FDA 510(k) Premarket Notification

K860298 STABLELINE

CLEARED

Apr 2, 1986
64 days to decision

K860298 · Product code: **KRI** · Cardiovascular
Source: https://www.510kdatabase.net/k860298/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessory Equipment, Cardiopulmonary Bypass (KRI)
Date received	Jan 28, 1986
Decision date	Apr 2, 1986
Days to decision	64 days
Third-party review	No

APPLICANT

Company	**Cardio Metrics, Inc.**
Location	Houston, TX, US
Contact	MARY K NOONEN
510(k) history	23 submissions · 23 cleared · 1986-1997