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6 **IN THE UNITED STATES DISTRICT COURT**
7 **FOR THE DISTRICT OF ARIZONA**

8 IN RE: Bard Implanted Port Catheter
Products Liability Litigation

MDL No. 3081

9 THIS DOCUMENT RELATES TO:
10 **KAREN D. RICHARD**,

**FIRST AMENDED
MASTER SHORT-FORM
COMPLAINT AND JURY
TRIAL DEMAND**

11 _____,
12 Plaintiff(s),

Case 2:25-cv-04036-DGC

13 v.

14 Becton Dickinson and Company, et al.,
15 Defendants.
16

17 Plaintiff(s) named below, for their Complaint against Defendants named below,
18 incorporate(s) by reference the Second Amended Master Complaint in MDL 3081 (Doc.
19 1889). Pursuant to Case Management Order No. 7, as amended (Docs. 112, 145, 526,
20 2343), this Short-Form Complaint adopts the allegations, claims, and relief as set forth in
21 the Second Amended Master Complaint (Doc. 1889). As set forth below, Plaintiff(s) may
22 include (a) additional claims and allegations against Defendants, as set forth in Paragraph
23 15 or an additional sheet attached hereto; and/or (b) additional claims and allegations
24 against other Defendants, as set forth in Paragraph 5 or an additional sheet attached hereto.
25 Plaintiff(s) further allege(s) as follows:
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1 **I. PLAINTIFF(S)**

2 1. Name of Plaintiff/Decedent implanted with Bard Implanted Port Catheter
3 Product (“Device”) (first, middle, and last name):

4 Karen D. Richard

5 2. Name of Plaintiff/Decedent’s spouse (if bringing a loss-of-consortium claim):
6 N/A

7 3. Other Plaintiff and capacity (*i.e.*, administrator, executor, guardian, conservator,
8 representative, survivor, etc.), if any:

9 N/A

10 **II. DEFENDANT(S)**

11 4. Plaintiff(s) name(s) the following Defendant(s) in this action:

12 Becton, Dickinson and Company

13 C.R. Bard, Inc.

14 Bard Access Systems, Inc.

15 Bard Peripheral Vascular, Inc.

16 5. Plaintiff(s) contend(s) that additional parties may be liable or responsible for
17 Plaintiff(s)’ damages alleged herein. Such additional parties and their
18 citizenship are as follows:

19 _____
20 _____
21 _____

22 **III. JURISDICTION AND VENUE**

23 6. City and State of domicile of each Plaintiff at time of filing Plaintiff(s)’ initial
24 Complaint:

25 Wilmington, NC

26 7. City and State of residence of Plaintiff/Decedent at the time of Device
27 placement:

28 Wilmington, NC

1 8. City and State of residence of Plaintiff/Decedent at the time of alleged injury
2 for which a claim is asserted:

3 Wilmington, NC

4 9. Basis for jurisdiction:

5 Diversity of citizenship (28 U.S.C. § 1332(a))

6 Other: _____

7 a. Other allegations of jurisdiction and venue not expressed in Master
8 Complaint:

9 _____
10 _____

11 10. Designated forum (United States District Court and Division, if applicable) in
12 which Plaintiff asserts personal jurisdiction and venue would be proper absent
13 direct filing in this MDL:

14 United States District Court for the Eastern District of North Carolina

15 **IV. PRODUCT USE AND INJURY**

16 11. Plaintiff/Decedent was implanted with the following Device(s) and alleges that
17 the Device(s) caused their injuries¹:

18 BardPort M.R.I. Implantable Port

19 BardPort M.R.I. Low-Profile Implantable Port

20 BardPort Titanium Dome Implantable Port

21 BardPort Titanium Implantable Port

22 M.R.I. Plastic Dual Lumen Port

23 M.R.I. Ultra SlimPort Implantable Port

24 Peritoneal Titanium Port

25 PowerFlow Implantable Apheresis IV Port

26 PowerPort ClearVUE isp Implantable Port

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28 ¹ Check all that apply. See Exhibit A for additional information regarding the
corresponding model numbers/product codes for these Devices.

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- PowerPort ClearVUE Slim Implantable Port
- PowerPort duo M.R.I. Implantable Port
- PowerPort Implantable Port
- PowerPort isp Implantable Port
- PowerPort isp M.R.I. Implantable Port
- PowerPort M.R.I. Implantable Port
- PowerPort Slim Implantable Port
- PowerPort VUE M.R.I. Implantable Port
- PowerPort VUE Titanium Implantable Port
- SlimPort Dual-Lumen Rosenblatt Implantable Port
- Titanium Low-Profile Port
- Titanium SlimPort Implantable Port
- Vaccess CT Low-Profile Titanium Power-Injectable Port
- Vaccess CT Power-Injectable Implantable Port
- X-Port isp M.R.I. Implantable Port
- X-Port Low-Profile Titanium Port
- Other: _____

12. Date(s) of implantation as to the foregoing Device(s):

8/23/2023

13. Model number(s)/product code(s), if available, for the foregoing Device(s):

1708061

14. Complication(s) alleged to have occurred from use of the foregoing Device(s):

- Catheter fracture
- Infection
- Thrombosis
- Other: _____

1 **V. CAUSES OF ACTION**

2 15. Plaintiff(s) adopt(s) in this Short-Form Complaint the following claims and
3 allegations asserted in the Master Long-Form Complaint:

- 4 Count I: Design Defect – Strict Liability
- 5 Count II: Design Defect – Negligence
- 6 Count III: Failure to Warn/Instruct – Strict Liability
- 7 Count IV: Failure to Warn/Instruct – Negligence
- 8 Count V: Manufacturing Defect – Strict Liability
- 9 Count VI: Manufacturing Defect – Negligence
- 10 Count VII: Breach of Express Warranty
- 11 Count VIII: Breach of Implied Warranty
- 12 Count IX: Negligent Misrepresentation
- 13 Count X: Fraudulent Misrepresentation
- 14 Count XI: Fraudulent Concealment
- 15 Count XII: Consumer Fraud and/or Unfair and Deceptive Trade Practices
- 16 Count XIII: Unjust Enrichment
- 17 Count XIV: Loss of Consortium
- 18 Count XV: Wrongful Death
- 19 Count XVI: Survival
- 20 Timeliness and Tolling of Statutes of Limitation and Repose
- 21 Punitive Damages
- 22 Count XVII: Other _____

23 If additional claim(s) against Defendant(s) are alleged in Count XVII
24 above, the facts supporting such claim(s) must be pleaded. Plaintiff(s)
25 assert(s) the following factual allegations:

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16. Jury Trial demanded for all issues so triable?

Yes

No

WHEREFORE, Plaintiff(s) pray(s) for relief and judgment against Defendants and all such further relief that this Court deems equitable and just as set forth in the Master Long Form Complaint and Jury Demand and any additional relief to which Plaintiff(s) may be entitled.

Dated: 11/10/2025

Respectfully submitted,

/s/ David Kuttles

David Kuttles, Esq.

Admitted Pro Hac Vice

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EXHIBIT A

<u>Brand Name</u>	<u>Model Number/Product Code</u>
BardPort M.R.I. Implantable Port	0602610, 0602620, 0602640, 0602650, 0602660, 0602670, 0602680, 0602690, 0602830, 0602833, 0602840, 0602843, 0605400, 0605420, 0607173
BardPort M.R.I. Low-Profile Implantable Port	0603830, 0603840, 0603870, 0603880, 6603880
BardPort Titanium Dome Implantable Port	0602850, 0602860, 0602870
BardPort Titanium Implantable Port	0602230, 0602240, 0602270, 0602290, 0603000, 0602820, 0605300, 0605320, 0607301, 0607302, 0602210, 0602260, 0602280, 0602810
M.R.I. Plastic Dual Lumen Port	0603500, 0605920, 0605930, 0607100, 0607200, 0615460
M.R.I. Ultra SlimPort Implantable Port	0605640, 0655640
Peritoneal Titanium Port	0603000, 0603006
PowerFlow Implantable Apheresis IV Port	A710962
PowerPort ClearVUE isp Implantable Port	1606052, 1606062, 1606362, 1606382, 1608052, 1608062, 1608362, 1608382, 1666362, 1668362, 1676300, 5606362, 5608062, 5608362, 5666362, 5668362, CP00004

1 2 3 4 5 6	PowerPort ClearVUE Slim Implantable Port	1616000, 1616001, 1616070, 1616071, 1616300, 1616380, 1618000, 1618001, 1618070, 1618300, 1618380, 1676301, 1678300, 1678301, 5616000, 5616300, 5618000, 5618300, 5676300, 5676301, 5678300, 5678301, CP00005
7	PowerPort duo M.R.I. Implantable Port	1829500, 1829570, 5829500, 5829502
8 9 10	PowerPort Implantable Port	1708000, 1708001, 1708070, 1708071, 1709600, 1709601, 1759600, 1759601, 1778000, 1778001, 1778070, 1778071
11 12 13 14 15 16	PowerPort isp Implantable Port	1706050, 1706051, 1706060, 1706061, 1708050, 1708051, 1708060, 1708061, 1708160, 1708550, 1708551, 1708560, 1708561, 4708060, 4708061, 4708560, 4708561, CP00001, CP00002, CP00003, CP00009
17 18 19 20 21 22	PowerPort isp M.R.I. Implantable Port	1806050, 1806051, 1806060, 1806061, 1808050, 1808051, 1808060, 1808061, 1808069, 1808360, 1808550, 1808551, 1808560, 1808561, 1809660, 1809661, 1859660, 1859661, 4808060, 4808061, 4808560, 4808561, 9808560
23 24 25 26	PowerPort M.R.I. Implantable Port	1808000, 1808001, 1808002, 1808070, 1808071, 1808300, 1809600, 1809601, 1809670, 1859600, 1859601, 1878000, 1878001, 1878070, 1878071

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1		1716000, 1716001, 1716070, 1716071,
2	PowerPort Slim Implantable Port	1716080, 1718000, 1718001, 1718070,
3		1718500, 1718501, 1718570, 1718571,
4		CP00008
5	PowerPort VUE M.R.I. Implantable Port	1806052, 1806062, 1808052, 1808062
6	PowerPort VUE Titanium Implantable	
7	Port	1706052, 1706062, 1708052, 1708062
8	SlimPort Dual-Lumen Rosenblatt	
9	Implantable Port	0604970, 0624970, 0654970
10		
11	Titanium Low-Profile Port	0602180, 0602190, 0605490, 0605510,
12		0606100, 0606150, 0606200
13	Titanium SlimPort Implantable Port	0605550, 0605560, 0655510
14	Vaccess CT Low-Profile Titanium	
15	Power-Injectable Port	7360000, 7360001, 7380000
16	Vaccess CT Power-Injectable	
17	Implantable Port	7460000, 7480000, 7496000
18		
19	X-Port isp M.R.I. Implantable Port	0607500, 0607510, 0607520, 0607530,
20		0607540, 0607550, 0607555, 0657500,
21		0657510, 0657520, 0657525, 7707540,
22		7757540
23	X-Port Low-Profile Titanium Port	0655870, 0605840, 0605850