FDA 510(k) Premarket Notification

K062478 CLAVIS

CLEARED

Feb 8, 2007
168 days to decision

K062478 · 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