

**K213444 RELIEF Ureteral Stent Kit**Mar 17, 2022  
143 days to decisionK213444 · Product code: **FAD** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k213444/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Oct 25, 2021
Decision date	Mar 17, 2022
Days to decision	143 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	Model: RS-001 - 6 Fr x 24cm, RELIEF Ureteral Stent Kit; Model: RS-002 - 6 Fr x 26cm

**APPLICANT**

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Company	<b>Ureteral Stent Company</b>
Location	Chagrin Falls, OH, US
Contact	Mike Bunker
510(k) history	2 submissions · 2 cleared · 2022-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k213444/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 26, 2026