FDA 510(k) Premarket Notification

# K232920 RELIEF™ Ureteral Stent Kit

**CLEARED**

Mar 22, 2024
185 days to decision

K232920 · Product code: **FAD** · Gastroenterology & Urology
Source: 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 |