

**K760182 SHEATH, RESECTOSCOPE**Dec 2, 1976  
149 days to decisionK760182 · Product code: **FDC** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k760182/>**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	Dec 2, 1976
Days to decision	149 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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Device record: <https://www.510kdatabase.net/k760182/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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