DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION

DISTRICT ADDRESS AND PHONE NUMBER	DATE(S) OF INSPECTION
550 W. Jackson Blvd., Suite 800 Chicago, IL 60661-4716 (312)353-5863 Fax:(312)596-4187	2/17/2026-2/20/2026
	FEI NUMBER 3009542956

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED
Rajsekhar Jayaraman, Owner

FIRM NAME	STREET ADDRESS
BURTON MEDICAL	2300 W Windsor Ct Ste C
CITY, STATE, ZIP CODE, COUNTRY	TYPE ESTABLISHMENT INSPECTED
Addison, IL 60101-1491	Manufacturer

This document lists observations made by the FDA representative(s) during the inspection of your facility. They are inspectional observations, and do not represent a final Agency determination regarding your compliance. If you have an objection regarding an observation, or have implemented, or plan to implement, corrective action in response to an observation, you may discuss the objection or action with the FDA representative(s) during the inspection or submit this information to FDA at the address above. If you have any questions, please contact FDA at the phone number and address above.

DURING AN INSPECTION OF YOUR FIRM I OBSERVED:

OBSERVATION 1

The information required by the Global Unique Device Identification Database for each medical device version or model required to bear a unique device identifier (UDI) was not provided.

Specifically, your firm has not registered the APEX LED device with the Global Unique Device Identification Database.

Since January 1st, 2024, your firm has manufactured [redacted] (b) (4) Class II APEX LED, a mounted surgical ceiling light, intended to provide illumination for surgeries, procedures, emergency rooms, trauma centers, and other health care facilities.

SEE REVERSE OF THIS PAGE

EMPLOYEE(S) SIGNATURE
Hammam A Abdallah, Investigator

X Hammam A Abdallah
Investigator
Signed By: Hammam A. Abdallah
-5
Date Signed: 02-20-2026
16:39:47

DATE ISSUED
2/20/2026

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Annotations to Observations

Observation 1: Promised to correct

SEE REVERSE OF THIS PAGE	**EMPLOYEE(S) SIGNATURE** Hammam A Abdallah, Investigator	X Hammam A Abdallah Investigator Signed By: Hammam A. Abdallah Date Signed: 02-20-2026 16:39:47	**DATE ISSUED** 2/20/2026

The observations of objectionable conditions and practices listed on the front of this form are reported:

1. Pursuant to Section 704(b) of the Federal Food, Drug and Cosmetic Act, or
2. To assist firms inspected in complying with the Acts and regulations enforced by the Food and Drug Administration.

Section 704(b) of the Federal Food, Drug, and Cosmetic Act (21 USC 374(b)) provides:

"Upon completion of any such inspection of a factory, warehouse, consulting laboratory, or other establishment, and prior to leaving the premises, the officer or employee making the inspection shall give to the owner, operator, or agent in charge a report in writing setting forth any conditions or practices observed by him which, in his judgment, indicate that any food, drug, device, or cosmetic in such establishment (1) consists in whole or in part of any filthy, putrid, or decomposed substance, or (2) has been prepared, packed, or held under insanitary conditions whereby it may have become contaminated with filth, or whereby it may have been rendered injurious to health. A copy of such report shall be sent promptly to the Secretary."