

**K082023 REPROCESSED ELECTROPHYSIOLOGY CATHETERS**Sep 16, 2008  
62 days to decisionK082023 · Product code: **NLH** · Cardiovascular  
Source: <https://www.510kdatabase.net/k082023/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Recording, Electrode, Reprocessed (NLH)
Date received	Jul 16, 2008
Decision date	Sep 16, 2008
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ascent Healthcare Solutions</b>
Location	Phoenix, AZ, US
Contact	AMANDA BABCOCK
510(k) history	21 submissions · 21 cleared · 2006-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k082023/> 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 14, 2026