

K823361 SWITCHING UNIT ELECTRODE CATHETERSJan 14, 1983
66 days to decisionK823361 · Product code: **DRF** · CardiovascularSource: <https://www.510kdatabase.net/k823361/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF) |
| Date received | Nov 9, 1982 |
| Decision date | Jan 14, 1983 |
| Days to decision | 66 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Webster Laboratories, Inc. |
| Location | Mchenry, IL, US |
| 510(k) history | 5 submissions · 5 cleared · 1981-1985 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k823361/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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