FDA 510(k) Premarket Notification

K882887 PROGEL(TM), MODELS 8630 AND 8635

CLEARED

Sep 29, 1988
80 days to decision

K882887 · 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