FDA 510(k) Premarket Notification

K871850 TELEFLEX STEERABLE GUIDEWIRE

CLEARED

Jan 27, 1988
260 days to decision

K871850 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k871850/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	May 12, 1987
Decision date	Jan 27, 1988
Days to decision	260 days
Third-party review	No

APPLICANT

Company	**Teleflexmedical, Inc.**
Location	Jeffrey, NH, US
Contact	RICHARD GARDNER
510(k) history	64 submissions · 61 cleared · 1985-2024