FDA 510(k) Premarket Notification

K041762 FINISHING WIRE SUPPORTRAK

CLEARED

Aug 6, 2004
37 days to decision

K041762 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k041762/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 30, 2004
Decision date	Aug 6, 2004
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Guidant Corporation**
Location	St. Paul, MN, US
Contact	JENNIFER TANG
510(k) history	15 submissions · 12 cleared · 2003-2006