Dear Ms. Lozano:

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

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

**Compliance with all Conditions of Participation.**

This determination is based on your agency’s compliance with CMS waiver assurances at the Condition of Participation level. The attached QMB Report of Findings indicates Standard Level deficiencies identified and requires implementation of a Plan of Correction.

**Plan of Correction:**
The attached Report of Findings identifies the Standard Level and/or Condition of Participation deficiencies found during your agency’s compliance review. You are required to complete and implement a Plan of Correction.
agency has a total of 45 business days (10 business days to submit your POC for approval and 35 days to implement your *approved* Plan of Correction) from the receipt of this letter.

**Submission of your Plan of Correction:**
Please submit your agency’s Plan of Correction in the space on the two right columns of the Report of Findings. *(See attachment “A” for additional guidance in completing the Plan of Correction)*.

Within 10 business days of receipt of this letter your agency Plan of Correction must be submitted to the parties below:

1. Quality Management Bureau, Attention: Plan of Correction Coordinator  
   5301 Central Ave. NE Suite 400 Albuquerque, NM 87108  
2. Developmental Disabilities Supports Division Regional Office for region of service surveyed

Upon notification from QMB that your *Plan of Correction has been approved*, you must implement all remedies and corrective actions to come into compliance. If your Plan of Correction is denied, you must resubmit a revised plan as soon as possible for approval, as your POC approval and all remedies must be completed within 45 business days of the receipt of this letter.

Failure to submit your POC within the allotted 10 business days or complete and implement your Plan of Correction within the total 45 business days allowed may result in the imposition of a $200 per day Civil Monetary Penalty until it is received, completed and/or implemented.

**Request for Informal Reconsideration of Findings (IRF):**
If you disagree with a finding of deficient practice, you have 10 business days upon receipt of this notice to request an IRF. Submit your request for an IRF in writing to:

QMB Deputy Bureau Chief  
5301 Central Ave NE Suite #400  
Albuquerque, NM  87108  
Attention: IRF request

See Attachment "C" for additional guidance in completing the request for Informal Reconsideration of Findings. The request for an IRF will not delay the implementation of your Plan of Correction which must be completed within 45 total business days (10 business days to submit your POC for approval and 35 days to implement your *approved* Plan of Correction). Providers may not appeal the nature or interpretation of the standard or regulation, the team composition or sampling methodology. If the IRF approves the modification or removal of a finding, you will be advised of any changes.

Please call the Plan of Correction Coordinator at 505-222-8647 if you have questions about the Report of Findings or Plan of Correction. Thank you for your cooperation and for the work you perform.

Sincerely,

*Deb Russell, BS*

Deb Russell, BS  
Team Lead/Healthcare Surveyor  
Division of Health Improvement  
Quality Management Bureau
### Survey Process Employed:

**Entrance Conference Date:** January 30, 2012  
**Present:**  
Casa Alegre (MEND, Inc.)  
Vicky Lozano, Administrator  
Linda Lozano, Assistant Administrator  

**DOH/DHI/QMB**  
Deb Russell, BS, Team Lead/Healthcare Surveyor  
Mari Chavez, BSW, Healthcare Surveyor  

**Exit Conference Date:** January 31, 2012  
**Present:**  
Casa Alegre (MEND, Inc.)  
Vicky Lozano, Administrator  
Linda Lozano, Assistant Administrator  

**DOH/DHI/QMB**  
Deb Russell, BS, Team Lead/Healthcare Surveyor  
Mari Chavez, BSW, Healthcare Surveyor  

**DDSD - Southeast Regional Office**  
Jessica Renteria, Regional Office Manager (by telephone conference)  

| **Total Homes Visited** | **Number:** 2  
| **Supported Homes Visited** | **Number:** 2 (Note: Individuals #1, 3 & 4 share a residence)  
| **Administrative Locations Visited** | **Number:** 1  
| **Total Sample Size** | **Number:** 4  
| 1 - *Jackson* Class Members  
3 - Non-*Jackson* Class Members  
4 - Supported Living  
4 - Adult Habilitation  
3 - Supported Employment  
| **Persons Served Records Reviewed** | **Number:** 4  
| **Persons Served Interviewed** | **Number:** 2  
| **Persons Served Observed** | **Number:** 2 (2 Individuals did not respond to surveyor questions)  
| **Direct Support Personnel Interviewed** | **Number:** 3  
| **Direct Support Personnel Records Reviewed** | **Number:** 18  
| **Service Coordinator Records Reviewed** | **Number:** 2  
| **Administrative Files Reviewed** |  
- Billing Records  
- Medical Records  
- Incident Management Records  
- Personnel Files  
- Training Records  
- Agency Policy and Procedure  

Survey Report #: Q12.03.D0161.SE.001.RTN.01
• Caregiver Criminal History Screening Records
• Employee Abuse Registry
• Human Rights Notes and/or Meeting Minutes Evacuation Drills
• Quality Assurance / Improvement Plan

CC: Distribution List:  
DOH - Division of Health Improvement  
DOH - Developmental Disabilities Supports Division  
DOH - Office of Internal Audit  
HSD - Medical Assistance Division
Attachment A

Provider Instructions for Completing the QMB Plan of Correction (POC) Process

Introduction:
After a QMB Compliance Survey, your QMB Report of Findings will be sent to you via e-mail.

Each provider must develop and implement a Plan of Correction (POC) that identifies specific quality assurance and quality improvement activities the agency will implement to correct deficiencies and prevent continued non compliance.

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

Agencies must fully implement their approved Plan of Correction within 45 business days (10 business days to submit your POC for approval and 35 days to implement your approved Plan of Correction) from the date they receive the QMB Report of Findings. (Providers who fail to complete a POC within the 45 business days allowed shall be referred to the IRC for possible actions or sanctions.)

If you have questions about the Plan of Correction process, call the QMB Plan of Correction Coordinator at 505-222-8647 or email at George.Perrault@state.nm.us. Requests for technical assistance must be requested through your DDSD Regional Office.

The POC process cannot resolve disputes regarding findings. If you wish to dispute a finding on the official Report of Findings, you must file an Informal Reconsideration of Findings (IRF) request within ten (10) business days of receiving your report. Please note that you must still submit a POC for findings that are in question (see Attachment “C”).

Instructions for Completing Agency POC:

Required Content
Your Plan of Correction should provide a step-by-step description of the methods to correct each deficient practice to prevent recurrence and information that ensures the regulation cited is in compliance. The remedies noted in your POC are expected to be added to your Agency’s required, annual Quality Assurance Plan.

If a deficiency has already been corrected, the plan should state how it was corrected, the completion date (date the correction was accomplished), and how possible recurrence of the deficiency will be prevented.

The Plan of Correction must address the required six CMS core elements to address each deficiency of the POC:

1. How the specific and realistic corrective action will be accomplished for individuals found to have been affected by the deficient practice.
2. How the agency will identify other individuals who have the potential to be affected by the same deficient practice, and how the agency will act to protect individuals in similar situations.
3. What QA measures will be put into place or systemic changes made to ensure that the deficient practice will not recur
4. Indicate how the agency plans to monitor its performance to make sure that solutions are sustained. The agency must develop a QA plan for ensuring that correction is achieved and
sustained. This QA plan must be implemented, and the corrective action evaluated for its effectiveness. The plan of correction is integrated into the agency quality assurance system; and

5. Include dates when corrective action will be completed. The corrective action completion dates must be acceptable to the State.

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

The following details should be considered when developing your POC:

- Details about how and when Consumer, Personnel and Residential files are audited by Agency personnel to ensure they contain required documents;
- Information about how Medication Administration Records are reviewed to verify they contain all required information before they are distributed, as they are being used, and after they are completed;
- Your processes for ensuring that all staff are trained in Core Competencies, Incident Reporting, and Individual-Specific service requirements, etc;
- How accuracy in Billing documentation is assured;
- How health, safety is assured;
- For Case Management Providers, how ISPs are reviewed to verify they meet requirements, how the timeliness of LOC packet submissions and consumer visits are tracked;
- Your process for ensuring that all staff are trained in Core Competencies, Incident Reporting, and Individual-Specific service requirements, etc;
- How accuracy in Billing documentation is assured;
- How health, safety is assured;
- For Case Management Providers, how ISPs are reviewed to verify they meet requirements, how the timeliness of LOC packet submissions and consumer visits are tracked;
- Your process for ensuring that all staff are trained in Core Competencies, Incident Reporting, and Individual-Specific service requirements, etc;
- Details about Quality Targets in various areas, current status, analyses about why targets were not met, and remedies implemented.

**Note:** Instruction or in-service of staff alone may not be a sufficient plan of correction. This is a good first step toward correction, but additional steps should be taken to ensure the deficiency is corrected and will not recur.

**Completion Dates**

- The plan of correction must include a completion date (entered in the far right-hand column) for each finding. Be sure the date is realistic in the amount of time your Agency will need to correct the deficiency; not to exceed 45 total business days.
- Direct care issues should be corrected immediately and monitored appropriately.
- Some deficiencies may require a staged plan to accomplish total correction.
- Deficiencies requiring replacement of equipment, etc., may require more time to accomplish correction but should show reasonable time frames.

**Initial Submission of the Plan of Correction Requirements**

1. The Plan of Correction must be completed on the official QMB Survey Report of Findings/Plan of Correction Form and received by QMB within ten (10) business days from the date you received the report of findings.
2. For questions about the POC process, call the POC Coordinator, George Perrault at 505-222-8647 for assistance.
3. For Technical Assistance (TA) in developing or implementing your POC, contact your local DDSD Regional Office.
4. Submit your POC to George Perrault, POC Coordinator in any of the following ways:
   a. Electronically at George.Perrault@state.nm.us (preferred method)
   b. Fax to 505-222-8661, or
   c. Mail to POC Coordinator, 5301 Central Avenue SW, Suite 400, Albuquerque, NM 87108
5. Do not submit supporting documentation (evidence of compliance) to QMB until after your POC has been approved by the POC Coordinator.
6. QMB will notify you when your POC has been “approve” or “denied.”
a. During this time, whether your POC is “approved,” or “denied,” you will have a maximum of 45 business days from the date of receipt of your Report of Findings to correct all survey deficiencies.

b. If your POC is denied, it must be revised and resubmitted as soon as possible, as the 45 business day limit is in effect.

c. If your POC is denied a second time your agency may be referred to the Internal Review Committee.

d. You will receive written confirmation when your POC has been approved by QMB and a final deadline for completion of your POC.

7. Failure to submit your POC within 10 business days without prior approval of an extension by QMB will result in a referral to the Internal Review Committee and the possible implementation of monetary penalties and/or sanctions.

**POC Document Submission Requirements**

Once your POC has been approved by the QMB Plan of Correction Coordinator you must submit copies of documents as evidence that all deficiencies have been corrected, as follows.

1. Your internal documents are due within a **maximum** of 45 business days of receipt of your Report of Findings.

2. You may submit your documents by postal mail (paper hard copy or on a disc), fax, or electronically (scanned and attached to e-mails).

3. All submitted documents **must be annotated**; please be sure the tag numbers and Identification numbers are indicated on each document submitted. Documents which are not annotated with the Tag number and Identification number may not be accepted.

4. Do not submit original documents; Please provide copies or scanned electronic files for evidence. Originals must be maintained in the agency file(s) per DDSD Standards.

5. In lieu of some documents, you may submit copies of file or home audit forms that clearly indicate cited deficiencies have been corrected, other attestations of correction must be approved by the Plan of Correction Coordinator prior to their submission.

6. For billing deficiencies, you must submit:
   a. Evidence of an internal audit of billing documentation for a sample of individuals and timeframes;
   b. Copies of “void and adjust” forms submitted to correct all over-billed or unjustified units billed identified during your internal audit.

Revisions, Modifications or Extensions to your Plan of Correction (post QMB approval) must be made in writing and submitted to the Plan of Correction Coordinator at QMB, prior to the due date and are approved on a case-by-case basis. No changes may be made to your POC or the timeframes for implementation without written approval of the POC Coordinator.
QMB Determinations of Compliance

- “Compliance with Conditions of Participation”
  The QMB determination of “Compliance with Conditions of Participation,” indicates that a provider is in compliance with all ‘Conditions of Participation,’ (CoP) but may have standard level deficiencies (deficiencies which are not at the condition level) out of compliance. The agency has obtained a level of compliance such that there is a minimal potential for harm to individuals’ health and safety. To qualify for a determination of Compliance with Conditions of Participation, the provider must be in compliance with all Conditions of Participation.

- “Partial-Compliance with Conditions of Participation”
  The QMB determination of “Partial-Compliance with Conditions of Participation” indicates that a provider is out of compliance with one (1) to three (3) ‘Conditions of Participation.’ This partial-compliance, if not corrected, may result in a serious negative outcome or the potential for more than minimal harm to individuals’ health and safety. The Agency may also have standard level deficiencies (deficiencies which are not at the condition level).

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

- “Non-Compliant with Conditions of Participation”:
  The QMB determination of “Non-Compliance with Conditions of Participation,” indicates a provider is significantly out of compliance with Conditions of Participation and/or has:
  - Four (4) Conditions of Participation out of compliance.
  - Multiple findings of widespread non-compliance with any standard or regulation with a significant potential for more than minimal harm.
  - Any finding of actual harm or Immediate Jeopardy.

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

  Providers receiving a repeat determination of 'Non-Compliance’ will be referred by Quality Management Bureau to the Internal Review Committee (IRC) for consideration of remedies and possible actions.
Guidelines for the Provider
Informal Reconsideration of Finding (IRF) Process

Introduction:
Throughout the QMB Survey process, surveyors are openly communicating with providers. Open communication means surveyors have clarified issues and/or requested missing information before completing the review through the use of the signed/dated “Document Request,” or “administrative Needs,” etc. forms. Regardless, there may still be instances where the provider disagrees with a specific finding. Providers may use the following process to informally dispute a finding.

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

The following limitations apply to the IRF process:
- The request for an IRF and all supporting evidence must be received within 10 business days.
- Findings based on evidence requested during the survey and not provided may not be subject to reconsideration.
- The supporting documentation must be new evidence not previously reviewed or requested by the survey team.
- Providers must continue to complete their Plan of Correction during the IRF process.
- Providers may not request an IRF to challenge the sampling methodology.
- Providers may not request an IRF based on disagreement with the nature of the standard or regulation.
- Providers may not request an IRF to challenge the team composition.
- Providers may not request an IRF to challenge the DHI/QMB determination of compliance or the length of their DDSD provider contract.

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

The IRF Committee will review the request, the Provider will be notified in writing of the ruling; no face-to-face meeting will be conducted.

When a Provider requests that a finding be reconsidered, it does not stop or delay the Plan of Correction process. Providers must continue to complete the Plan of Correction, including the finding in dispute regardless of the IRF status. If a finding is removed or modified, it will be noted and removed or modified from the Report of Findings. It should be noted that in some cases a Plan of Correction may be completed prior to the IRF process being completed. The provider will be notified in writing on the decisions of the IRF committee.
### Standard of Care

**CMS Assurance – Service Plans: ISP Implementation** – *Services are delivered in accordance with the service plan, including type, scope, amount, duration and frequency specified in the service plan.*

<table>
<thead>
<tr>
<th>Tag # 1A32 &amp; 6L14 ISP Implementation</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>NMAC 7.26.5.16.C and D Development of the ISP. Implementation of the ISP. The ISP shall be implemented according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan.</td>
<td>Based on record review, the Agency failed to implement the ISP according to the timelines determined by the IDT and as specified in the ISP for each stated desired outcomes and action plan for 1 of 4 individuals.</td>
</tr>
<tr>
<td>C. The IDT shall review and discuss information and recommendations with the individual, with the goal of supporting the individual in attaining desired outcomes. The IDT develops an ISP based upon the individual's personal vision statement, strengths, needs, interests and preferences. The ISP is a dynamic document, revised periodically, as needed, and amended to reflect progress towards personal goals and achievements consistent with the individual's future vision. This regulation is consistent with standards established for individual plan development as set forth by the commission on the accreditation of rehabilitation facilities (CARF) and/or other program accreditation approved and adopted by the developmental disabilities division and the department of health. It is the policy of the developmental disabilities division (DDD), that to the extent permitted by</td>
<td></td>
</tr>
<tr>
<td>Provider: State your Plan of Correction for the findings in this Tag above this line.</td>
<td></td>
</tr>
<tr>
<td>Enter your Quality Assurance/Quality Improvement processes below the line.</td>
<td></td>
</tr>
</tbody>
</table>

**Evidence**

- Per Individuals ISP the following was found with regards to the implementation of ISP Outcomes:

#### Administrative Files Reviewed:

- **Supported Living Data Collection/Data Tracking/Progress with regards to ISP Outcomes:**
  - Individual #4
  - Per Live Outcome; Actions Steps for “Participate/Play in Exercise Game” is to be completed 5 times per week. Evidence indicated it was not being completed at the required frequency indicated in the ISP for November 2011 and December 2011.
funding, each individual receive supports and services that will assist and encourage independence and productivity in the community and attempt to prevent regression or loss of current capabilities. Services and supports include specialized and/or generic services, training, education and/or treatment as determined by the IDT and documented in the ISP.

D. The intent is to provide choice and obtain opportunities for individuals to live, work and play with full participation in their communities. The following principles provide direction and purpose in planning for individuals with developmental disabilities.

[05/03/94; 01/15/97; Recompiled 10/31/01]
<table>
<thead>
<tr>
<th>Tag # 6L14 Residential Case File</th>
<th>Standard Level Deficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>Developmental Disabilities (DD) Waiver Service Standards effective 4/1/2007</td>
<td>Based on record review, the Agency failed to maintain a complete and confidential case file in the residence for 2 of 4 Individuals receiving Supported Living Services.</td>
</tr>
<tr>
<td>CHAPTER 6. VIII. COMMUNITY LIVING SERVICE PROVIDER AGENCY REQUIREMENTS</td>
<td>The following was not found, incomplete and/or not current:</td>
</tr>
<tr>
<td>A. Residence Case File: For individuals receiving Supported Living or Family Living, the Agency shall maintain in the individual’s home a complete and current confidential case file for each individual. For individuals receiving Independent Living Services, rather than maintaining this file at the individual's home, the complete and current confidential case file for each individual shall be maintained at the agency's administrative site. Each file shall include the following:</td>
<td></td>
</tr>
<tr>
<td>(1) Complete and current ISP and all supplemental plans specific to the individual;</td>
<td></td>
</tr>
<tr>
<td>(2) Complete and current Health Assessment Tool;</td>
<td></td>
</tr>
<tr>
<td>(3) Current emergency contact information, which includes the individual’s address, telephone number, names and telephone numbers of residential Community Living Support providers, relatives, or guardian or conservator, primary care physician’s name(s) and telephone number(s), pharmacy name, address and telephone number and dentist name, address and telephone number, and health plan;</td>
<td></td>
</tr>
<tr>
<td>(4) Up-to-date progress notes, signed and dated by the person making the note for at least the past month (older notes may be transferred to the agency office);</td>
<td></td>
</tr>
<tr>
<td>(5) Data collected to document ISP Action Plan implementation</td>
<td></td>
</tr>
<tr>
<td>(6) Progress notes written by direct care staff</td>
<td></td>
</tr>
</tbody>
</table>

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

Enter your Quality Assurance/Quality Improvement processes below the line.
and by nurses regarding individual health status
and physical conditions including action taken in
response to identified changes in condition for at
least the past month;
(7) Physician’s or qualified health care providers
written orders;
(8) Progress notes documenting implementation
of a physician’s or qualified health care
provider’s order(s);
(9) Medication Administration Record (MAR) for
the past three (3) months which includes:
(a) The name of the individual;
(b) A transcription of the healthcare
practitioners prescription including the
brand and generic name of the medication;
(c) Diagnosis for which the medication is
prescribed;
(d) Dosage, frequency and method/route of
delivery;
(e) Times and dates of delivery;
(f) Initials of person administering or assisting
with medication; and
(g) An explanation of any medication
irregularity, allergic reaction or adverse
effect.
(h) For PRN medication an explanation for the
use of the PRN must include:
(i) Observable signs/symptoms or
circumstances in which the medication
is to be used, and
(ii) Documentation of the
effectiveness/result of the PRN
delivered.
(i) A MAR is not required for individuals
participating in Independent Living Services
who self-administer their own medication.
However, when medication administration
is provided as part of the Independent
Living Service a MAR must be maintained
at the individual’s home and an updated
copy must be placed in the agency file on a
weekly basis.
(10) Record of visits to healthcare practitioners including any treatment provided at the visit and a record of all diagnostic testing for the current ISP year; and
(11) Medical History to include: demographic data, current and past medical diagnoses including the cause (if known) of the developmental disability and any psychiatric diagnosis, allergies (food, environmental, medications), status of routine adult health care screenings, immunizations, hospital discharge summaries for past twelve (12) months, past medical history including hospitalizations, surgeries, injuries, family history and current physical exam.
### Standard of Care

**CMS Assurance – Health and Welfare** – The state, on an ongoing basis, identifies, addresses and seeks to prevent occurrences of abuse, neglect and exploitation. Individuals shall be afforded their basic human rights. The provider supports individuals to access needed healthcare services in a timely manner.

### Tag # 1A15.2 & 5I09 - Healthcare Documentation

**Standard Level Deficiency**

Based on record review, the Agency failed to maintain the required documentation in the Individuals Agency Record as required per standard for 1 of 4 individual. The following were not found, incomplete and/or not current:

- **Health Care Plans**
- **Bowel & Bladder**

  Individual #3 - According to Electronic Comprehensive Health Assessment Tool the individual is required to have a plan. No evidence of a plan found.

**Provider:**

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

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

<table>
<thead>
<tr>
<th>Standard of Care</th>
<th>Deficiencies</th>
<th>Agency Plan of Correction, On-going QA/QI &amp; Responsible Party</th>
<th>Date Due</th>
</tr>
</thead>
</table>
| **CMS Assurance – Health and Welfare** – The state, on an ongoing basis, identifies, addresses and seeks to prevent occurrences of abuse, neglect and exploitation. Individuals shall be afforded their basic human rights. The provider supports individuals to access needed healthcare services in a timely manner.** | Based on record review, the Agency failed to maintain the required documentation in the Individuals Agency Record as required per standard for 1 of 4 individual. The following were not found, incomplete and/or not current:  
  - **Health Care Plans**  
  - **Bowel & Bladder**  
  Individual #3 - According to Electronic Comprehensive Health Assessment Tool the individual is required to have a plan. No evidence of a plan found. | | |
arranges for the individual’s Primary Care Practitioner (PCP) to voluntarily complete these assessments in lieu of the agency nurse. Agency nurses may also complete these assessments in collaboration with the Primary Care Practitioner if they believe such consultation is necessary for an accurate assessment. Family Living Provider Agencies have the option of having the subcontracted caregiver complete the HAT instead of the nurse or PCP, if the caregiver is comfortable doing so. However, the agency nurse must be available to assist the caregiver upon request.

(c) For newly allocated individuals, the HAT and the MAAT must be completed within seventy-two (72) hours of admission into direct services or two weeks following the initial ISP, whichever comes first.

(d) For individuals already in services, the HAT and the MAAT must be completed at least fourteen (14) days prior to the annual ISP meeting and submitted to all members of the interdisciplinary team. The HAT must also be completed at the time of any significant change in clinical condition and upon return from any hospitalizations. In addition to annually, the MAAT must be completed at the time of any significant change in clinical condition, when a medication regime or route change requires delivery by licensed or certified staff, or when an individual has completed additional training designed to improve their skills to support self-administration (see DDSD Medication Assessment and Delivery Policy).

(e) Nursing assessments conducted to determine current health status or to evaluate a change in clinical condition must be documented in a signed progress note that includes time and date as well as subjective information including the individual complaints, signs and symptoms noted by staff, family members or other team members; objective
information including vital signs, physical examination, weight, and other pertinent data for the given situation (e.g., seizure frequency, method in which temperature taken); assessment of the clinical status, and plan of action addressing relevant aspects of all active health problems and follow up on any recommendations of medical consultants.

(2) Health related plans
(a) For individuals with chronic conditions that have the potential to exacerbate into a life-threatening situation, a medical crisis prevention and intervention plan must be written by the nurse or other appropriately designated healthcare professional.
(b) Crisis prevention and intervention plans must be written in user-friendly language that is easily understood by those implementing the plan.
(c) The nurse shall also document training regarding the crisis prevention and intervention plan delivered to agency staff and other team members, clearly indicating competency determination for each trainee.
(d) If the individual receives services from separate agencies for community living and community inclusion services, nurses from each agency shall collaborate in the development of and training delivery for crisis prevention and intervention plans to assure maximum consistency across settings.
(3) For all individuals with a HAT score of 4, 5 or 6, the nurse shall develop a comprehensive healthcare plan that includes health related supports identified in the ISP (The healthcare plan is the equivalent of a nursing care plan; two separate documents are not required nor recommended):
(a) Each healthcare plan must include a statement of the person’s healthcare needs and list measurable goals to be achieved through implementation of the healthcare plan.
Needs statements may be based upon supports needed for the individual to maintain a current strength, ability or skill related to their health, prevention measures, and/or supports needed to remediate, minimize or manage an existing health condition.

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

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

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

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

(f) Healthcare plans must be updated to reflect relevant discharge orders whenever an individual returns to services following a hospitalization.

(g) All crisis prevention and intervention plans and healthcare plans shall include the individual’s name and date on each page and
shall be signed by the author.

(h) Crisis prevention and intervention plans as well as healthcare plans shall be reviewed by the nurse at least quarterly, and updated as needed.

(4) General Nursing Documentation
(a) The nurse shall complete legible and signed progress notes with date and time indicated that describe all interventions or interactions conducted with individuals served as well as all interactions with other healthcare providers serving the individual. All interactions shall be documented whether they occur by phone or in person.
(b) For individuals with a HAT score of 4, 5 or 6, or who have identified health concerns in their ISP, the nurse shall provide the interdisciplinary team with a quarterly report that indicates current health status and progress to date on health related ISP desired outcomes and action plans as well as progress toward goals in the healthcare plan.
### Standard of Care

**Deficiencies**

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

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#### CMS Assurance – Financial Accountability

*State financial oversight exists to assure that claims are coded and paid for in accordance with the reimbursement methodology specified in the approved waiver.*

#### TAG #1A12 All Services Reimbursement (No Deficiencies)


**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 Living (Supported Living) and Community Inclusion (Adult Habilitation & Supported Employment) services was reviewed for 4 of 4 individuals. *Progress notes and billing records supported billing activities for the months of October, November & December 2011.*
Dear Ms. Lozano:

The Division of Health Improvement Quality Management Bureau received, reviewed and approved the documents you submitted for your Plan of Correction. The documents you provided verified that survey deficiencies were corrected.

**Your Plan of Correction is closed.**

To maintain ongoing compliance with standards and regulations, continue to use the Quality Improvement/Quality Assurance processes in your Plan of Correction, including:

- Consumer files audited monthly to ensure that Outcome frequencies are being met according to each individual’s ISP, and that required documents are in place
Consistent implementation of your Quality Assurance/Quality Improvement processes will enable you to identify and promptly respond to problems, enhance your service delivery, and result in fewer deficiencies in future QMB surveys.

Thank you for your cooperation with the Plan of Correction process, and for the work you and your team perform.

Sincerely,

George Perrault, MBA
Plan of Correction Coordinator

Cc: DHI
    DDSD