

**K770569 CATHETER, EMBOLECTOMY, UMI**Apr 5, 1977  
11 days to decisionK770569 · Product code: **DXE** · CardiovascularSource: <https://www.510kdatabase.net/k770569/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Mar 25, 1977
Decision date	Apr 5, 1977
Days to decision	11 days
Third-party review	No

**APPLICANT**

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Company	<b>Universal Medical Instrument Corp.</b>
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1977-1996

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Device record: <https://www.510kdatabase.net/k770569/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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