FDA 510(k) Premarket Notification

K861951 MODIFIED TELEFLEX MEDICAL INTRODUCER CATHETER

CLEARED

Jun 19, 1986
30 days to decision

K861951 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k861951/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	May 20, 1986
Decision date	Jun 19, 1986
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Teleflex, Inc.**
Location	Jeffrey, NH, US
Contact	GARY C BULLARD
510(k) history	10 submissions · 10 cleared · 1986-2016

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k861951/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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