FDA 510(k) Premarket Notification

K810859 SINGLE USE SELDINGER/COURNAND PERC. NDL

CLEARED

Apr 11, 1981
12 days to decision

K810859 · Product code: **DRE** · Cardiovascular
Source: https://www.510kdatabase.net/k810859/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	Mar 30, 1981
Decision date	Apr 11, 1981
Days to decision	12 days
Third-party review	No

APPLICANT

Company	**Procedure Products, Inc.**
Location	Mchenry, IL, US
510(k) history	16 submissions · 16 cleared · 1981-2017