

K812640 URE SIL EMBOLECTOMY/THROMBECTOMY CATHOct 2, 1981
16 days to decisionK812640 · Product code: **DXE** · CardiovascularSource: <https://www.510kdatabase.net/k812640/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Sep 16, 1981
Decision date	Oct 2, 1981
Days to decision	16 days
Third-party review	No

APPLICANT

Company	Uresil Corp.
Location	Mchenry, IL, US
510(k) history	45 submissions · 44 cleared · 1981-2001

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Device record: <https://www.510kdatabase.net/k812640/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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