

K882887 PROGEL(TM), MODELS 8630 AND 8635Sep 29, 1988
80 days to decisionK882887 · Product code: **GXY** · Neurology
Source: <https://www.510kdatabase.net/k882887/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Jul 11, 1988
Decision date	Sep 29, 1988
Days to decision	80 days
Third-party review	No

APPLICANT

Company	Medical Devices, Inc.
Location	Mchenry, IL, US
Contact	BRUCE MACFARLANE,PHD
510(k) history	49 submissions · 47 cleared · 1977-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k882887/>; 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 21, 2026