FDA 510(k) Premarket Notification

K022764 7F LAUNCHER GUIDE CATHETER

CLEARED

Aug 28, 2002
7 days to decision

K022764 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k022764/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Aug 21, 2002
Decision date	Aug 28, 2002
Days to decision	7 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Vascular**
Location	Walker, MI, US
Contact	FRED L BOUCHER
510(k) history	475 submissions · 453 cleared · 1977-2023