FDA 510(k) Premarket Notification

K810206 DUAL TYPE ELECTRODE

CLEARED

Mar 20, 1981
53 days to decision

K810206 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k810206/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Jan 26, 1981
Decision date	Mar 20, 1981
Days to decision	53 days
Third-party review	No

APPLICANT

Company	**Professional Instruments Co.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1981-1987