

**K761031 ELECATH ARTERIAL EMBOLECTOMY CATHETER**Dec 13, 1976  
28 days to decisionK761031 · Product code: **DXE** · CardiovascularSource: <https://www.510kdatabase.net/k761031/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Nov 15, 1976
Decision date	Dec 13, 1976
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Electro-Catheter Corp.</b>
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1976-1995

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Device record: <https://www.510kdatabase.net/k761031/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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