FDA 510(k) Premarket Notification

K903554 NORTECH MODEL 87202204 ELECTRODE

CLEARED

Aug 30, 1990
23 days to decision

K903554 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k903554/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Aug 7, 1990
Decision date	Aug 30, 1990
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Medtronic Andover Medical, Inc.**
Location	Lowell, MA, US
Contact	JANICE M PEVIDE
510(k) history	6 submissions · 6 cleared · 1985-1991