FDA 510(k) Premarket Notification

K993698 SLIMCATH-DX MODELS, 3SUXXXX, 3SBXXXX, AND FCXXXX

CLEARED

Aug 16, 2000
288 days to decision

K993698 · 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