FDA 510(k) Premarket Notification

K854610 VASCULAR ACCESS PORT/CATH CAT.#VP-XXXX VASPORT SER

CLEARED

May 8, 1986
171 days to decision

K854610 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k854610/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Nov 18, 1985
Decision date	May 8, 1986
Days to decision	171 days
Third-party review	No

APPLICANT

Company	**Gish Biomedical, Inc.**
Location	Mchenry, IL, US
Contact	JEANNE PIERSON
510(k) history	75 submissions · 75 cleared · 1983-2009

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k854610/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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