

K760184 ELEMENT, WORKINGDec 2, 1976
149 days to decisionK760184 · Product code: **FDC** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k760184/>**SUBMISSION DETAILS**

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

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

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Device record: <https://www.510kdatabase.net/k760184/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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