FDA 510(k) Premarket Notification

# K833433 PHYSIO-PROBE-HIGH PERMEABILITY CATH.

**CLEARED**

Dec 29, 1983
85 days to decision

K833433 · Product code: **DQY** · Cardiovascular
Source: 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 |