FDA 510(k) Premarket Notification

# K830600 REFERENCE & MONITORING ELECTRODE

**CLEARED**

Mar 24, 1983
28 days to decision

K830600 · Product code: **GXY** · Neurology
Source: https://www.510kdatabase.net/k830600/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrode, Cutaneous (GXY) |
| Date received | Feb 24, 1983 |
| Decision date | Mar 24, 1983 |
| Days to decision | 28 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Precision Cap Systems** |
| Location | Mchenry, IL, US |
| 510(k) history | 7 submissions · 7 cleared · 1983-1983 |