

K973639 ELECTRODE SEMI-FLOTATION CATHETER BY J-LLOYD MEDICAL, INC.Apr 20, 1998
208 days to decisionK973639 · Product code: DRF · Cardiovascular
Source: <https://www.510kdatabase.net/k973639/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Sep 24, 1997
Decision date	Apr 20, 1998
Days to decision	208 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	J-Lloyd Medical, Inc.
Location	West Berlin, NJ, US
Contact	JAMES L SKAGGS
510(k) history	16 submissions · 16 cleared · 1993-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k973639/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 4, 2026