

K780454 ACTIVE-LIFEApr 21, 1978
32 days to decisionK780454 · Product code: **EXB** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k780454/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Ostomy (EXB)
Date received	Mar 20, 1978
Decision date	Apr 21, 1978
Days to decision	32 days
Third-party review	No

APPLICANT

Company	E. R. Squibb & Sons, Inc.
Location	New York, NY, US
510(k) history	32 submissions · 32 cleared · 1977-1982

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Device record: <https://www.510kdatabase.net/k780454/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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