FDA 510(k) Premarket Notification

K900756 ITC INTRODUCER CATHETER

CLEARED

Aug 8, 1990
173 days to decision

K900756 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k900756/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Feb 16, 1990
Decision date	Aug 8, 1990
Days to decision	173 days
Third-party review	No

APPLICANT

Company	**Interventional Therapeutics Corp.**
Location	Santa Barbara, CA, US
Contact	JULIE D BELL
Website	https://www.itcmed.com
510(k) history	28 submissions · 27 cleared · 1987-1996

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k900756/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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