

K883561 BARD(R) URETERAL STENT W/SUTURENov 15, 1988
88 days to decisionK883561 · Product code: **FAD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k883561/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Stent, Ureteral (FAD) |
| Date received | Aug 19, 1988 |
| Decision date | Nov 15, 1988 |
| Days to decision | 88 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |

APPLICANT

| | |
|----------------|---|
| Company | C.R. Bard, Inc. |
| Location | Covington, GA, US |
| Contact | DONNA J WILSON |
| Website | https://www.bd.com |
| 510(k) history | 645 submissions · 609 cleared · 1976-2026 |

C.R. Bard, Inc. is a developer, manufacturer, and marketer of medical technologies headquartered in Covington, US. The company specializes in vascular medicine, urology, oncology, and surgical specialty devices. C.R. Bard maintains a strong FDA 510(k) regulatory track record with FDA 510(k) cleared devices from total submissions spanning 1976 to 2026. The company's portfolio encompasses cardiovascular devices, gastroenterology and urology products, and general surgical technologies. Recent clearances include temporary pacing electrode catheters, thrombectomy systems, and ...
