FDA 510(k) Premarket Notification

K760894 VACUUM RELIEF VALVE

CLEARED

Dec 9, 1976
44 days to decision

K760894 · Product code: **DWD** · Cardiovascular
Source: 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