

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

PRINTED: 06/09/2025
FORM APPROVED
OMB NO. 0938-0391

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTION		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 155077	(X2) MULTIPLE CONSTRUCTION A. BUILDING 01 B. WING _____		(X3) DATE SURVEY COMPLETED R 06/03/2025
NAME OF PROVIDER OR SUPPLIER ENVIVE OF INDIANAPOLIS			STREET ADDRESS, CITY, STATE, ZIP CODE 45 BEACHWAY DR INDIANAPOLIS, IN 46224		
(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
{E 000}	Initial Comments A Post Survey Revisit (PSR) to the Emergency Preparedness Survey conducted on 03/05/25 was conducted by the Indiana Department of Health in accordance with 42 CFR 483.73. Survey Date: 06/03/25 Facility Number: 000032 Provider Number: 155077 AIM Number: 100273330 At this PSR survey to the Emergency Preparedness survey, Envive of Indianapolis was found in compliance with Emergency Preparedness Requirements for Medicare and Medicaid Participating Providers and Suppliers, 42 CFR 483.73. The facility has 184 certified beds. At the time of the survey, the census was 104.	{E 000}			
{K 000}	Quality Review completed on 06/05/25 INITIAL COMMENTS A Post Survey Revisit (PSR) to the Life Safety Code Recertification and State Licensure Survey conducted on 03/05/25 was conducted by the Indiana Department of Health in accordance with 42 CFR 483.90(a). Survey Date: 06/03/25 Facility Number: 000032 Provider Number: 155077 AIM Number: 100273330 At this PSR survey, Envive of Indianapolis was	{K 000}			

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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{K 000}	Continued From page 1 found in compliance with Requirements for Participation in Medicare/Medicaid, 42 CFR Subpart 483.90(a), Life Safety from Fire and the 2012 Edition of the National Fire Protection Association (NFPA) 101, Life Safety Code (LSC), Chapter 19, Existing Health Care Occupancies and 410 IAC 16.2. This one story facility was determined to be of Type III (211) construction and was fully sprinklered. The facility has a fire alarm system with smoke detection in the corridors, in all areas open to the corridor and in rooms 11 through 19 in the C Wing. The facility has battery operated smoke detectors in all other resident sleeping rooms. The facility has a capacity of 184 and had a census of 104 at the time of this survey. All areas where residents have customary access were sprinklered. The facility has four detached buildings providing storage services and one detached building housing an emergency generator which were each not sprinklered.	{K 000}			
{K 363} SS=E	Quality Review completed on 06/05/25 Corridor - Doors CFR(s): NFPA 101 Corridor - Doors Doors protecting corridor openings in other than required enclosures of vertical openings, exits, or hazardous areas resist the passage of smoke and are made of 1 3/4 inch solid-bonded core wood or other material capable of resisting fire for at least 20 minutes. Doors in fully sprinklered smoke compartments are only required to resist the passage of smoke. Corridor doors and doors to rooms containing flammable or combustible	{K 363}			

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{K 363}	Continued From page 2 materials have positive latching hardware. Roller latches are prohibited by CMS regulation. These requirements do not apply to auxiliary spaces that do not contain flammable or combustible material. Clearance between bottom of door and floor covering is not exceeding 1 inch. Powered doors complying with 7.2.1.9 are permissible if provided with a device capable of keeping the door closed when a force of 5 lbf is applied. There is no impediment to the closing of the doors. Hold open devices that release when the door is pushed or pulled are permitted. Nonrated protective plates of unlimited height are permitted. Dutch doors meeting 19.3.6.3.6 are permitted. Door frames shall be labeled and made of steel or other materials in compliance with 8.3, unless the smoke compartment is sprinklered. Fixed fire window assemblies are allowed per 8.3. In sprinklered compartments there are no restrictions in area or fire resistance of glass or frames in window assemblies. 19.3.6.3, 42 CFR Parts 403, 418, 460, 482, 483, and 485 Show in REMARKS details of doors such as fire protection ratings, automatics closing devices, etc. This REQUIREMENT is not met as evidenced by:	{K 363}			
{K 921} SS=F	Electrical Equipment - Testing and Maintenance CFR(s): NFPA 101 Electrical Equipment - Testing and Maintenance Requirements The physical integrity, resistance, leakage current, and touch current tests for fixed and	{K 921}	TEMPORARY WAIVER APPROVED UNTIL 10/31/25		

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{K 921}	Continued From page 3 portable patient-care related electrical equipment (PCREE) is performed as required in 10.3. Testing intervals are established with policies and protocols. All PCREE used in patient care rooms is tested in accordance with 10.3.5.4 or 10.3.6 before being put into service and after any repair or modification. Any system consisting of several electrical appliances demonstrates compliance with NFPA 99 as a complete system. Service manuals, instructions, and procedures provided by the manufacturer include information as required by 10.5.3.1.1 and are considered in the development of a program for electrical equipment maintenance. Electrical equipment instructions and maintenance manuals are readily available, and safety labels and condensed operating instructions on the appliance are legible. A record of electrical equipment tests, repairs, and modifications is maintained for a period of time to demonstrate compliance in accordance with the facility's policy. Personnel responsible for the testing, maintenance and use of electrical appliances receive continuous training. 10.3, 10.5.2.1, 10.5.2.1.2, 10.5.2.5, 10.5.3, 10.5.6, 10.5.8 This REQUIREMENT is not met as evidenced by:	{K 921}	TEMPORARY WAIVER APPROVED UNTIL 10/31/25		