

K833303 DUOFLO 6FDec 27, 1983
95 days to decisionK833303 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k833303/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Percutaneous (DQY) |
| Date received | Sep 23, 1983 |
| Decision date | Dec 27, 1983 |
| Days to decision | 95 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Argon Medical Corp. |
| Location | Mchenry, IL, US |
| 510(k) history | 27 submissions · 27 cleared · 1976-1991 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k833303/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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