FDA 510(k) Premarket Notification

K891992 TENZCARE STIMULATOR 6890

CLEARED

Oct 12, 1989
198 days to decision

K891992 · 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