FDA 510(k) Premarket Notification

**CLEARED**

# K021736 BARD VENTRALEX HERNIA PATCH, MODELS 0010302, 0010301

Jul 16, 2002
49 days to decision

K021736 · Product code: **FTL** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k021736/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Mesh, Surgical, Polymeric (FTL) |
| Date received | May 28, 2002 |
| Decision date | Jul 16, 2002 |
| Days to decision | 49 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **C.R. Bard, Inc.** |
| Location | Covington, GA, US |
| Contact | BRAIN A KANERVIKO |
| 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 ...