FDA 510(k) Premarket Notification

K903906 PERCUTANEOUS CATHETER INTRODUCER KIT

CLEARED

Sep 26, 1990
34 days to decision

K903906 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k903906/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Aug 23, 1990
Decision date	Sep 26, 1990
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Intec Medical, Inc.**
Location	Blue Springs, MO, US
Contact	KENNETH A SPECTOR
510(k) history	17 submissions · 17 cleared · 1980-1991