K230669 L-Cath™ Single and Dual Lumen Catheters, L-Cath™ Midline Catheters

CLEARED

Nov 30, 2023
265 days to decision

K230669 · Product code: **LJS** · General Hospital
Source: https://www.510kdatabase.net/k230669/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Mar 10, 2023
Decision date	Nov 30, 2023
Days to decision	265 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Argon Medical Devices**
Location	Athens, TX, US
Contact	Scott Bishop
510(k) history	5 submissions · 5 cleared · 2021-2023

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k230669/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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