

K840671 CARDIAC CATHETERIZATION PACKMar 12, 1984
20 days to decisionK840671 · Product code: **DRE** · CardiovascularSource: <https://www.510kdatabase.net/k840671/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	Feb 21, 1984
Decision date	Mar 12, 1984
Days to decision	20 days
Third-party review	No

APPLICANT

Company	Professional Medical Services
Location	Mchenry, IL, US
510(k) history	19 submissions · 18 cleared · 1983-1992

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Device record: <https://www.510kdatabase.net/k840671/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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