FDA 510(k) Premarket Notification

# K812432 HEALTH-UP

**CLEARED**

Nov 6, 1981
73 days to decision

K812432 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k812432/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ) |
| Date received | Aug 25, 1981 |
| Decision date | Nov 6, 1981 |
| Days to decision | 73 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Wfx Industries, Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 1 submissions · 1 cleared · 1981-1981 |