

K760894 VACUUM RELIEF VALVEDec 9, 1976
44 days to decisionK760894 · Product code: **DWD** · CardiovascularSource: <https://www.510kdatabase.net/k760894/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suction Control, Intracardiac, Cardiopulmonary Bypass (DWD)
Date received	Oct 26, 1976
Decision date	Dec 9, 1976
Days to decision	44 days
Third-party review	No

APPLICANT

Company	Delta Medical Industries
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1976-1984

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Device record: <https://www.510kdatabase.net/k760894/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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