FDA 510(k) Premarket Notification

K051917 DIVER C.E. CATHETER

CLEARED

Aug 8, 2005
24 days to decision

K051917 · Product code: **QEZ** · Cardiovascular
Source: https://www.510kdatabase.net/k051917/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Aspiration Thrombectomy Catheter (QEZ)
Date received	Jul 15, 2005
Decision date	Aug 8, 2005
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Invatec Innovative Technologies, S.R.L.**
Location	Plymouth, MN, US
Contact	STEPHANIE K ISGRIGG ROBINSON
510(k) history	7 submissions · 7 cleared · 2005-2009