

**K801650 DESERET EMBOLECTOMY/THROMBECTOMY CATH.**Aug 4, 1980  
17 days to decisionK801650 · Product code: **DXE** · CardiovascularSource: <https://www.510kdatabase.net/k801650/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Jul 18, 1980
Decision date	Aug 4, 1980
Days to decision	17 days
Third-party review	No

**APPLICANT**

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Company	<b>Deseret Medical, Inc.</b>
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1979-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k801650/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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