NEW MEXICO
Department of Health
Division of Health Improvement

MICHELLE LUJAN GRISHAM
Governor

DAVID R. SCRASE, M.D.
Acting Cabinet Secretary

Date:	October 20, 2022
To:	Emad Elmaoued, Executive Director
Provider:	ADID Care, INC
Address:	5115 Copper Ave NE
State/Zip:	Albuquerque, New Mexico 87108
E-mail Address:	emad@adidcare.com
Region:	Metro & Northeast
Routine Survey:	February 21 – March 3, 2022
Verification Survey:	September 19 – 28, 2022
Program Surveyed:	Developmental Disabilities Waiver
Service Surveyed:	Supported Living, Family Living, Customized In-Home Supports, Customized Community Supports
Survey Type:	Verification
Team Leader:	Kayla R. Benally, BSW, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau
Team Members:	Elizabeth Vigil, Healthcare Surveyor, Division of Health Improvement/Quality Management Bureau

Dear Mr. Emad Elmaoued;

The Division of Health Improvement/Quality Management Bureau has completed a Verification survey of the services identified above. The purpose of the survey was to determine compliance with your Plan of Correction submitted to DHI regarding the *Routine Survey on February 21 – March 3, 2022*.

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

Partial Compliance with Standard Level Tags and Conditions of Participation Level Tags: This determination is based on noncompliance with one to five (1 – 5) Condition of Participation Level Tags *(refer to Attachment D for details)*. The attached QMB Report of Findings indicates Standard Level and Condition of Participation Level deficiencies identified and requires completion and implementation of a Plan of Correction.

The following tags are identified as Condition of Participation Level:

- Tag # 1A09 Medication Delivery Routine Medication Administration **(New / Repeat Findings)**
- Tag # 1A09.2 Medication Delivery Nurse Approval for PRN Medication **(New Findings)**

The following tags are identified as Standard Level:

- Tag # 1A09.1.0 Medication Delivery PRN Medication Administration **(New Findings)**

However, due to the new/repeat deficiencies your agency will be required to contact your DDSD Regional Office for technical assistance and follow up and complete the Plan of Correction document attached at the end of this report. Please respond to the Plan of Correction Coordinator within 10 business days of receipt of this letter.

Plan of Correction:
The attached Report of Findings identifies the new/repeat Standard Level deficiencies found during your agency's verification compliance review. You are required to complete and implement a Plan of Correction. Your agency has a total of 10 business days from the receipt of this letter. The Plan of Correction must include the following:

1. Evidence your agency has contacted your DDSD Regional Office for technical assistance;
2. A Plan of Correction detailing Quality Assurance/Quality Improvement processes to prevent your agency from receiving deficiencies in the future. Please use the format provided at the end of this report;
3. Documentation verifying that newly cited deficiencies have been corrected.

Submission of your Plan of Correction:
Please submit your agency's Plan of Correction and documentation verifying correction of survey deficiencies within 10 business days of receipt of this letter to the parties below:

1. **Quality Management Bureau, Attention: Plan of Correction Coordinator**
   5301 Central Ave. NE Suite 400, New Mexico 87108
   MonicaE.Valdez@state.nm.us

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

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

Please call the Plan of Correction Coordinator Monica Valdez at 505-273-1930 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,

Kayla R. Benally, BSW

Kayla R. Benally, BSW
Team Lead/Healthcare Surveyor
Division of Health Improvement
Quality Management Bureau

Survey Process Employed:

Administrative Review Start Date:	September 19, 2022
Contact:	**ADID Care, INC** Emad Elmaoued, Executive Director **DOH/DHI/QMB** Kayla R. Benally, BSW, Team Lead/Healthcare Surveyor
On-site Entrance Conference Date:	*Entrance conference was waived by provider*
Exit Conference Date:	September 28, 2022
Present:	**ADID Care, INC** Emad Elmaoued, Executive Director **DOH/DHI/QMB** Kayla R. Benally, BSW, Team Lead/Healthcare Surveyor Elizabeth Vigil, Healthcare Surveyor Wolf Krusemark, BFA, Healthcare Surveyor Supervisor **DDSD - Metro and NE Regional Office** Linda Clark, Metro Assistant Regional Manager Angela Pacheco, NE Regional Director
Total Sample Size:	8 0 - *Jackson* Class Members 8 - Non-*Jackson* Class Members 3 - Supported Living 4 - Family Living 1 - Customized In-Home Supports 4 - Customized Community Supports
Persons Served Records Reviewed	8
Direct Support Personnel Records Reviewed	67
Direct Support Personnel Interviewed during Routine Survey	8 *(Note: Interviews conducted by video / phone due to COVID-19 Public Health Emergency)*
Substitute Care/Respite Personnel Records Reviewed	9
Service Coordinator Records Reviewed	1
Nurse Interview completed during Routine Survey	1

Administrative Processes and Records Reviewed:

- Medicaid Billing/Reimbursement Records for all Services Provided
- Accreditation Records
- Oversight of Individual Funds

- Individual Medical and Program Case Files, including, but not limited to:
    - °Individual Service Plans
    - °Progress on Identified Outcomes
    - °Healthcare Plans
    - °Medication Administration Records
    - °Medical Emergency Response Plans
    - °Therapy Evaluations and Plans
    - °Healthcare Documentation Regarding Appointments and Required Follow-Up
    - °Other Required Health Information
- Internal Incident Management Reports and System Process / General Events Reports
- Personnel Files, including nursing and subcontracted staff
- Staff Training Records, Including Competency Interviews with Staff
- Agency Policy and Procedure Manual
- Caregiver Criminal History Screening Records
- Consolidated Online Registry/Employee Abuse Registry
- Human Rights Committee Notes and Meeting Minutes
- Evacuation Drills of Residences and Service Locations
- 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
NM Attorney General's Office

Attachment B

Department of Health, Division of Health Improvement
QMB Determination of Compliance Process

The Division of Health Improvement, Quality Management Bureau (QMB) surveys compliance of the Developmental Disabilities Waiver (DDW) standards and other state and federal regulations. For the purpose of the LCA / CI survey the CMS waiver assurances have been grouped into four (4) Service Domains: Plan of Care (ISP Implementation); Qualified Providers; Health, Welfare and Safety; and Administrative Oversight (note that Administrative Oversight listed in this document is not the same as the CMS assurance of Administrative Authority. Used in this context it is related to the agency's operational policies and procedures, Quality Assurance system and Medicaid billing and reimbursement processes.)

The QMB Determination of Compliance process is based on provider compliance or non-compliance with standards and regulations identified during the on-site survey process and as reported in the QMB Report of Findings. All areas reviewed by QMB have been agreed to by DDSD and DHI/QMB and are reflective of CMS requirements. All deficiencies (non-compliance with standards and regulations) are identified and cited as either a Standard level deficiency or a Condition of Participation level deficiency in the QMB Reports of Findings. All deficiencies require corrective action when non-compliance is identified.

Each deficiency in your Report of Findings has been predetermined to be a Standard Level Deficiency, a Condition of Participation Level Deficiency, if below 85% compliance or a non-negotiable Condition of Participation Level Deficiency. Your Agency's overall Compliance Determination is based on a Scope and Severity Scale which takes into account the number of Standard and Condition Level Tags cited as well as the percentage of Individuals affected in the sample.

Conditions of Participation (CoPs)

CoPs are based on the Centers for Medicare and Medicaid Services, Home and Community-Based Waiver required assurances, in addition to the New Mexico Developmental Disability Waiver (DDW) Service Standards. The Division of Health Improvement (DHI), in conjunction with the Developmental Disability Support Division (DDSD), has identified certain deficiencies that have the potential to be a Condition of Participation Level, if the tag falls below 85% compliance based on the number of people affected. Additionally, there are what are called non-negotiable Conditions of Participation, regardless if one person or multiple people are affected. In this context, a CoP is defined as an essential / fundamental regulation or standard, which when out of compliance directly affects the health and welfare of the Individuals served. If no deficiencies within a Tag are at the level of a CoP, it is cited as a Standard Level Deficiency.

Service Domains and CoPs for <u>Living Care Arrangements and Community Inclusion</u> are as follows:

<u>Service Domain: Service Plan: ISP Implementation -</u> *Services are delivered in accordance with the service plan, including type, scope, amount, duration and frequency specified in the service plan.*

Potential Condition of Participation Level Tags, if compliance is below 85%:

- **1A08.3 –** Administrative Case File: Individual Service Plan / ISP Components
- **1A32 –** Administrative Case File: Individual Service Plan Implementation
- **LS14 –** Residential Service Delivery Site Case File (ISP and Healthcare Requirements)
- **IS14 –** CCS / CIES Service Delivery Site Case File (ISP and Healthcare Requirements)

<u>Service Domain: Qualified Providers -</u> *The State monitors non-licensed/non-certified providers to assure adherence to waiver requirements. The State implements its policies and procedures for verifying that provider training is conducted in accordance with State requirements and the approved waiver.*

Potential Condition of Participation Level Tags, if compliance is below 85%:

- **1A20 -** Direct Support Personnel Training
- **1A22 -** Agency Personnel Competency

- **1A37 –** Individual Specific Training

Non-Negotiable Condition of Participation Level Tags (one or more Individuals are cited):
- **1A25.1 –** Caregiver Criminal History Screening
- **1A26.1 –** Consolidated On-line Registry Employee Abuse Registry

Service Domain: Health, Welfare and Safety - *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.*

Potential Condition of Participation Level Tags, if compliance is below 85%:
- **1A08.2 –** Administrative Case File: Healthcare Requirements & Follow-up
- **1A09 –** Medication Delivery Routine Medication Administration
- **1A09.1 –** Medication Delivery PRN Medication Administration
- **1A15.2 –** Administrative Case File: Healthcare Documentation (Therap and Required Plans)

Non-Negotiable Condition of Participation Level Tags (one or more Individuals are cited):
- **1A05 –** General Requirements / Agency Policy and Procedure Requirements
- **1A07 –** Social Security Income (SSI) Payments
- **1A09.2 –** Medication Delivery Nurse Approval for PRN Medication
- **1A15 –** Healthcare Coordination - Nurse Availability / Knowledge
- **1A31 –** Client Rights/Human Rights
- **LS25.1 –** Residential Reqts. (Physical Environment - Supported Living / Family Living / Intensive Medical Living)

Attachment C

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 received in writing to the QMB Bureau Chief **within 10 business days** of receipt of the final Report of Findings **(*Note: No extensions are granted for the IRF)*.**
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: https://nmhealth.org/about/dhi/cbp/irf/
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 IRF process, email the IRF Chairperson, Valerie V. Valdez at valerie.valdez@state.nm.us for assistance.

The following limitations apply to the IRF process:
- The written 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 received 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.

Attachment D

QMB Determinations of Compliance

Compliance:

The QMB determination of *Compliance* indicates that a provider has either no deficiencies found during a survey or that no deficiencies at the Condition of Participation Level were found. 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*, the provider must have received no Conditions of Participation Level Deficiencies and have a minimal number of Individuals on the sample affected by the findings indicated in the Standards Level Tags.

Partial-Compliance with Standard Level Tags:

The QMB determination of *Partial-Compliance with Standard Level Tags* indicates that a provider is in compliance with all Condition of Participation Level deficiencies but is out of compliance with a certain percentage of Standard Level deficiencies. This partial-compliance, if not corrected, may result in a negative outcome or the potential for more than minimal harm to individuals' health and safety. There are two ways to receive a determination of Partial Compliance with Standard Level Tags:

1. Your Report of Findings includes 16 or fewer Standards Level Tags with between 75% and 100% of the survey sample affected in any tag.

2. Your Report of Findings includes 17 or more Standard Level Tags with between 50% to 74% of the survey sample affected in any tag.

Partial-Compliance with Standard Level Tags and Condition of Participation Level Tags:

The QMB determination of *Partial-Compliance with Standard Level Tags and Condition of Participation Level Tags* indicates that a provider is out of compliance with one to five (1 – 5) Condition of Participation Level Tags. 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.

Non-Compliance:

The QMB determination of *Non-Compliance* indicates a provider is significantly out of compliance with both Standard Level deficiencies and Conditions of Participation level deficiencies. This non-compliance, if not corrected, may result in a serious negative outcome or the potential for more than minimal harm to individuals' health and safety. There are three ways an agency can receive a determination of Non-Compliance:

1. Your Report of Findings includes 17 or more total Tags with 0 to 5 Condition of Participation Level Tags with 75% to 100% of the survey sample affected in any Condition of Participation Level tag.

2. Your Report of Findings includes any amount of Standard Level Tags with 6 or more Condition of Participation Level Tags.

Compliance Determination	Weighting						
	LOW		MEDIUM			HIGH	
Total Tags:	up to 16	17 or more	up to 16	17 or more	Any Amount	17 or more	Any Amount
	and	and	and	and	And/or	and	And/or
COP Level Tags:	0 COP	0 COP	0 COP	0 COP	1 to 5 COP	0 to 5 CoPs	6 or more COP
	and	and	and	and		and	
Sample Affected:	0 to 74%	0 to 49%	75 to 100%	50 to 74%		75 to 100%	
"Non-Compliance"						**17 or more** Total Tags with **75 to 100%** of the Individuals in the sample cited in any CoP Level tag.	**Any Amount** of Standard Level Tags and **6 or more** Conditions of Participation Level Tags.
"Partial Compliance with Standard Level tags and Condition of Participation Level Tags"					**Any Amount** Standard Level Tags, plus **1 to 5** Conditions of Participation Level tags.		
"Partial Compliance with Standard Level tags"			**up to 16** Standard Level Tags with **75 to 100%** of the individuals in the sample cited in any tag.	**17 or more** Standard Level Tags with **50 to 74%** of the individuals in the sample cited any tag.			
"Compliance"	**Up to 16** Standard Level Tags with **0 to 74%** of the individuals in the sample cited in any tag.	**17 or more** Standard Level Tags with **0 to 49%** of the individuals in the sample cited in any tag.					

Agency: **ADID Care, INC – Metro and Northeast**
Program: Developmental Disabilities Waiver
Service: Supported Living, Family Living, Customized In-Home Supports and Customized Community Supports
Survey Type: Verification
Routine Survey: **February 21 – March 3, 2022**
Verification Survey: **September 19 – 28, 2022**

Standard of Care	Routine Survey Deficiencies February 21 – March 3, 2022	Verification Survey New and Repeat Deficiencies September 19 – 28, 2022
Service Domain: 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 # 1A09 Medication Delivery Routine Medication Administration	**Condition of Participation Level Deficiency**	**Condition of Participation Level Deficiency**
Developmental Disabilities (DD) Waiver Service Standards 2/26/2018; Re-Issue: 12/28/2018; Eff 1/1/2019 **Chapter 20: Provider Documentation and Client Records 20.6 Medication Administration Record (MAR):** A current Medication Administration Record (MAR) must be maintained in all settings where medications or treatments are delivered. Family Living Providers may opt not to use MARs if they are the sole provider who supports the person with medications or treatments. However, if there are services provided by unrelated DSP, ANS for Medication Oversight must be budgeted, and a MAR must be created and used by the DSP. Primary and Secondary Provider Agencies are responsible for: 1. Creating and maintaining either an electronic or paper MAR in their service setting. Provider Agencies may use the MAR in Therap, but are not mandated to do so. 2. Continually communicating any changes about medications and treatments between Provider Agencies to assure health and safety. 7. Including the following on the MAR: a. The name of the person, a transcription of the physician's or licensed health care provider's orders including the brand and generic names for all ordered routine and PRN medications or treatments, and the diagnoses for which the medications or	After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur. Medication Administration Records (MAR) were reviewed for the month of January 2022. Based on record review, 1 of 4 individuals had Medication Administration Records (MAR), which contained missing medications entries and/or other errors: Individual #4 January 2022 Medication Administration Records contained missing entries. No documentation found indicating reason for missing entries: • Butenafine HCL 1% cream (2 times daily) – Blank 1/23 (10:00 AM) • Carbamazepine 200 mg (2 times daily) – Blank 1/23 (8:00 AM) • Escitalopram Oxalate 20 mg (1 time daily) – Blank 1/23 (8:00 AM) • Genteal Tears Severe .3% (2 times daily) – Blank 1/23 (10:00 AM)	**New / Repeat Findings:** After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur. Medication Administration Records (MAR) were reviewed for the month of August 2022. Based on record review, 1 of 3 individuals had Medication Administration Records (MAR), which contained missing medications entries and/or other errors: Individual #6 August 2022 No Physician's Orders were found for medications listed on the Medication Administration Records for the following medications: • Oyster Shell Calcium 500 mg (1 time daily)

treatments are prescribed; b. The prescribed dosage, frequency and method or route of administration; times and dates of administration for all ordered routine or PRN prescriptions or treatments; over the counter (OTC) or "comfort" medications or treatments and all self-selected herbal or vitamin therapy; c. Documentation of all time limited or discontinued medications or treatments; d. The initials of the individual administering or assisting with the medication delivery and a signature page or electronic record that designates the full name corresponding to the initials; e. Documentation of refused, missed, or held medications or treatments; f. Documentation of any allergic reaction that occurred due to medication or treatments; and g. For PRN medications or treatments: i. instructions for the use of the PRN medication or treatment which must include observable signs/symptoms or circumstances in which the medication or treatment is to be used and the number of doses that may be used in a 24-hour period; ii. clear documentation that the DSP contacted the agency nurse prior to assisting with the medication or treatment, unless the DSP is a Family Living Provider related by affinity of consanguinity; and iii. documentation of the effectiveness of the PRN medication or treatment. **Chapter 10 Living Care Arrangements** ***10.3.4 Medication Assessment and Delivery:*** Living Supports Provider Agencies must support and comply with: 1. the processes identified in the DDSD AWMD	• Lisinopril 10 mg (1 time daily) – Blank 1/23 (9:00 AM) • Quetiapine Fumarate 25 mg (1 time daily) – Blank 1/23 (8:00 AM) • Vitamin D3 50 mcg (1 time daily) – Blank 1/23 (9:00 AM)	

training; 2. the nursing and DSP functions identified in the Chapter 13.3 Part 2- Adult Nursing Services; 3. all Board of Pharmacy regulations as noted in Chapter 16.5 Board of Pharmacy; and 4. documentation requirements in a Medication Administration Record (MAR) as described in Chapter 20.6 Medication Administration Record (MAR). **NMAC 16.19.11.8 MINIMUM STANDARDS:** A. MINIMUM STANDARDS FOR THE DISTRIBUTION, STORAGE, HANDLING AND RECORD KEEPING OF DRUGS: (d) The facility shall have a Medication Administration Record (MAR) documenting medication administered to residents, **including over-the-counter medications.** This documentation shall include: (i) Name of resident; (ii) Date given; (iii) Drug product name; (iv) Dosage and form; (v) Strength of drug; (vi) Route of administration; (vii) How often medication is to be taken; (viii) Time taken and staff initials; (ix) Dates when the medication is discontinued or changed; (x) The name and initials of all staff administering medications. **Model Custodial Procedure Manual** ***D. Administration of Drugs*** Unless otherwise stated by practitioner, patients will not be allowed to administer their own medications. Document the practitioner's order authorizing the self-administration of medications.		

All PRN (As needed) medications shall have complete detail instructions regarding the administering of the medication. This shall include: ➢ symptoms that indicate the use of the medication, ➢ exact dosage to be used, and ➢ the exact amount to be used in a 24-hour period.		

Tag # 1A09.1.0 Medication Delivery PRN Medication Administration		Standard Level Deficiency
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 10 Living Care Arrangements (LCA):** **10.3.5 Medication Assessment and Delivery:** Living Supports Provider Agencies must support and comply with: 1. the processes identified in the DDSD AWMD training; 2. the nursing and DSP functions identified in the Chapter 13.3 Adult Nursing Services; 3. all Board of Pharmacy regulations as noted in Chapter 16.5 Board of Pharmacy; and 4. documentation requirements in a Medication Administration Record (MAR) as described in Chapter 20 20.6 Medication Administration Record (MAR) **Chapter 20 Provider Documentation and Client Records: 20.6 Medication Administration Record (MAR):** Administration of medications apply to all provider agencies of the following services: living supports, customized community supports, community integrated employment, intensive medical living supports. 1. Primary and secondary provider agencies are to utilize the Medication Administration Record (MAR) online in Therap. 2. Providers have until November 1, 2022, to have a current Electronic Medication Administration Record online in Therap in all settings where medications or treatments are delivered. 3. Family Living Providers may opt not to use MARs if they are the **sole** provider who supports the person and are related by affinity or consanguinity. However, if there are services provided by unrelated DSP, ANS for Medication Oversight must be budgeted, a MAR online in Therap must be created and used by the DSP. 4. Provider Agencies must configure and use the MAR when assisting with medication.	**NA**	**New Findings:** Medication Administration Records (MAR) were reviewed for the month of August 2022. Based on record review, 2 of 3 individuals had PRN Medication Administration Records (MAR), which contained missing elements as required by standard: Individual #6 August 2022 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Ibuprofen 200 mg – PRN – 8/27 (given 1 time) • Loperamide 2 mg – PRN – 8/4 - 6 (given 1 time) Individual #7 August 2022 No Effectiveness was noted on the Medication Administration Record for the following PRN medication: • Acetaminophen 325 mg – PRN – 8/3 (given 2 times), 8/4 - 5, 9 - 12 (given 1 time) • Chlorpheniramine 4 mg – PRN – 8/3, 5, 9 – 12, 16 – 19, 23 - 26, 30 - 31 (given 1 time)

5. Provider Agencies Continually communicating any changes about medications and treatments between Provider Agencies to assure health and safety. 6. Provider agencies must include the following on the MAR: a. The name of the person, a transcription of the physician's or licensed health care provider's orders including the brand and generic names for all ordered routine and PRN medications or treatments, and the diagnoses for which the medications or treatments are prescribed. b. The prescribed dosage, frequency and method or route of administration; times and dates of administration for all ordered routine and PRN medications and other treatments; all over the counter (OTC) or "comfort" medications or treatments; all self-selected herbal preparation approved by the prescriber, and/or vitamin therapy approved by prescriber. c. Documentation of all time limited or discontinued medications or treatments. d. The initials of the person administering or assisting with medication delivery. e. Documentation of refused, missed, or held medications or treatments. f. Documentation of any allergic reaction that occurred due to medication or treatments. g. For PRN medications or treatments including all physician approved over the counter medications and herbal or other supplements: i. instructions for the use of the PRN medication or treatment which must include observable signs/symptoms or circumstances in which the medication or treatment is to be used and the number of doses that may be used in a 24-hour period; ii. clear follow-up detailed documentation that the DSP contacted the agency nurse prior to assisting with the medication or treatment; and iii. documentation of the effectiveness of the PRN medication or treatment.		

NMAC 16.19.11.8 MINIMUM STANDARDS:
A. MINIMUM STANDARDS FOR THE DISTRIBUTION, STORAGE, HANDLING AND RECORD KEEPING OF DRUGS:
(d) The facility shall have a Medication Administration Record (MAR) documenting medication administered to residents, **including over-the-counter medications.** This documentation shall include:

(i) Name of resident;
(ii) Date given;
(iii) Drug product name;
(iv) Dosage and form;
(v) Strength of drug;
(vi) Route of administration;
(vii) How often medication is to be taken;
(viii) Time taken and staff initials;
(ix) Dates when the medication is discontinued or changed;
(x) The name and initials of all staff administering medications.

Model Custodial Procedure Manual
D. Administration of Drugs
Unless otherwise stated by practitioner, patients will not be allowed to administer their own medications.
Document the practitioner's order authorizing the self-administration of medications.

All PRN (As needed) medications shall have complete detail instructions regarding the administering of the medication. This shall include:

- symptoms that indicate the use of the medication,
- exact dosage to be used, and
- the exact amount to be used in a 24-hour period.

Tag # 1A09.2 Medication Delivery Nurse Approval for PRN Medication		Condition of Participation Level Deficiency
Developmental Disabilities Waiver Service Standards Eff 11/1/2021 **Chapter 10 Living Care Arrangements (LCA): 10.3.5 Medication Assessment and Delivery:** Living Supports Provider Agencies must support and comply with: 1. the processes identified in the DDSD AWMD training; 2. the nursing and DSP functions identified in the Chapter 13.3 Adult Nursing Services; 3. all Board of Pharmacy regulations as noted in Chapter 16.5 Board of Pharmacy; and 4. documentation requirements in a Medication Administration Record (MAR) as described in Chapter 20 20.6 Medication Administration Record (MAR) **Chapter 13 Nursing Services: 13.2 General Nursing Services Requirements and Scope of Services:** The following general requirements are applicable for all RNs and LPNs in the DD Waiver. This section represents the scope of nursing services. Refer to Chapter 10 Living Care Arrangements (LCA) for residential provider agency responsibilities related to nursing. Refer to Chapter 11.6 Customized Community Supports (CCS) for agency responsibilities related to nursing. **13.3.2.3 Medication Oversight:** Medication Oversight by a DD Waiver nurse is required in Family Living when a person lives with a non-related Family Living provider; for all JCMs; and whenever non-related DSP provide AWMD medication supports. 1. The nurse must respond to calls requesting delivery of PRN medications from AWMD trained DSP, non-related Family Living providers. 2. Family Living providers related by affinity or consanguinity (blood, adoption, or marriage) are not required to contact the nurse prior to assisting with delivery of a PRN medication.	**NA**	**New Findings:** After an analysis of the evidence it has been determined there is a significant potential for a negative outcome to occur. Based on record review, the Agency did not maintain documentation of PRN authorization as required by standard for 1 of 3 Individuals. Individual #6 August 2022 No documentation of the verbal authorization from the Agency nurse prior to each administration / assistance of PRN medication was found for the following PRN medication: • Ibuprofen 200 mg – PRN – 8/27 (given 1 time) • Loperamide 2 mg – PRN – 8/6 (given 1 time)

13.2.8.1.3 Assistance with Medication Delivery by Staff (AWMD): For people who do not meet the criteria to self-administer medications independently or with physical assistance, trained staff may assist with medication delivery if: 1. Criteria in the MAAT are met. 2. Current written consent has been obtained from the person/guardian/surrogate healthcare decision maker. 3. There is a current Primary Care Practitioner order to receive AWMD by staff. 4. Only AWMD trained staff, in good standing, may support the person with this service. 5. All AWMD trained staff must contact the on-call nurse prior to assisting with a PRN medication of any type. a. Exceptions to this process must comply with the DDSD Emergency Medication list as part of a documented MERP with evidence of DSP training to skill level.		

Standard of Care	Routine Survey Deficiencies February 21 - March 3, 2022	Verification Survey New and Repeat Deficiencies September 19 – 28, 2022
Service Domain: 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.*		
Tag # 1A08 Administrative Case File (Other Required Documents)	**Standard Level Deficiency**	**COMPLETE**
Tag # LS14 Residential Service Delivery Site Case File (ISP and Healthcare Requirements) ***(Upheld by IRF)***	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # LS14.1 Residential Service Delivery Site Case File (Other Req. Documentation) ***(Upheld by IRF)***	**Standard Level Deficiency**	**COMPLETE**
Service Domain: Qualified Providers – *The State monitors non-licensed/non-certified providers to assure adherence to waiver requirements. The State implements its policies and procedures for verifying that provider training is conducted in accordance with State requirements and the approved waiver.*		
Tag # 1A22 Agency Personnel Competency	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # 1A25.1 Caregiver Criminal History Screening	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # 1A26 Consolidated On-line Registry Employee Abuse Registry	**Standard Level Deficiency**	**COMPLETE**
Tag # 1A43.1 General Events Reporting: Individual Reporting	**Standard Level Deficiency**	**COMPLETE**
Service Domain: 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 # 1A08.2 Administrative Case File: Healthcare Requirements & Follow-up	**Standard Level Deficiency**	**COMPLETE**
Tag # 1A09.1 Medication Delivery PRN Medication Administration	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # 1A15 Healthcare Coordination - Nurse Availability / Knowledge	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # 1A15.2 Administrative Case File: Healthcare Documentation (Therap and Required Plans)	**Condition of Participation Level Deficiency**	**COMPLETE**
Tag # 1A27.2 Duty to Report IRs Filed During On-Site and/or IRs Not Reported by Provider	**Standard Level Deficiency**	**COMPLETE**
Tag # 1A50.1 Individual: Scope of Services ***(Individual Interviews)***	**Standard Level Deficiency**	**COMPLETE**
Tag # LS25 Residential Health & Safety (Supported Living / Family Living / Intensive Medical Living)	**Standard Level Deficiency**	**COMPLETE**
Service Domain: Medicaid Billing/Reimbursement – *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 # IS30 Customized Community Supports Reimbursement	**Standard Level Deficiency**	**COMPLETE**
Tag # LS26 Supported Living Reimbursement	**Standard Level Deficiency**	**COMPLETE**

	Verification Survey Plan of Correction, On-going QA/QI and Responsible Party	Completion Date
Tag # 1A09 Medication Delivery Routine Medication Administration	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)***:** → **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to affect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)***:** →	
Tag # 1A09.1.0 Medication Delivery PRN Medication Administration	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)***:** → **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to affect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)***:** →	

Tag # 1A09.2 Medication Delivery Nurse Approval for PRN Medication	**Provider:** **State your Plan of Correction for the deficiencies cited in this tag here** *(How is the deficiency going to be corrected? This can be specific to each deficiency cited or if possible an overall correction?)***: →** **Provider:** **Enter your ongoing Quality Assurance/Quality Improvement processes as it related to this tag number here** *(What is going to be done? How many individuals is this going to affect? How often will this be completed? Who is responsible? What steps will be taken if issues are found?)***: →**	

NEW MEXICO
Department of Health
Division of Health Improvement

MICHELLE LUJAN GRISHAM
Governor

DAVID R. SCRASE, M.D.
Acting Cabinet Secretary

Date: November 21, 2022

To: Emad Elmaoued, Executive Director

Provider: ADID Care, INC
Address: 5115 Copper Ave NE
State/Zip: Albuquerque, New Mexico 87108

E-mail Address: emad@adidcare.com

Region: Metro & Northeast
Routine Survey: February 21 – March 3, 2022
Verification Survey: September 19 – 28, 2022
Program Surveyed: Developmental Disabilities Waiver

Service Surveyed: Supported Living, Family Living, Customized In-Home Supports, Customized Community Supports

Survey Type: Verification

Dear Mr. Elmaoued:

The Division of Health Improvement/Quality Management Bureau has received, reviewed and approved the supporting documents you submitted for your Plan of Correction. The documents you provided verified that all previously cited survey Deficiencies have been corrected.

The Plan of Correction process is now complete.

Furthermore, your agency is now determined to be in Compliance with all Conditions of Participation.

To maintain ongoing compliance with standards and regulations, continue to use the Quality Assurance (self-auditing) processes you described in your Plan of Correction.

Consistent use of these Quality Assurance processes will enable you to identify and promptly respond to problems, enhance your service delivery, and result in fewer deficiencies cited in future QMB surveys.

Thank you for your cooperation with the Plan of Correction process, for striving to come into compliance with standards and regulations, and for helping to provide the health, safety and personal growth of the people you serve.

Sincerely,

Monica Valdez, BS

Monica Valdez, BS
Healthcare Surveyor Advanced/Plan of Correction Coordinator
Quality Management Bureau/DHI

Q.23.1.DDW.D4455.5/2.VER.09.22.325