FDA 510(k) Premarket Notification

K883880 KATZEN THROMBOLYSIS GUIDEWIRE

CLEARED

Apr 6, 1989
205 days to decision

K883880 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k883880/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Sep 13, 1988
Decision date	Apr 6, 1989
Days to decision	205 days
Third-party review	No

APPLICANT

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