

K914441 MENS 2000-DDec 17, 1991
78 days to decisionK914441 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k914441/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026