FDA 510(k) Premarket Notification

K904025 MYOTRACK, MODEL HH 515

CLEARED

Mar 1, 1991
183 days to decision

K904025 · Product code: **BXN** · Anesthesiology
Source: https://www.510kdatabase.net/k904025/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Battery-powered (BXN)
Date received	Aug 30, 1990
Decision date	Mar 1, 1991
Days to decision	183 days
Third-party review	No

APPLICANT

Company	**Myotrack Co.**
Location	Van Nuys, CA, US
Contact	FREDERIC P TORRES
510(k) history	1 submissions · 1 cleared · 1991-1991