

**K030187 REPROCESSED ELECTROPHYSIOLOGY CATHETER**Sep 24, 2003  
246 days to decisionK030187 · Product code: **NLH** · Cardiovascular  
Source: <https://www.510kdatabase.net/k030187/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030187/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 19, 2026