

**K791489 VITAL CATHETER**Oct 26, 1979  
84 days to decisionK791489 · Product code: **FIE** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k791489/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Aug 3, 1979
Decision date	Oct 26, 1979
Days to decision	84 days
Third-party review	No

**APPLICANT**

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Company	<b>Hospal Medical Corp.</b>
Location	Mchenry, IL, US
510(k) history	55 submissions · 55 cleared · 1977-1989

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

Device record: <https://www.510kdatabase.net/k791489/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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