FDA 510(k) Premarket Notification

K790316 ELECTRODE, ELECTROFLEX

CLEARED

Mar 27, 1979
40 days to decision

K790316 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k790316/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Feb 15, 1979
Decision date	Mar 27, 1979
Days to decision	40 days
Third-party review	No

APPLICANT

Company	**Andover Medical, Inc.**
Location	Mchenry, IL, US
Website	https://www.andovermedical.com
510(k) history	21 submissions · 21 cleared · 1976-1991