FDA 510(k) Premarket Notification

K820516 ARTERIAL CANALIZATION DEVICE

CLEARED

Apr 15, 1982
49 days to decision

K820516 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k820516/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Feb 25, 1982
Decision date	Apr 15, 1982
Days to decision	49 days
Third-party review	No

APPLICANT

Company	**Alimed, Inc.**
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1979-1996