

K833433 PHYSIO-PROBE-HIGH PERMEABILITY CATH.Dec 29, 1983
85 days to decisionK833433 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k833433/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Oct 5, 1983
Decision date	Dec 29, 1983
Days to decision	85 days
Third-party review	No

APPLICANT

Company	Research Industries Corp.
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1983-1995

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Device record: <https://www.510kdatabase.net/k833433/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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