FDA 510(k) Premarket Notification

K931424 MAXIMA I

CLEARED

Jun 4, 1993
74 days to decision

K931424 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k931424/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Mar 22, 1993
Decision date	Jun 4, 1993
Days to decision	74 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Staodyne, Inc.**
Location	Longmont, CO, US
Contact	THOMAS W ROBIRDS
510(k) history	9 submissions · 9 cleared · 1992-1995