FDA 510(k) Premarket Notification

K840175 DOUBLE LUMEN INFUSION CATHETERS

CLEARED

Feb 10, 1984
24 days to decision

K840175 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k840175/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jan 17, 1984
Decision date	Feb 10, 1984
Days to decision	24 days
Third-party review	No

APPLICANT

Company	**Medical Components, Inc.**
Location	Mchenry, IL, US
510(k) history	63 submissions · 55 cleared · 1980-2025