

K993698 SLIMCATH-DX MODELS, 3SUXXXX, 3SBXXXX, AND FCXXXXAug 16, 2000
288 days to decisionK993698 · Product code: **DRF** · Cardiovascular
Source: <https://www.510kdatabase.net/k993698/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Nov 2, 1999
Decision date	Aug 16, 2000
Days to decision	288 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Mogul Enterprises
Location	San Jose, CA, US
Contact	JAMIL MOGUL
510(k) history	3 submissions · 3 cleared · 2000-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k993698/> 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 3, 2026