

**K934782 AMMEX TUBING ORGANIZER**Nov 18, 1993  
43 days to decisionK934782 · 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	<b>Ammex Cardiopulmonary Corp.</b>
Location	Houston, TX, US
Contact	HILLARD E MILLER III
510(k) history	1 submissions · 1 cleared · 1993-1993

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k934782/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 6, 2026