FDA 510(k) Premarket Notification

K934782 AMMEX TUBING ORGANIZER

CLEARED

Nov 18, 1993
43 days to decision

K934782 · Product code: **KRI** · Cardiovascular
Source: https://www.510kdatabase.net/k934782/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessory Equipment, Cardiopulmonary Bypass (KRI)
Date received	Oct 6, 1993
Decision date	Nov 18, 1993
Days to decision	43 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ammex Cardiopulmonary Corp.**
Location	Houston, TX, US
Contact	HILLARD E MILLER III
510(k) history	1 submissions · 1 cleared · 1993-1993