

K050023 SECURELOCFeb 18, 2005
44 days to decisionK050023 · Product code: **DYB** · CardiovascularSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k050023/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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