FDA 510(k) Premarket Notification

K813577 PRESSURE INDICATOR-REGULATOR

CLEARED

Jan 29, 1982
37 days to decision

K813577 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k813577/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Dec 23, 1981
Decision date	Jan 29, 1982
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Advanced Catheter Systems**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1982-1982