

K924491 INTRODUCER, PERCUTANEOUS -- MODIFICATIONNov 18, 1992
90 days to decisionK924491 · Product code: **DYB** · CardiovascularSource: <https://www.510kdatabase.net/k924491/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Aug 20, 1992
Decision date	Nov 18, 1992
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Medamicus, Inc.
Location	Minneapolis, MN, US
Contact	DAVID A LIEBL
510(k) history	20 submissions · 20 cleared · 1989-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k924491/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026