FDA 510(k) Premarket Notification

K854018 ANGIOGRAPHY CATHETERS

CLEARED

Jan 16, 1986
108 days to decision

K854018 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k854018/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Sep 30, 1985
Decision date	Jan 16, 1986
Days to decision	108 days
Third-party review	No

APPLICANT

Company	**Micor, Inc.**
Location	Pittsburgh, PA, US
Contact	STEPHEN BRUSHEY
510(k) history	8 submissions · 8 cleared · 1986-2003