FDA 510(k) Premarket Notification

K911527 NOVOSTE GUIDING CATHETER

CLEARED

Apr 24, 1991
20 days to decision

K911527 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k911527/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Apr 4, 1991
Decision date	Apr 24, 1991
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Novoste Corp.**
Location	Miami, FL, US
Contact	LEROY S FORNEY
510(k) history	5 submissions · 5 cleared · 1988-1995