FDA 510(k) Premarket Notification

K896787 TRANSCUTANEOUS ELECTRICAL NERVE STIMULATION

CLEARED

Jul 30, 1990
238 days to decision

K896787 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k896787/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Dec 4, 1989
Decision date	Jul 30, 1990
Days to decision	238 days
Third-party review	No

APPLICANT

Company	**Saratoga Intl., Inc.**
Location	San Jose, CA, US
Contact	MIKE CHUANG
510(k) history	2 submissions · 2 cleared · 1990-1994