FDA 510(k) Premarket Notification

K843012 WIREGUIDE OR GUIDEWIRE

CLEARED

Nov 29, 1984
121 days to decision

K843012 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k843012/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jul 31, 1984
Decision date	Nov 29, 1984
Days to decision	121 days
Third-party review	No

APPLICANT

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