

K831104 TENZCARE MUSCLE STIMULATION ELECTRODESMay 9, 1983
34 days to decisionK831104 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k831104/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026