FDA 510(k) Premarket Notification

K790034 INTRODUCERS

CLEARED

Feb 1, 1979
29 days to decision

K790034 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k790034/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Jan 3, 1979
Decision date	Feb 1, 1979
Days to decision	29 days
Third-party review	No

APPLICANT

Company	**Clincal Instruments Corp.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1977-1991