FDA 510(k) Premarket Notification

K884354 IMAGER FLUSH CATHETERS

CLEARED

Apr 21, 1989
186 days to decision

K884354 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k884354/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Oct 17, 1988
Decision date	Apr 21, 1989
Days to decision	186 days
Third-party review	No

APPLICANT

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