FDA 510(k) Premarket Notification

K052414 REPROCESSED ELECTROPHYSIOLOGY CATHETER

CLEARED

Nov 16, 2005
75 days to decision

K052414 · Product code: **NLH** · Cardiovascular
Source: https://www.510kdatabase.net/k052414/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Recording, Electrode, Reprocessed (NLH)
Date received	Sep 2, 2005
Decision date	Nov 16, 2005
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Alliance Medical Corp.**
Location	Phoenix, AZ, US
Contact	ELIZABETH RENKEN
510(k) history	36 submissions · 36 cleared · 2001-2007

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k052414/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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