FDA 510(k) Premarket Notification

K882221 OVER THE NEEDLE CATHETER, ARTERIAL CATHETER NEEDLE

CLEARED

Aug 12, 1988
78 days to decision

K882221 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k882221/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	May 26, 1988
Decision date	Aug 12, 1988
Days to decision	78 days
Third-party review	No

APPLICANT

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