

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

STATEMENT OF DEFICIENCIES AND PLAN OF CORRECTIONS		(X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER: 15C0001102	(X2) MULTIPLE CONSTRUCTION A. BUILDING B. WING	(X3) DATE SURVEY COMPLETED 08/23/2023
NAME OF PROVIDER OR SUPPLIER VISION SURGICAL CENTER AT SPRINGHILL INC		STREET ADDRESS, CITY, STATE, ZIP CODE 302 W 14TH ST STE 100 B , JEFFERSONVILLE, Indiana, 47130		
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Q0000	INITIAL COMMENTS This visit was for a recertification survey of an Ambulatory Surgery Center. Facility Number: 002769 Survey Date: 08/14/2023 and 8/23/23 QA: 8/21/23	Q0000		
Q0100	ENVIRONMENT CFR(s): 416.44 The ASC must have a safe and sanitary environment, properly constructed, equipped, and maintained to protect the health and safety of patients. This CONDITION is NOT MET as evidenced by: Based on record review and interview, the facility failed to ensure the documentation for the annual testing of all devices connected to 1 of 1 fire alarm system was complete and accurate, failed to ensure 1 of 1 fire alarm system was maintained in accordance with 9.6.1.3 and failed to provide documentation for the testing of 1 of 1 Emergency Power Standby System in accordance with NFPA 110, Standard for Emergency and Standby Power Systems, Section 8.4.9, as required by NFPA 99 Health Care Facilities Code, Section 6.4.1.1.6.1... The cumulative effect of these systemic problems limited the facilities ability to ensure it provided care in a safe environment.	Q0100		
Q0104	SAFETY FROM FIRE CFR(s): 416.44(b)(1)-(3) (b) Standard: Safety from fire. (1) Except as otherwise	Q0104		

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 reverse for further 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.

LABORATORY DIRECTOR'S OR PROVIDER/SUPPLIER REPRESENTATIVE'S SIGNATURE	TITLE	(X6) DATE
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Q0104	<p>Continued from page 1</p> <p>provided in this section, the ASC must meet the provisions applicable to Ambulatory Health Care Occupancies, regardless of the number of patients served, and must proceed in accordance with the Life Safety Code (NFPA 101 and Tentative Interim Amendments TIA 12-1, TIA 12-2, TIA 12-3, and TIA 12-4).</p> <p>(2) In consideration of a recommendation by the State survey agency or Accrediting Organization or at the discretion of the Secretary, may waive, for periods deemed appropriate, specific provisions of the Life Safety Code, which would result in unreasonable hardship upon an ASC, but only if the waiver will not adversely affect the health and safety of the patients.</p> <p>(3) The provisions of the Life Safety Code do not apply in a State if CMS finds that a fire and safety code imposed by State law adequately protects patients in an ASC.</p> <p>This STANDARD is NOT MET as evidenced by:</p> <p>1. Based on record review and interview, the facility failed to ensure the documentation for the annual testing of all devices connected to 1 of 1 fire alarm system was complete and accurate. NFPA 72, National Fire Alarm Code, the 2010 Edition, at 14.6.2.4 requires a record of all inspections, testing, and maintenance shall be provided that includes the following information regarding tests and all the applicable information requested in Figure 14.6.2.4:</p> <p>(1) Date</p> <p>(2) Test frequency</p> <p>(3) Name of property</p> <p>(4) Address</p> <p>(5) Name of person performing inspection, maintenance, tests, or combination thereof, and affiliation, business address, and telephone number</p> <p>(6) Name, address, and representative of approving agency (ies)</p> <p>(7) Designation of the detector(s) tested</p> <p>(8) Functional test of detectors</p> <p>(9)*Functional test of required sequence of operations</p>	Q0104		

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Q0104	<p>Continued from page 2</p> <p>(10) Check of all smoke detectors</p> <p>(11) Loop resistance for all fixed-temperature, line-type heat detectors</p> <p>(12) Functional test of mass notification system control units</p> <p>(13) Functional test of signal transmission to mass notification systems</p> <p>(14) Functional test of ability of mass notification system to silence fire alarm notification appliances</p> <p>(15) Tests of intelligibility of mass notification system speakers</p> <p>(16) Other tests as required by the equipment manufacturer's published instructions</p> <p>(17) Other tests as required by the authority having jurisdiction</p> <p>(18) Signatures of tester and approved authority representative</p> <p>(19) Disposition of problems identified during test (e.g., system owner notified, problem corrected/successfully retested, device abandoned in place)</p> <p>This deficient practice could affect all occupants in the facility.</p> <p>2. Based on record review and interview, the facility failed to ensure 1 of 1 fire alarm system was maintained in accordance with 9.6.1.3. LSC 9.6.1.3 requires a fire alarm system to be installed, tested, and maintained in accordance with NFPA 70, National Electrical Code and NFPA 72, National Fire Alarm Code. NFPA 72, 7-3.2 requires testing shall be performed in accordance with the Table 14.4.5 Testing Frequencies. NFPA 72, 14.4.5.3.1 states sensitivity shall be checked within 1 year after installation. NFPA 72, 14.4.5.3.2 states sensitivity shall be checked every alternate year thereafter unless otherwise permitted by compliance with 14.4.5.3.3. NFPA 72, 14.4.5.3.5 states smoke detectors or smoke alarms found to have a sensitivity outside the listed and marked sensitivity range shall be cleaned and recalibrated or be replaced.</p> <p>Findings include:</p>	Q0104		

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Q0104	<p>Continued from page 3</p> <p>#1 Based on record review on 08/23/23 between 9:30 a.m. and 2:15 p.m. with the DON and Assistant DON present, the most recent four quarterly fire alarm system inspection/testing reports, which also included the annual fire alarm system inspection/testing report dated 08/18/22, 11/22/22, 02/23/23 and 05/23/23, only included the visual inspection and functional testing of 21 of the facility's 25 smoke detectors. The smoke detectors in the two OR's and two Post Op smoke detectors were not included as being inspected/tested during any of the four most recent fire alarm system inspections. This was acknowledged by the DON and Assistant DON at the time of record review.</p> <p>This finding was reviewed with the DON and Assistant DON during the exit conference.</p> <p>Findings include:</p> <p>#2 Based on record review on 08/23/23 between 9:30 a.m. and 2:15 p.m. with the DON and Assistant DON present, there was no smoke detector sensitivity test available for review for 25 of 25 smoke detectors. Based on interview at the time of record review, the DON confirmed no documentation for smoke detector sensitivity was available for review.</p> <p>This finding was reviewed with the DON and Assistant DON during the exit conference.</p>	Q0104		
Q0109	<p>EMERGENCY EQUIPMENT</p> <p>CFR(s): 416.44(d)</p> <p>(d) Standard: Emergency equipment. The ASC medical staff and governing body of the ASC coordinates, develops, and revises ASC policies and procedures to specify the types of emergency equipment required for use in the ASC's operating room. The equipment must meet the following requirements:</p> <p>(1) Be immediately available for use during emergency situations.</p> <p>(2) Be appropriate for the facility's patient population.</p> <p>(3) Be maintained by appropriate personnel.</p> <p>This STANDARD is NOT MET as evidenced by:</p>	Q0109		

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Q0109	<p>Continued from page 4</p> <p>Based on record review and interview, the facility failed to provide documentation for the testing of 1 of 1 Emergency Power Standby System in accordance with NFPA 110, Standard for Emergency and Standby Power Systems, Section 8.4.9, as required by NFPA 99 Health Care Facilities Code, Section 6.4.1.1.6.1. NFPA 110 Section 8.4.9 states that all Level 1 Emergency Power Systems shall be tested at least once within every three years. Where the assigned class is greater than 4 hours, it shall be permitted to terminate the test after 4 hours. NFPA 99 Section 6.4.1.1.6.1 states that Type 1 and Type 2 essential electrical system power sources shall be classified at Type 10, Class X, Level 1 generator sets. This deficient practice could affect all occupants in the facility.</p> <p>Findings include:</p> <p>Based on record review on 08/23/23 between 9:30 a.m. and 2:15 p.m. with the DON and Assistant DON present, the facility could not provide documentation of a four hour test of the emergency generator within the past 36 months for the emergency generator. This was confirmed by the DON at the time of record review.</p> <p>This finding was reviewed with the DON and Assistant DON during the exit conference.</p>	Q0109			