

K834602 SWITCHING UNIT ELECTRODE CATHETERMay 9, 1984
131 days to decisionK834602 · Product code: **DRF** · CardiovascularSource: <https://www.510kdatabase.net/k834602/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Dec 30, 1983
Decision date	May 9, 1984
Days to decision	131 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k834602/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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