FDA 510(k) Premarket Notification

K832058 ANGIO-FLO CONTIN-ARTERIAL FLUSH DEV.

CLEARED

Sep 12, 1983
77 days to decision

K832058 · Product code: **KRA** · Cardiovascular
Source: https://www.510kdatabase.net/k832058/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Continuous Flush (KRA)
Date received	Jun 27, 1983
Decision date	Sep 12, 1983
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**Parke-Davis Co.**
Location	Mchenry, IL, US
510(k) history	47 submissions · 47 cleared · 1976-1986

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k832058/ 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 May 27, 2026