FDA 510(k) Premarket Notification

K830058 MYOGRAPH 2000

CLEARED

Mar 7, 1983
59 days to decision

K830058 · 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	**Prothia USA, Inc.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1982-1983