FDA 510(k) Premarket Notification

K901417 ELECTROSTIM MST

CLEARED

Aug 28, 1990
155 days to decision

K901417 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k901417/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Mar 26, 1990
Decision date	Aug 28, 1990
Days to decision	155 days
Third-party review	No

APPLICANT

Company	**Promatek Industries, Ltd.**
Location	Quebec, H4p 2c9, CA
Contact	ARTHUR LEVINE
510(k) history	3 submissions · 2 cleared · 1987-1990

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k901417/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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