

**K052603 REPROCESSED DAIG LIVEWIRE
ELECTROPHYSIOLOGY CATHETERS**Nov 16, 2005
56 days to decisionK052603 · Product code: **NLH** · Cardiovascular
Source: <https://www.510kdatabase.net/k052603/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Recording, Electrode, Reprocessed (NLH)
Date received	Sep 21, 2005
Decision date	Nov 16, 2005
Days to decision	56 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.netDevice record: <https://www.510kdatabase.net/k052603/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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