FDA 510(k) Premarket Notification

CLEARED

K862540 LEFT VENTRICULAR VENT VALVE MODEL NO. GLV

Aug 29, 1986
58 days to decision

K862540 · Product code: **DWD** · Cardiovascular
Source: https://www.510kdatabase.net/k862540/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suction Control, Intracardiac, Cardiopulmonary Bypass (DWD)
Date received	Jul 2, 1986
Decision date	Aug 29, 1986
Days to decision	58 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/k862540/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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