FDA 510(k) Premarket Notification

K030800 ELECTROMED REUSABLE NEUROSTIMULATION ELECTRODES

CLEARED

Jun 11, 2003
90 days to decision

K030800 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k030800/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Mar 13, 2003
Decision date	Jun 11, 2003
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Eletromed, Inc.**
Location	Betheda, MD, US
Contact	LYDIA BAYNES
510(k) history	1 submissions · 1 cleared · 2003-2003