FDA 510(k) Premarket Notification

CLEARED

K990805 MODIFICATION TO 7F HYDROLYSER THROMBECTOMY CATHETER

Mar 15, 1999
4 days to decision

K990805 · Product code: **QEZ** · Cardiovascular
Source: https://www.510kdatabase.net/k990805/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Aspiration Thrombectomy Catheter (QEZ)
Date received	Mar 11, 1999
Decision date	Mar 15, 1999
Days to decision	4 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cordis Corp.**
Location	Mchenry, IL, US
Contact	ARIEL MACTAVISH
Website	https://cordis.com
510(k) history	315 submissions · 281 cleared · 1976-2014

Cordis Corp. is a medical device manufacturer based in McHenry, US. The company specializes in interventional cardiovascular and gastroenterology devices. Cordis has a substantial FDA 510(k) regulatory history spanning from 1976 to 2014. The company received FDA 510(k) clearances from total submissions. Its portfolio focuses primarily on cardiovascular devices and gastroenterology stent systems, including percutaneous transluminal angioplasty catheters, emboli capture guidewires, and self-expanding biliary stent systems. Notable cleared products include the FLEXSTENT Bili...