FDA 510(k) Premarket Notification

K831104 TENZCARE MUSCLE STIMULATION ELECTRODES

CLEARED

May 9, 1983
34 days to decision

K831104 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k831104/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Apr 5, 1983
Decision date	May 9, 1983
Days to decision	34 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

Company	**3M Company**
Location	White City, OR, US
Website	http://www.3m.com/
510(k) history	331 submissions · 322 cleared · 1976-2025