FDA 510(k) Premarket Notification

K962332 TENS/FES/NMES ELECTRODES

CLEARED

Jul 31, 1996
44 days to decision

K962332 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k962332/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jun 17, 1996
Decision date	Jul 31, 1996
Days to decision	44 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Uni-Patch, Inc.**
Location	Wabasha, MN, US
Contact	THOMAS MOORE
510(k) history	6 submissions · 6 cleared · 1987-1999