FDA 510(k) Premarket Notification

K863792 DURACATH, GISH MODEL NO. DVAC-XXXX

CLEARED

Dec 5, 1986
70 days to decision

K863792 · Product code: **BSO** · Anesthesiology
Source: https://www.510kdatabase.net/k863792/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Conduction, Anesthetic (BSO)
Date received	Sep 26, 1986
Decision date	Dec 5, 1986
Days to decision	70 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/k863792/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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