FDA 510(k) Premarket Notification

K973639 ELECTRODE SEMI-FLOTATION CATHETER BY J-LLOYD MEDICAL, INC.

CLEARED

Apr 20, 1998
208 days to decision

K973639 · 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