FDA 510(k) Premarket Notification

K863593 SARNS SAFETY LOOP

CLEARED

Nov 18, 1986
64 days to decision

K863593 · Product code: **DWD** · Cardiovascular
Source: https://www.510kdatabase.net/k863593/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suction Control, Intracardiac, Cardiopulmonary Bypass (DWD)
Date received	Sep 15, 1986
Decision date	Nov 18, 1986
Days to decision	64 days
Third-party review	No

APPLICANT

Company	**3M Health Care, Sarns**
Location	Mchenry, IL, US
Contact	JOSEPH W O'DONNELL
510(k) history	76 submissions · 76 cleared · 1976-1996