FDA 510(k) Premarket Notification

K871666 MODEL HP (OR MODEL 1001) TRANSCUTANEOUS STIMULATOR

CLEARED

Jun 25, 1987
58 days to decision

K871666 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k871666/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Apr 28, 1987
Decision date	Jun 25, 1987
Days to decision	58 days
Third-party review	No

APPLICANT

Company	**Oriental Medical Supplies, Inc.**
Location	Braintree, MA, US
Contact	THOMAS RIIHIMAKI
510(k) history	6 submissions · 6 cleared · 1985-1989