FDA 510(k) Premarket Notification

K780479 DELTA-FLOW

CLEARED

Nov 22, 1978
240 days to decision

K780479 · Product code: **KRA** · Cardiovascular
Source: 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