

**K904025 MYOTRACK, MODEL HH 515**Mar 1, 1991  
183 days to decisionK904025 · Product code: **BXN** · Anesthesiology  
Source: <https://www.510kdatabase.net/k904025/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Battery-powered (BXN)
Date received	Aug 30, 1990
Decision date	Mar 1, 1991
Days to decision	183 days
Third-party review	No

**APPLICANT**

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Company	<b>Myotrack Co.</b>
Location	Van Nuys, CA, US
Contact	FREDERIC P TORRES
510(k) history	1 submissions · 1 cleared · 1991-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904025/>; 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 26, 2026