FDA 510(k) Premarket Notification

K081871 UNIVERSAL ELECTRODES

CLEARED

Sep 29, 2008
90 days to decision

K081871 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k081871/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Jul 1, 2008
Decision date	Sep 29, 2008
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Neurometrix, Inc.**
Location	North Attleboro, MA, US
Contact	RAINER MAAS
510(k) history	18 submissions · 17 cleared · 1998-2022