FDA 510(k) Premarket Notification

K884375 TRANSLUMBAR AORTOGRAPHY NEEDLE CATHETER

CLEARED

Jan 11, 1989
85 days to decision

K884375 · Product code: **DQR** · Cardiovascular
Source: https://www.510kdatabase.net/k884375/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Catheter (DQR)
Date received	Oct 18, 1988
Decision date	Jan 11, 1989
Days to decision	85 days
Third-party review	No

APPLICANT

Company	**Argon Medical Corp.**
Location	Mchenry, IL, US
Contact	BRISTER, P.E.
510(k) history	27 submissions · 27 cleared · 1976-1991