FDA 510(k) Premarket Notification

K863790 J-CATH, GISH MODEL NO. JVAC-XXXX

CLEARED

Jun 24, 1987
271 days to decision

K863790 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k863790/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Sep 26, 1986
Decision date	Jun 24, 1987
Days to decision	271 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/k863790/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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