FDA 510(k) Premarket Notification

# K993229 ANALGESIC PULSER, MODEL AP-439

**CLEARED**

Feb 2, 2001
494 days to decision

K993229 · 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 |