FDA 510(k) Premarket Notification

K903017 MEGA-TENS DUAL CHANNEL T.E.N.S.

CLEARED

Oct 26, 1990
108 days to decision

K903017 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k903017/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jul 10, 1990
Decision date	Oct 26, 1990
Days to decision	108 days
Third-party review	No

APPLICANT

Company	**Futuremed Div. of Future Impex Corp.**
Location	Deerpark, NY, US
Contact	MIKE DAVIDSON
510(k) history	17 submissions · 17 cleared · 1985-1994