FDA 510(k) Premarket Notification

# K850439 DESERET EPIDURAL CATHETER

**CLEARED**

Mar 5, 1985
28 days to decision

K850439 · Product code: **BSO** · Anesthesiology
Source: https://www.510kdatabase.net/k850439/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Conduction, Anesthetic (BSO) |
| Date received | Feb 5, 1985 |
| Decision date | Mar 5, 1985 |
| Days to decision | 28 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Parke-Davis Co.** |
| Location | Mchenry, IL, US |
| Contact | CLEARY |
| 510(k) history | 47 submissions · 47 cleared · 1976-1986 |

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k850439/ 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 May 27, 2026