

K081871 UNIVERSAL ELECTRODESSep 29, 2008
90 days to decisionK081871 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081871/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026