FDA 510(k) Premarket Notification

K833322 TUBING CONNECTORS EC-2XXX SERIES

CLEARED

Dec 16, 1983
81 days to decision

K833322 · Product code: **DTL** · Cardiovascular
Source: https://www.510kdatabase.net/k833322/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Adaptor, Stopcock, Manifold, Fitting, Cardiopulmonary Bypass (DTL)
Date received	Sep 26, 1983
Decision date	Dec 16, 1983
Days to decision	81 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k833322/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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