

DEPARTMENT OF HEALTH AND HUMAN SERVICES
CENTERS FOR MEDICARE & MEDICAID SERVICES

PRINTED: 10/07/2013
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 09G027	(X2) MULTIPLE CONSTRUCTION A. BUILDING _____ B. WING _____		(X3) DATE SURVEY COMPLETED 09/27/2013
NAME OF PROVIDER OR SUPPLIER MY OWN PLACE			STREET ADDRESS, CITY, STATE, ZIP CODE 3215 20TH STREET, NE WASHINGTON, DC 20018		
(X4) ID PREFIX TAG	SUMMARY STATEMENT OF DEFICIENCIES (EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)	ID PREFIX TAG	PROVIDER'S PLAN OF CORRECTION (EACH CORRECTIVE ACTION SHOULD BE CROSS-REFERENCED TO THE APPROPRIATE DEFICIENCY)	(X5) COMPLETION DATE	
W 000	INITIAL COMMENTS A recertification survey was conducted from September 24, 2013 to September 27, 2013. A sample of two clients was selected from a population of four females with varying degrees of intellectual disabilities. This survey was initiated utilizing the fundamental survey process. The findings of the survey were based on observations in the home and one day program, interviews with direct support staff, nursing and administrative staff, as well as a review of client and administrative records, including incident reports. [Qualified mental retardation professional (QMRP) will be referred to as qualified intellectual disabilities professional (QIDP) within this report.]	W 000			
W 104	483.410(a)(1) GOVERNING BODY The governing body must exercise general policy, budget, and operating direction over the facility. This STANDARD is not met as evidenced by: Based on interview and record review, the governing body failed to have an effective system to ensure the timely procurement of a medication that required prior authorization, for one of two clients in the sample. (Client #1) The finding includes: On September 24, 2013, at 7:56 p.m., trained medication employee (TME) #1 was observed to administer Client #1 Calcium W/V 600 mg/400 tablet by mouth.	W 104			

Received 10/18/13
Department of Health
Health Regulation & Licensing Administration
Intermediate Care Facilities Division
899 North Capitol St., N.E.
Washington, D.C. 20002

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

TITLE

(X6) DATE

Any deficiency statement ending with an asterisk (*) denotes a deficiency which the institution may be excused from correcting providing it is determined that other safeguards provide sufficient protection to the patients. (See instructions.) Except for nursing homes, the findings stated above are disclosable 90 days following the date of survey whether or not a plan of correction is provided. For nursing homes, the above findings and plans of correction are disclosable 14 days following the date these documents are made available to the facility. If deficiencies are cited, an approved plan of correction is requisite to continued program participation.

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W 104	Continued From page 1 Interview with TME #1 on September 24, 2013, at 8:03 p.m. revealed that Client #1 was prescribed to receive Calcium W/V 600 mg/400 tablet in the morning and evening. Additionally, the client was prescribed to receive several other medications in the morning. On September 24, 2013, at 8:29 p.m., verification of the medication administration record (MAR) and the corresponding physician's orders, confirmed Client #1 was prescribed Calcium W/V 600 mg/400 tablet in the morning and evening for bone health. The MAR and the physician's revealed the client was prescribed Tyzeka F/C 600 mg tablet, 1 tablet by mouth every morning for chronic hepatitis B. Further review of the MAR revealed however, that Client #1 did not receive Tyzeka F/C 600 mg tablet, on September 14, and September 15, 2013 because it was not available. Interview with primary registered nurse (RN) #1 on September 27, 2013 at 11:40 a.m., confirmed that Client #1 did not receive the Tyzeka F/C 600 mg tablet medication as prescribed on the aforementioned dates, because the pharmacy did not deliver it on time. Primary RN #1 indicated that although the facility made efforts to correct the problem in the past, it remained a concern. [Note: The MAR revealed that the client also did not receive this medication on May 12, 13, 14, and 15, 2013 because it was not available.] On September 27, 2013, at 11:51 p.m., interview with the director of nursing (DON) revealed that in both instances the pharmacy reported that the delivery of the Tyzeka F/C 600 mg tablets was delayed because prior approval was not obtained	W 104	W104 Governing Body The QIDP in conjunction with the RN will ensure that an effective system are in place for the timely procurement of medications that requires prior authorization before the release of the medication by the facility by establishing a written protocol by 11/01/13. The RN will review and monitor medication on a monthly basis to ensure compliance.	11/01/13 Ongoing	

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W 104	Continued From page 2 to renew the prescriptions. According to the DON, the pharmacy did not notify the agency of the expiration dates of authorizations to purchase the medication. The DON revealed that this notification would alert the agency to request a timely updated authorization from the funding source, to ensure the client received the medication as prescribed. The DON indicated that the agency is continuing to investigate the problem in order to develop an effective system to ensure timely notification when prior authorization for medication is required. At the time of the survey, there was no evidence an effective system had been implemented to ensure the timely provision of each medication requiring prior authorization before the release of the medication to the facility.	W 104			
W 382	483.460(l)(2) DRUG STORAGE AND RECORDKEEPING The facility must keep all drugs and biologicals locked except when being prepared for administration. This STANDARD is not met as evidenced by: Based on observation and interview, the facility failed to ensure that all drugs and biological were kept locked, except when being prepared for administration for three of four clients in the sample (Clients #1, #3, and #4). The finding includes: On September 24, 2013, at 7:10 p.m., trained medication employee (TME) #1 announced to the surveyor that she was ready to administer the	W 382	W382 Drug Storage and Recordkeeping The QIDP/Program Manager in conjunction with the RN will ensure that all drugs and biological are kept locked, except when being prepared for administration by in-service all staff on medication management by 11/01/13. The RN will continue to observe staff on a quarterly basis to ensure compliance. The staff involved has received additional training on safe guarding medications during the administration process. In addition the staff involved will be monitored during the delivery of medication for a period of once a week, 3 consecutive weeks until they demonstrate error free sessions.	10/15/13 Ongoing	

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W 382	Continued From page 3 clients' evening medications. At 7:27 p.m., TME #1 was observed to administer Client #4 medications that included: Atorvastin Calcium 80 milligrams (mg.) tablet; Calcium w/v 600 mg.- 400 mg. tablet; Carbamazepine 200 mg. tablet; Folic Acid 0.4 mg. tablet, 2 tablets (.8 mg.); Trihexyphenidyl HCl 2 mg. tablet; and Zetia U-D 10 mg. tablet. At 7:35 p.m., TME #1 left the medication closet open and escorted Client #4 back to the living room. After the clients' medications were administered, the closet remained open while the TME escorted the clients from the medication area to the living room. Interview with TME #1 on September 24, 2013, at 8:29 p.m., and with primary registered nurse (RN) #1 on September 25, 2013, at 2:37 p.m., confirmed that the medication closet should be locked at all time except when medications are being prepared for administration.	W 382	

Health Regulation & Licensing Administration

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION	(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: HFD03-0238	(X2) MULTIPLE CONSTRUCTION A. BUILDING: _____ B. WING: _____	(X3) DATE SURVEY COMPLETED 09/27/2013
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1 000 INITIAL COMMENTS

1 000

A licensure survey was conducted from September 24, 2013 to September 27, 2013. A sample of two residents was selected from a population of four females with varying degrees of intellectual disabilities.

The findings of the survey were based on observations in the home and one day programs, interviews with direct support staff, nursing and administrative staff, as well as a review of resident and administrative records, including incident reports.

[Qualified mental retardation professional (QMRP) will be referred to as qualified intellectual disabilities professional (QIDP) within this report.]

1 075 3503.3(d) BEDROOMS AND BATHROOMS

1 075

Each bedroom shall be equipped with at least the following items for each resident:

(d) Night stand.

This Statute is not met as evidenced by: Based on observation and interview, the group home for individuals with intellectual disabilities (GHIID) failed to ensure that each bedroom was equipped with a nightstand for each resident in the bedroom of two residents. (Residents #1 and #3)

The finding includes:

Observation of the bedroom of Residents #1 and #3 on September 26, 2013, at 3:47 p.m., revealed one nightstand placed between the beds of the

11/01/13

Ongoing

1075 Bedrooms and Bathrooms

Each bedroom shall be equipped with at least the following items

(d) Night stand

1. The QIDP/Program Manager will ordered a night stand. The QIDP/Program Manager will monitor residence weekly using the Environmental Compliance Form to ensure that all furnishings are in good working order and repaired/replaced as needed.

Health Regulation & Licensing Administration

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

TITLE

(X6) DATE

STATE FORM

6899

MHLR11

If continuation sheet 1 of 5

Director Health Services 10-17-13

Health Regulation & Licensing Administration

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I 075	Continued From page 1 residents. Interview with facility's program director (PD) #1 during the observation on September 26, 2013, at 3:37 p.m., confirmed there was only one night stand for the bedroom. At the time of the survey, the facility failed to ensure that a nightstand was provided as required for the use of each client in their bedroom. [This is a repeat deficiency.]	I 075		
I 090	3504.1 HOUSEKEEPING The interior and exterior of each GHMRP shall be maintained in a safe, clean, orderly, attractive, and sanitary manner and be free of accumulations of dirt, rubbish, and objectionable odors. This Statute is not met as evidenced by: Based on observation, and interview, the group home for individuals with intellectual disabilities (GHIID) failed to maintain the environment in accordance with the needs of four of four residents of the facility. (Residents #1, #2, #3, and #4. The findings include: On September 26, 2013, at 3:02 p.m., the facility's coordinator (FC) #1 accompanied the surveyor to conduct an inspection of the environment. The following concerns were identified:	I 090		

If continuation sheet 3 of 5

If continuation sheet 4 of 5

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[illegible]