FDA 510(k) Premarket Notification

K905773 PERCUTANEOUS TRANSLUMINAL ANGIOPLASTY CATHETER

CLEARED

Jan 30, 1991
34 days to decision

K905773 · Product code: **LIT** · Cardiovascular
Source: https://www.510kdatabase.net/k905773/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Angioplasty, Peripheral, Transluminal (LIT)
Date received	Dec 27, 1990
Decision date	Jan 30, 1991
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Mallinckrodt Medical**
Location	St Louis, MO, US
Contact	DAVID E BROWN
510(k) history	43 submissions · 42 cleared · 1990-2008

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k905773/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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