FDA 510(k) Premarket Notification

K860429 MEADOX SURGIMED GUIDE CATHETER

CLEARED

Apr 23, 1986
78 days to decision

K860429 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k860429/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Feb 4, 1986
Decision date	Apr 23, 1986
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Meadox Surgimed, Inc.**
Location	Mchenry, IL, US
Contact	JAN AOYAGI
510(k) history	24 submissions · 24 cleared · 1984-1995