

K232920 RELIEF™ Ureteral Stent KitMar 22, 2024
185 days to decisionK232920 · Product code: **FAD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k232920/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Sep 19, 2023
Decision date	Mar 22, 2024
Days to decision	185 days
Third-party review	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

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k232920/> Data sourced from FDA 510(k) public records (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. - Generated May 13, 2026