAL FINDINGS: Reimbursement Deficiencies**

**BILLING**
**TAG #1A12**


**Chapter 1. III. PROVIDER AGENCY DOCUMENTATION OF SERVICE DELIVERY AND LOCATION**

**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:

1. Date, start and end time of each service encounter or other billable service interval;
2. A description of what occurred during the encounter or service interval; and
3. The signature or authenticated name of staff providing the service.

Billing for Community Inclusion (Supported Employment) services was reviewed for 8 of 8 individuals. Progress notes and billing records supported billing activities for the months of April, May and June 2010.