

K843261 ANGIOSYSTEMS STANDARD ARTERIOGRAM & CARDSep 11, 1984
22 days to decisionK843261 · Product code: **DRE** · CardiovascularSource: <https://www.510kdatabase.net/k843261/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	Aug 20, 1984
Decision date	Sep 11, 1984
Days to decision	22 days
Third-party review	No

APPLICANT

Company	Angiosystems, Inc.
Location	Mchenry, IL, US
510(k) history	14 submissions · 11 cleared · 1984-1991

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Device record: <https://www.510kdatabase.net/k843261/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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