

K964433 AQUABOND ELECTRODEJan 27, 1997
83 days to decisionK964433 · Product code: **GXY** · Neurology
Source: <https://www.510kdatabase.net/k964433/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Nov 5, 1996
Decision date	Jan 27, 1997
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medibond , Ltd.
Location	Efrat, IL
Contact	ELI M ORBACH
510(k) history	1 submissions · 1 cleared · 1997-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k964433/> 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 June 30, 2026