FDA 510(k) Premarket Notification

CLEARED

K020316 CORDIS TRAPEASE PERMANENT VENA CAVA FILTER WITH THE VISEASE ANGIOGRAPHIC VESSEL DILATOR

Mar 20, 2002
49 days to decision

K020316 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k020316/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jan 30, 2002
Decision date	Mar 20, 2002
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary
Other names	MODELS 466-P306AU & 466-P306BU

APPLICANT

Company	**Cordis Corp.**
Location	Mchenry, IL, US
Contact	KAREN WILK
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...