

K874138 ACCUCOM 2Jun 1, 1988
232 days to decisionK874138 · Product code: **DPT** · Cardiovascular
Source: <https://www.510kdatabase.net/k874138/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Probe, Blood-flow, Extravascular (DPT) |
| Date received | Oct 13, 1987 |
| Decision date | Jun 1, 1988 |
| Days to decision | 232 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Datascope Corp. |
| Location | Mchenry, IL, US |
| Contact | ARTHUR FRIEDMAN |
| 510(k) history | 136 submissions · 135 cleared · 1976-2019 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k874138/>; 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 June 20, 2026