

**K760174 ADAPTER, RESECTOSCOPE**Aug 30, 1976  
55 days to decisionK760174 · Product code: **FDC** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k760174/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resectoscope, Working Element (FDC)
Date received	Jul 6, 1976
Decision date	Aug 30, 1976
Days to decision	55 days
Third-party review	No

**APPLICANT**

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Company	<b>V. Mueller O.V. Baxter Healthcare Corp.</b>
Location	Mchenry, IL, US
510(k) history	41 submissions · 41 cleared · 1976-1981

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

Device record: <https://www.510kdatabase.net/k760174/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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