

K790327 ANGIO-FLOFeb 26, 1979
10 days to decisionK790327 · Product code: **KRA** · CardiovascularSource: <https://www.510kdatabase.net/k790327/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Continuous Flush (KRA)
Date received	Feb 16, 1979
Decision date	Feb 26, 1979
Days to decision	10 days
Third-party review	No

APPLICANT

Company	Deseret Medical, Inc.
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1979-1991

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k790327/> 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 1, 2026