CLEARED

K022071 EMBOL-X AORTIC FILTERS,ETRA-SMALL,SMALL,MEDIUM,LARGE,EXTRA LARGE,MODELS CF26000.CF29000,CF32000,CF35000,CF4000

Feb 14, 2003
233 days to decision

K022071 · Product code: **DTM** · Cardiovascular
Source: https://www.510kdatabase.net/k022071/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Blood, Cardiopulmonary Bypass, Arterial Line (DTM)
Date received	Jun 26, 2002
Decision date	Feb 14, 2003
Days to decision	233 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Embol-X, Inc.**
Location	Mountain View, CA, US
Contact	ED LEE
510(k) history	4 submissions · 4 cleared · 1999-2003

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k022071/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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