FDA 510(k) Premarket Notification

K800555 PARTNER SEVEN DISP. BUBBLE OXYGENATOR

CLEARED

Apr 4, 1980
23 days to decision

K800555 · Product code: **DTZ** · Cardiovascular
Source: https://www.510kdatabase.net/k800555/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oxygenator, Cardiopulmonary Bypass (DTZ)
Date received	Mar 12, 1980
Decision date	Apr 4, 1980
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Cardiovascular Research, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1984