FDA 510(k) Premarket Notification

K030187 REPROCESSED ELECTROPHYSIOLOGY CATHETER

CLEARED

Sep 24, 2003
246 days to decision

K030187 · Product code: **NLH** · Cardiovascular
Source: https://www.510kdatabase.net/k030187/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Recording, Electrode, Reprocessed (NLH)
Date received	Jan 21, 2003
Decision date	Sep 24, 2003
Days to decision	246 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Alliance Medical, Inc.**
Location	Santa Ynez, CA, US
Contact	MOIRA BARTON
510(k) history	11 submissions · 11 cleared · 1994-2006