FDA 510(k) Premarket Notification

K790327 ANGIO-FLO

CLEARED

Feb 26, 1979
10 days to decision

K790327 · Product code: **KRA** · Cardiovascular
Source: 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