FDA 510(k) Premarket Notification

K860430 THE NAVIGATOR STEERABLE GUIDE WIRE

CLEARED

Jul 18, 1986
164 days to decision

K860430 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k860430/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 4, 1986
Decision date	Jul 18, 1986
Days to decision	164 days
Third-party review	No

APPLICANT

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