FDA 510(k) Premarket Notification

K896671 SARNS MEMBRANE OXYGENATOR W/INTEGRAL RESER. #9461

CLEARED

Feb 14, 1990
79 days to decision

K896671 · Product code: **DTZ** · Cardiovascular
Source: https://www.510kdatabase.net/k896671/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oxygenator, Cardiopulmonary Bypass (DTZ)
Date received	Nov 27, 1989
Decision date	Feb 14, 1990
Days to decision	79 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

Company	**3M Company**
Location	White City, OR, US
Contact	D SHIMOKOCHI
Website	http://www.3m.com/
510(k) history	331 submissions · 322 cleared · 1976-2025