

K840017 VERTEX PERCUTANEOUS CATHETER INTRODUCFeb 4, 1984
31 days to decisionK840017 · Product code: **DRE** · CardiovascularSource: <https://www.510kdatabase.net/k840017/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	Jan 4, 1984
Decision date	Feb 4, 1984
Days to decision	31 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/k840017/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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