FDA 510(k) Premarket Notification

K823361 SWITCHING UNIT ELECTRODE CATHETERS

CLEARED

Jan 14, 1983
66 days to decision

K823361 · Product code: **DRF** · Cardiovascular
Source: https://www.510kdatabase.net/k823361/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Nov 9, 1982
Decision date	Jan 14, 1983
Days to decision	66 days
Third-party review	No

APPLICANT

Company	**Webster Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1981-1985