

**K233402 Bard 3DMax Mesh**Nov 1, 2023  
28 days to decisionK233402 · Product code: **FTL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k233402/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Oct 4, 2023
Decision date	Nov 1, 2023
Days to decision	28 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Davol Inc., Subsidiary of C. R. Bard, Inc.</b>
Location	Warwick, RI, US
Contact	Richard Spaziano
510(k) history	3 submissions · 3 cleared · 2023-2025

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k233402/> 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 25, 2026