

K062478 CLAVISFeb 8, 2007
168 days to decisionK062478 · Product code: **BXN** · Anesthesiology
Source: <https://www.510kdatabase.net/k062478/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Stimulator, Nerve, Battery-powered (BXN)
Date received	Aug 24, 2006
Decision date	Feb 8, 2007
Days to decision	168 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Medtronic A/S
Location	Skovlunde, DK
Contact	ANN-CHRISTINE PROVOOST
510(k) history	1 submissions · 1 cleared · 2007-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k062478/> Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026