FDA 510(k) Premarket Notification

K800741 CENTRAL VEIN CATHETERIZATION KIT

CLEARED

Apr 16, 1980
13 days to decision

K800741 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k800741/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Apr 3, 1980
Decision date	Apr 16, 1980
Days to decision	13 days
Third-party review	No

APPLICANT

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