FDA 510(k) Premarket Notification

K862028 ULTRAHESIVE II POSTOP ELECTRODE

CLEARED

Aug 1, 1986
65 days to decision

K862028 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k862028/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	May 28, 1986
Decision date	Aug 1, 1986
Days to decision	65 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