

**K892410 FOGARTY(R) THRU-LUMEN EMBOLECTOMY  
CATHETER**Jul 20, 1989  
104 days to decisionK892410 · Product code: **DXE** · Cardiovascular  
Source: <https://www.510kdatabase.net/k892410/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Apr 7, 1989
Decision date	Jul 20, 1989
Days to decision	104 days
Third-party review	No

**APPLICANT**

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Company	<b>Baxter Healthcare Corp</b>
Location	Mchenry, IL, US
Contact	J KRATTENMAKER
510(k) history	505 submissions · 496 cleared · 1977-2019

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k892410/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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