

K801702 SUR-FIT SYSTEM PRE-OPERATION SETAug 27, 1980
34 days to decisionK801702 · Product code: **EXB** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k801702/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Ostomy (EXB)
Date received	Jul 24, 1980
Decision date	Aug 27, 1980
Days to decision	34 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/k801702/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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