FDA 510(k) Premarket Notification

K914441 MENS 2000-D

CLEARED

Dec 17, 1991
78 days to decision

K914441 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k914441/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Sep 30, 1991
Decision date	Dec 17, 1991
Days to decision	78 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Monad Corp.**
Location	Pomona, CA, US
Contact	MARK HANKINS
510(k) history	9 submissions · 9 cleared · 1989-1991