

K864503 MODIFIED RLV-2100 BJan 6, 1987
53 days to decisionK864503 · Product code: **DWD** · CardiovascularSource: <https://www.510kdatabase.net/k864503/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suction Control, Intracardiac, Cardiopulmonary Bypass (DWD)
Date received	Nov 14, 1986
Decision date	Jan 6, 1987
Days to decision	53 days
Third-party review	No

APPLICANT

Company	American Omni Medical, Inc.
Location	Costa Mesa, CA, US
Contact	GEORGE G SIPOSS
510(k) history	20 submissions · 20 cleared · 1985-1993

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