FDA 510(k) Premarket Notification

K881832 DS-200 AND DS-400 T.E.N.S.

CLEARED

Jul 1, 1988
63 days to decision

K881832 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k881832/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Apr 29, 1988
Decision date	Jul 1, 1988
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**Delta Marketing, Inc.**
Location	Muskogee, OK, US
Contact	DAVID R WATERS
510(k) history	1 submissions · 1 cleared · 1988-1988