FDA 510(k) Premarket Notification

K050023 SECURELOC

CLEARED

Feb 18, 2005
44 days to decision

K050023 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k050023/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Jan 5, 2005
Decision date	Feb 18, 2005
Days to decision	44 days
Third-party review	No
Summary / Statement	Summary
Other names	ADVANCINTRODUCER

APPLICANT

Company	**Specialized Health Products, Inc.**
Location	Bountiful, UT, US
Contact	Mark Nelson
510(k) history	4 submissions · 4 cleared · 1996-2005