

Division of Health Improvement

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 4014	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____	(X3) DATE SURVEY COMPLETED 01/26/2017
NAME OF PROVIDER OR SUPPLIER SUNNY DAY ASSISTED LIVING		STREET ADDRESS, CITY, STATE, ZIP CODE 205 DAY STREET GALLUP, NM 87301		
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A 000	Initial Comments The following deficiencies were cited as a result of an Initial survey completed on 01/26/17 for the state requirements of 7 NMAC 8.2, Regulations for Assisted Living Facilities.	A 000		
A 034	7 NMAC 8.2.34 Custodial Drug Permits CUSTODIAL DRUG PERMITS: A facility with two (2) or more residents that is licensed pursuant to this rule and that assists with self-administration or safeguards medications for residents shall have a current custodial drug permit issued by the state board of pharmacy. A. Procurement, labeling and storage. The facility shall provide assistance to the resident in obtaining the necessary medications, treatment and medical supplies as identified in the ISP. The facility shall procure, label and store medications for residents who require assistance with self-administration of medication in compliance with state and federal laws. (1) All medications, including non-prescription drugs, shall be stored in a locked compartment or in a locked room, as approved by the board of pharmacy and the key shall be in the care of the administrator or designee. (2) Internal medication shall be kept separate from external medications. Drugs to be taken by mouth shall be separated from all other delivery forms. (3) A separate, locked refrigerator shall be provided by the facility for medications. The refrigerator temperature shall be kept in compliance with the state board of pharmacy requirements for medications. (4) All medications, including non-prescription medications, shall be stored in separate compartments for each resident and all medications shall be labeled with the resident's	A 034		

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LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE

TITLE

(X6) DATE

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A 034	Continued From page 1 name. (5) A resident may be permitted to keep his or her own medication in a locked compartment in his or her room for self-administration, if the physician's order deems it appropriate. (6) The facility shall not require the residents to purchase medications from any particular pharmacy. (7) Medical gases (oxygen) and equipment used for the administration of inhalation therapy and for resuscitative purposes shall comply with the national fire protection association (NFPA) 99. (8) A proof of use record shall be maintained separately for each schedule II through IV drug (controlled substances). The proof of use sheet shall document: (a) the type and strength of the schedule II through IV drugs; (b) the date and time staff assisted with self-administration; (c) the resident ' s name; (d) the prescriber ' s name; (e) the dose; (f) the signature of the person assisting with delivery of the medication; and (g) the balance of medication remaining. (9) Any remaining medication discontinued by a physician ' s order, or upon discharge or death of the resident shall be inventoried and moved to a separate locked storage container. Such discontinued medications shall be destroyed upon the next quarterly visit by the consulting pharmacist in accordance with 16.19.11.10 NMAC. (10) The record of medication destruction shall be signed by the administrator or designee and the pharmacist and shall be kept on file at the facility. B. Consulting pharmacist. The facility shall	A 034		

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A 034	Continued From page 2 maintain records demonstrating that the consulting pharmacist provides the following oversight and guidance. (1) Reviews the medication regimen as needed, but at least quarterly/every three (3) months, to determine that all medications and records are accurate and current. All irregularities shall be reported to the administrator of the facility and these irregularities shall be resolved by the administrator within seventy-two (72) hours. (2) A system of records of receipt and disposition of all drugs in sufficient detail to enable an accurate reconciliation. (3) Consultation shall be provided on all aspects of pharmacy services in the facility, including reference information regarding side effects and, when needed, physician consultation in cases involving the use of psychotropic medications. (4) The consulting pharmacist will be responsible for assuring that the facility meets all requirements for storage, labeling, destruction and documentation of medications as required by the state board of pharmacy, 16.19.11.10 NMAC and 7.8.2 NMAC. [7.8.2.34 NMAC - Rp, 7.8.2.35 NMAC, 01/15/2010] This REQUIREMENT is not met as evidenced by: 7.8.2.34 A. (7) Reference NFPA 99, 1999 Edition Section 8-3.1.11.3 (Signs) Reference NFPA 99, 1999 Edition Section 4-3.1.1.2 Storage Requirements (Location, Construction, Arrangement). (a) * Nonflammable Gases (Any Quantity;	A 034		

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A 034	Continued From page 3 In-Storage, Connected, or Both) 1. Sources of heat in storage locations shall be protected or located so that cylinders or compressed gases shall not be heated to the activation point of integral safety devices. In no case shall the temperature of the cylinders exceed 130°F (54°C). Care shall be exercised when handling cylinders that have been exposed to freezing temperatures or containers that contain cryogenic liquids to prevent injury to the skin. 2. * Enclosures shall be provided for supply systems cylinder storage or manifold locations for oxidizing agents such as oxygen and nitrous oxide. Such enclosures shall be constructed of an assembly of building materials with a fire-resistive rating of at least 1 hour and shall not communicate directly with anesthetizing locations. Other nonflammable (inert) medical gases may be stored in the enclosure. Flammable gases shall not be stored with oxidizing agents. Storage of full or empty cylinders is permitted. Such enclosures shall serve no other purpose. 3. Provisions shall be made for racks or fastenings to protect cylinders from accidental damage or dislocation. 4. The electric installation in storage locations or manifold enclosures for nonflammable medical gases shall comply with the standards of NFPA 70, National Electrical Code, for ordinary locations. Electric wall fixtures, switches, and receptacles shall be installed in fixed locations not less than 152 cm (5 ft) above the floor as a precaution against their physical damage. 5. Storage locations for oxygen and nitrous oxide shall be kept free of flammable materials [see also 4-3.1.1.2(a)7]. 6. Cylinders containing compressed gases and containers for volatile liquids shall be kept	A 034		

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A 034	Continued From page 4 away from radiators, steam piping, and like sources of heat. 7. Combustible materials, such as paper, cardboard, plastics, and fabrics, shall not be stored or kept near supply system cylinders or manifolds containing oxygen or nitrous oxide. Racks for cylinder storage shall be permitted to be of wooden construction. Wrappers shall be removed prior to storage. Exception: Shipping crates or storage cartons for cylinders. 8. When cylinder valve protection caps are supplied, they shall be secured tightly in place unless the cylinder is connected for use. 9. Containers shall not be stored in a tightly closed space such as a closet [see 8-2.1.2.3(c)]. Reference NFPA 99, 1999 Edition Section 8-3.1.11.3 (Signs) Based on observation and interview, the facility failed to ensure that: 1. Oxygen cylinder tanks were stored correctly, securely and protected from accidental damage or dislocation. 2. Oxygen warning signs were posted outside the room where oxygen is being used. These deficient practices may cause injury or death to all 27 (R #s 1-27) residents identified on the resident census provided by the Administrator on 01/24/17, if an oxygen cylinder tank were to fall over damaging the valve and causing compressed oxygen to burst out and if there are no warning signs posted informing there is "oxygen in use" in the event there is a fire. The findings are: A. On 01/25/17 at 11:05 am, during a tour of the facility an observation of R# 1's room revealed that three (3) oxygen cylinder tanks were stored	A 034		

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A 034	Continued From page 5 in the closet and no "oxygen in use" sign was posted on the door. B. On 1/25/17 at 11:15 am, during an interview with the Administrator, he confirmed that three (3) oxygen cylinder tanks were stored in R# 1's room closet and no "oxygen in use" sign was posted on the door.	A 034			
A 035	7 NMAC 8.2.35 Medication MEDICATIONS: Administration of medications or staff assistance with self-administration of medications shall be in accordance with state and federal laws. No medications, including over-the-counter medications, PRN (when needed) medications, or treatment shall be started, changed or discontinued by the facility without an order from the physician, physician assistant or nurse practitioner and with entry into the resident's record. A. State board of nursing licensed or certified health care professionals are responsible for the administration of medications. Administration may only be performed by these individuals. B. Facility staff may assist a resident with the self-administration of medications if written consent by the resident is given to the administrator of the facility or the administrator's designee. If the resident is incapable of giving consent, the surrogate decision maker named in accordance with New Mexico law may give written consent for assistance with self-administration of medications. All staff that assist with self-administration of medications shall have successfully completed a state approved assistance with self-administration of medication training program or be licensed or certified by the state board of nursing.	A 035			

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A 035	Continued From page 6 C. PRN (pro re nada) medication. (1) Physician or physician extender 's orders for PRN medications shall clearly indicate the circumstances in which they are to be used, the number of doses that may be given in a 24-hour period and indicate under what circumstances the primary care practitioner (PCP) is to be notified. (2) The utilization of PRN medications shall be reviewed routinely. Frequent or escalating use of PRN medications shall be reported to the PCP. D. Only a licensed nurse (RN or LPN) shall administer any medications or conduct any invasive procedures provided by the following routes: intravenous (IV), subcutaneous (SQ), intramuscular (IM), vaginal or rectal. Only a licensed nurse shall administer non-premixed nebulizer treatments. E. The facility shall have medication reference material that contains information relating to drug interactions and side effects on the premises. Staff that assist in the self-administration of medications shall know interactions or possible side effects that might occur. F. Medications prescribed for one resident shall not be used for another resident. G. Medication assistance record (MAR). For residents who are not independent and require assistance with self administration, the facility shall have a MAR that documents the details of the residents' medication, including PRN and over-the-counter medication that is assisted with self-administration by qualified staff or administered to the resident by licensed or certified staff. The information in the MAR shall include: (1) the resident's name; (2) any known allergies to medication that the resident has; (3) the name of the resident's PCP or the	A 035		

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A 035	Continued From page 7 prescriber of the medication; (4) the diagnosis or reason for the medication; (5) the name of the medication, including the drug product brand name and the generic name; (6) notation if the medication is a schedule II-IV drug; (7) the dosage of the medication; (8) the strength of the medication; (9) the frequency or how often the medication is to be taken or given; (10) the route of delivery for the medication (mouth, eye, ear, other); (11) the method of delivery for the medication (pills, drops, IM injection, other); (12) the date that the medication was started or discontinued; (13) any change in the medication order; (14) pre-medication information (i.e., pulse, respiration, blood pressure, blood sugar) as required by the medication order; (15) the date and time that the medication is self-administered, administered with assistance or is administered; (16) the initials and signature of the person assisting with or administering the medication; (17) the desired results obtained from or problems encountered with the medication (pain relieved, allergic reaction, etc.); (18) any refused dose of medication; (19) any missed dose of medication; and (20) any medication error. H. No medication shall be stopped or started without specific orders from the primary care physician. I. If a resident refuses to take a prescribed medication, it shall be documented and the facility shall report it to the prescriber. J. A suspected adverse reaction to a medication shall be documented on the MAR and reported	A 035		

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A 035	Continued From page 8 immediately to the PCP and the resident's surrogate decision maker. If applicable, emergency medical treatment shall be arranged. Documentation of the event shall be kept in the resident's record. K. Prescription medication, other than blister packs and unit dose containers, shall be kept in the original container with a pharmacy label that includes the following: (1) the resident's name; (2) the name of the medication; (3) the date that the prescription was issued; (4) the prescribed dosage and the instructions for administration of the medication; and (5) the name and title of the prescriber. L. Any medication that is removed from the pharmacy container or blister pack shall be given immediately and documented by the staff that assisted with the medication delivery. M. The facility shall report all medication errors to the physician, documentation of medication errors and the prescriber's response shall be kept in the resident's record. N. The facility shall develop and follow a written policy for unused, outdated, or recalled medications kept in the facility in accordance with 16.19.11.10 NMAC (AS AMENDED). [7.8.2.35 NMAC - Rp, 7.8.2.35 NMAC, 01/15/2010] This REQUIREMENT is not met as evidenced by: 7.8.2.35 G (5) (12) Based on record review and interview, the facility failed to ensure that the Medication Administration Record (MAR) for 4 (R #s 1-4) of 4	A 035		

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A 035	Continued From page 9 (R #s 1-4) residents who were reviewed for medication accuracy had complete and accurate MARs. If the information on the MAR is not complete, accurate, and includes all required information, then medication errors could occur resulting in residents being harmed or require medical treatment. The findings are: A. Record review of R #1's MARs dated December 2016 and January 2017 revealed they did not contain: 1. The generic name for the medication. 2. The date that the medication was started or discontinued. B. Record review of R #2's MARs dated December 2016 and January 2017 revealed they did not contain: 1. The generic name for the medication. 2. The date that the medication was started or discontinued. C. Record review of R #3's MARs dated December 2016 and January 2017 revealed they did not contain: 1. The generic name for the medication. 2. The date that the medication was started or discontinued. D. Record review of R #4's MARs dated December 2016 and January 2017 revealed they did not contain: 1. The generic name for the medication. 2. The date that the medication was started or discontinued. E. On 01/25/17 at 2:40 pm, during an interview with the Registered Nurse, confirmed the missing data from R #s 1 - 4's MARs.	A 035		

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A 037	Continued From page 10	A 037		
A 037	7 NMAC 8.2.37 Laundry Services LAUNDRY SERVICES: A. General requirements. The facility shall provide laundry services for the residents, either on the premises or through a commercial laundry and linen service. (1) On-site laundry facilities shall be located in areas separate from the resident units and shall be provided with necessary washing and drying equipment. (2) Soiled laundry shall be kept separate from clean laundry, unless the laundry facility is provided for resident use only. (3) Staff shall handle, store, process and transport linens with care to prevent the spread of infectious and communicable disease. (4) Soiled laundry shall not be stored in the kitchen or dining areas. The building design and layout shall ensure the separation of laundry room from kitchen and dining areas. An exterior route to the laundry room is not an acceptable alternative, unless it is completely enclosed. (5) In new construction or newly licensed facilities with more than fifteen (15) residents, washers shall be in separate rooms from dryers. The rooms with washers shall have negative air pressure from the other facility rooms. (6) All linens shall be changed as needed and at least weekly or when a new resident is to occupy the bed. (7) The mattress pad, blankets and bedspread shall be laundered as needed and at least once per month or when a new resident is to occupy the bed. (8) Bath linens consisting of hand towel, bath towel and washcloth shall be changed as needed and at least weekly. (9) There shall be a clean, dry, well ventilated	A 037		

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A 037	Continued From page 11 storage area provided for clean linen. (10) Facility laundry supplies and cleaning supplies shall not be kept in the same storage areas used for the storage of foods and clean storage and shall be kept in a secured room or cabinet. B. Residents may do their own laundry, if it is their preference and they are capable of doing so, or if it is part of their skill-building for independent living and is documented as part of their ISP. [7.8.2.37 NMAC - Rp, 7.8.2.39 NMAC, 01/15/2010] This REQUIREMENT is not met as evidenced by: 7.8.2.37 A (9) Based on observation and interview the facility failed to ensure that the clean linens were stored in a well ventilated closet. This deficient practice has the potential to affect all 27 (R #1-27) residents identified on the resident census list provided by the Administrator on 01/24/17 to become ill from bacteria contaminated linens if the clean linens are not stored in a well ventilated closet. The findings are: A. On 01/24/17 at 2:45 pm, during observation, the clean linen closet was observed to not be ventilated. B. On 01/24/17 at 2:50 pm, during interview with the Administrator, he confirmed that the linen closet was not ventilated.	A 037		