

**K800750 STOMAHESIVE STERILE WAFER**May 8, 1980  
34 days to decisionK800750 · Product code: **EXB** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k800750/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Ostomy (EXB)
Date received	Apr 4, 1980
Decision date	May 8, 1980
Days to decision	34 days
Third-party review	No

**APPLICANT**

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Company	<b>Squibb &amp; Sons, Inc.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1976-1982

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

Device record: <https://www.510kdatabase.net/k800750/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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