FDA 510(k) Premarket Notification

K800703 MODIFIED CATHETER PACKAGE

CLEARED

Apr 8, 1980
12 days to decision

K800703 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k800703/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Mar 27, 1980
Decision date	Apr 8, 1980
Days to decision	12 days
Third-party review	No

APPLICANT

Company	**Hancock Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1978-1980

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k800703/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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