

K810859 SINGLE USE SELDINGER/COURNAND PERC. NDLApr 11, 1981
12 days to decisionK810859 · Product code: **DRE** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k810859/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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