FDA 510(k) Premarket Notification

CLEARED

K010638 NEUROPLUS, MODEL A10040, A10041, A10042, A10043

May 3, 2001
59 days to decision

K010638 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k010638/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Mar 5, 2001
Decision date	May 3, 2001
Days to decision	59 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Vermont Medical, Inc.**
Location	Bellows Falls, VT, US
Contact	MARC FILLION
510(k) history	9 submissions · 9 cleared · 1978-2003