FDA 510(k) Premarket Notification

K790083 CATHETER, ARTERIAL LINE

CLEARED

Jan 25, 1979
13 days to decision

K790083 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k790083/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jan 12, 1979
Decision date	Jan 25, 1979
Days to decision	13 days
Third-party review	No

APPLICANT

Company	**Namic**
Location	Walker, MI, US
510(k) history	8 submissions · 8 cleared · 1979-1996