

K934787 SILICORE SEMI-FLOATERFeb 24, 1995
507 days to decisionK934787 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k934787/>**SUBMISSION DETAILS**

| | |
|-----------------------|---------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrode, Pacemaker, Temporary (LDF) |
| Date received | Oct 5, 1993 |
| Decision date | Feb 24, 1995 |
| Days to decision | 507 days |
| Third-party review | No |
| Summary / Statement | Statement |

APPLICANT

| | |
|----------------|---|
| Company | Electro-Catheter Corp. |
| Location | Mchenry, IL, US |
| Contact | LORRAINE T MONTEMURRO |
| 510(k) history | 35 submissions · 35 cleared · 1976-1995 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k934787/> 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