

K912685 MEM-4204-4-8, MEB-4204 &4208 EVOKE POTENT MEAS SYSFeb 26, 1992
254 days to decisionK912685 · Product code: **GWF** · Neurology
Source: <https://www.510kdatabase.net/k912685/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Jun 17, 1991
Decision date	Feb 26, 1992
Days to decision	254 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Nihon Kohden America, Inc.
Location	Foothill Ranch, CA, US
Contact	DIANA BEAL
510(k) history	166 submissions · 163 cleared · 1979-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k912685/> 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 24, 2026