FDA 510(k) Premarket Notification

K893658 PERCUTANEOUS INTRODUCER

CLEARED

Aug 3, 1989
80 days to decision

K893658 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k893658/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	May 15, 1989
Decision date	Aug 3, 1989
Days to decision	80 days
Third-party review	No

APPLICANT

Company	**Medamicus, Inc.**
Location	Minneapolis, MN, US
Contact	DENNIS MADISON
510(k) history	20 submissions · 20 cleared · 1989-2004