

**K802173 URIHESIVE STRIPS**Sep 26, 1980  
17 days to decisionK802173 · Product code: **KNX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k802173/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Sep 9, 1980
Decision date	Sep 26, 1980
Days to decision	17 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k802173/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 29, 2026