

K891992 TENZCARE STIMULATOR 6890Oct 12, 1989
198 days to decisionK891992 · Product code: **GZJ** · Neurology
Source: <https://www.510kdatabase.net/k891992/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Mar 28, 1989
Decision date	Oct 12, 1989
Days to decision	198 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k891992/> 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