FDA 510(k) Premarket Notification

K870907 INTRODUCER, PERCUTANEOUS

CLEARED

Apr 9, 1987
35 days to decision

K870907 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k870907/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Mar 5, 1987
Decision date	Apr 9, 1987
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Med Fusion Systems, Inc.**
Location	Mchenry, IL, US
Contact	RUFFIN BOOTH
510(k) history	22 submissions · 22 cleared · 1977-1987