

K780479 DELTA-FLOWNov 22, 1978
240 days to decisionK780479 · Product code: **KRA** · CardiovascularSource: <https://www.510kdatabase.net/k780479/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Continuous Flush (KRA)
Date received	Mar 27, 1978
Decision date	Nov 22, 1978
Days to decision	240 days
Third-party review	No

APPLICANT

Company	Tri-Delta Intl.
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1978-1978

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k780479/> Data sourced from FDA 510(k) public records (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. - Generated July 9, 2026