

K823758 CENTRAL VEIN CATHORIZATION KITFeb 23, 1983
71 days to decisionK823758 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k823758/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Intravascular, Diagnostic (DQO) |
| Date received | Dec 14, 1982 |
| Decision date | Feb 23, 1983 |
| Days to decision | 71 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Access Devices, Inc. |
| Location | Walker, MI, US |
| 510(k) history | 3 submissions · 3 cleared · 1982-1983 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k823758/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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