FDA 510(k) Premarket Notification

K822243 GUIDING CATHETER

CLEARED

Aug 24, 1982
28 days to decision

K822243 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k822243/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jul 27, 1982
Decision date	Aug 24, 1982
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Interventional Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1982-1988