

**K780062 CATHETER/HEMOSTATIC/GERIATRIC**Feb 14, 1978  
33 days to decisionK780062 · Product code: **EZL** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k780062/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Retention Type, Balloon (EZL)
Date received	Jan 12, 1978
Decision date	Feb 14, 1978
Days to decision	33 days
Third-party review	No

**APPLICANT**

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Company	<b>Kendall Research Center</b>
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1976-1980

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

Device record: <https://www.510kdatabase.net/k780062/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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