

K993229 ANALGESIC PULSER, MODEL AP-439Feb 2, 2001
494 days to decisionK993229 · Product code: **GZJ** · Neurology
Source: <https://www.510kdatabase.net/k993229/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Sep 27, 1999
Decision date	Feb 2, 2001
Days to decision	494 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Johari Electro-Tech Co.
Location	Jodhpur-342003, IN
Contact	NISHA JOHARI
510(k) history	1 submissions · 1 cleared · 2001-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k993229/> 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