FDA 510(k) Premarket Notification

K821816 ANGIOGRAPHIC CATHETER

CLEARED

Aug 24, 1982
64 days to decision

K821816 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k821816/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jun 21, 1982
Decision date	Aug 24, 1982
Days to decision	64 days
Third-party review	No

APPLICANT

Company	**Medi-Tech, Inc.**
Location	Mchenry, IL, US
510(k) history	36 submissions · 35 cleared · 1978-1996