

**K823583 MYOTEST NERVE STIMULATOR**Dec 30, 1982  
24 days to decisionK823583 · Product code: **BXN** · AnesthesiologySource: <https://www.510kdatabase.net/k823583/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Battery-powered (BXN)
Date received	Dec 6, 1982
Decision date	Dec 30, 1982
Days to decision	24 days
Third-party review	No

**APPLICANT**

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Company	<b>Prothia USA, Inc.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1982-1983

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k823583/> 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 27, 2026