

**K830058 MYOGRAPH 2000**Mar 7, 1983  
59 days to decisionK830058 · Product code: **BXN** · Anesthesiology  
Source: <https://www.510kdatabase.net/k830058/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Battery-powered (BXN)
Date received	Jan 7, 1983
Decision date	Mar 7, 1983
Days to decision	59 days
Third-party review	No

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k830058/> 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