FDA 510(k) Premarket Notification

# K930865 STAODYN MAXIMA III

**CLEARED**

May 13, 1993
84 days to decision

K930865 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k930865/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ) |
| Date received | Feb 18, 1993 |
| Decision date | May 13, 1993 |
| Days to decision | 84 days |
| Third-party review | No |
| Summary / Statement | Statement |

## APPLICANT

| | |
|---|---|
| Company | **Staodyne, Inc.** |
| Location | Longmont, CO, US |
| Contact | THOMAS W ROBIRDS |
| 510(k) history | 9 submissions · 9 cleared · 1992-1995 |

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k930865/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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