IN THE UNITED STATES DISTRICT COURT FOR THE DISTRICT OF ARIZONA

MDL No. 3081 In Re Bard Implanted Port Catheter Products Liability Litigation

In completing this <u>Plaintiff Profile Form</u>, you are under oath and must provide information that is true and correct to the best of your knowledge. The Plaintiff Profile Form shall be completed in accordance with the requirements set forth in the applicable Case Management Order.

1. CASE INFORMATION		
Caption:		Date:
Docket No.:		
		aformation, including email:
	2. PLAIN	TIFF INFORMATION
Full legal name of Pla Product ("Device"):	nintiff/Decedent im	planted with Bard Implanted Port Catheter
Former name:		
Address:		
Date of birth:		
Social security no. (la	st four digits only)):
Occupation:		
Spouse:		
Is Spouse making a cl	aim for loss of cor	nsortium?
Yes	No	
Representative name,	, if applicable:	
P oprosoptotivo rolotic	anchin to Dlaintiff	Dogodont:

3. DEVICE INFORMATION Name of Bard Implanted Port Catheter Product ("Device"): Model Number/Product Code: Lot Number: Provide the medical record, your medical alert card, or other documentation showing your **Device Product Code and Lot Number.** Medical records attached Medical alert card attached Other documentation showing Product Code and Lot Number attached Please check all the reasons why you believe your Device was implanted: **Blood draws Blood transfusions Chemotherapy delivery Immunotherapy delivery** IV fluid delivery IV antibiotics Parenteral nutrition Other – please describe below: Provide the name and address of the doctor who implanted the Device and the

hospital/medical facility at which the Device was implanted:

Doctor:	
Hospital/Medical Facility:	

Provide medical records for the implant of the Device.

Medical Records attached

*NOTE: If you are alleging injuries related to more than one Device, complete Sections 3-8 for each Device and attach additional pages as needed.

4. FAILURE MODE ALLEGED

Please check all failure mode(s) that you allege apply to the Device and attach medical records that show the failure mode:

CATHETER-RELATED FAILURE MODES:

Catheter-related infection
Type of infection:
Thrombosis in or around catheter
Occlusion of the catheter
Fracture of catheter without migration of a fragment
Fracture of catheter with migration of a fragment to(state location in your body)
Other – state in detail:
None (not making a catheter-related claim)

For each catheter-related complication identified above, state the date you were first diagnosed with such complication and state the name of the medical provider who diagnosed and/or treated the complication:

For each catheter-related complication identified above, provide medical records relating to the first diagnosis of each complication.

Medical records attached

PORT-BODY/RESERVOIR-RELATED FAILURE MODES:

Port-body/reservoir-related infection
Type of infection:
Thrombosis of port body/reservoir
Occlusion of port body/reservoir
Erosion or wound complications at the port-body site
Other – state in detail:
None (not making a port-body/reservoir-related claim)
For each port-body/reservoir-related complication identified above, state the date you were first diagnosed with such complication and state the name of the medical provider who diagnosed and/or treated the complication:
For each port-body/reservoir-related complication identified above, provide medical records relating to the first diagnosis of each complication.

Medical records attached

5. REMOVAL INFORMATION

* This Section is limited to removal of the Device as a whole. Information regarding fractures and removal of fracture remnants should be provided in Section 7.

Has yo	our Device identifie	d in Section 3 been removed?
	Yes	No
		name(s) and address(es) of the doctor(s) who removed your pital/medical facility where the removal/attempted removal
	Doctor:	
	Hospital/Medical l	acility:
	Date of removal: _	
Provid	le medical records f	or the removal/attempted removal and the procedure involved.
	Medical record	s attached
Was th	ne Device identified	in Section 3 preserved after removal?
	Yes	No
	,	me and address of the person or institution in possession of the
Do you	ı have photographs	and/or video of the removed Device or of the removal procedure?
	Yes photograp	ns. If yes, produce color copies of the photos.
	Photographs at	tached
	Yes video. If y	es, retain the video.
	No	

6. SUBSEQUENT DEVICE If your Device identified in Section 3 was removed, was a subsequent device implanted? No Yes. State date of implant of replacement device: Was it replaced with a Bard Port Catheter Device? If yes, provide: Product Name: ____ Product Code: Lot Number: If no, provide the name of replacement device: 7. CATHETER FRAGMENTS Do you claim that the catheter of your Device fractured? Yes No If you answered YES, answer the below questions in this Section. If you answered NO, skip the rest of Section 7 and go below to Section 8 - "Outcome Attributed to Device."

Are any catheter fragments retained in your body?

Have any catheter fragments been removed from your body?

Yes

No

Yes

No

Unknown

Unknown

If yes, identify the location(s) within your body of each retained catheter fragment.

If any catheter fragment has been removed (or a doctor has attempted to remove it), please check all that apply regarding the removal procedure(s):

Removed percutaneously

Removed via open-chest procedure

Removed via alternative open procedure

Attempted but unsuccessful removal percutaneously

Attempted but unsuccessful removal via open-chest procedure

Attempted but unsuccessful removal via alternative open procedure

If any catheter fragment has been removed or if there has been an attempt to remove, state the following for each removal/attempt:

Doctor:	
Hospital/Medical Facility:	
Date:	
Doctor:	
Date:	
Doctor:	
Hospital/Medical Facility:	
Date:	

Provide medical records that provide the date(s) of removal (or attempted removal), the location (in your body) of the fractured fragments, and the procedure(s) performed to remove (or attempt to remove) the fragments.

Medical records attached

Do you have photographs and/or video of the removed Device or fragments or of the removal procedure?

Yes photographs. If yes, produce color copies of the photos.

Photographs attached

Yes video. If yes, retain the video.

No

8. OUTCOME ATTRIBUTED TO DEVICE

·	nat you are currently suffering from any bodily uries related to the Device identified in Section 3:
Yes	
No	
If your answer is "Yes," please list a and describe the medical treatment	ll symptoms and injuries you claim to have suffered received to address them:
Of the injuries/symptoms you listed the current time:	above, which do you claim to be suffering from at

Plaintiff reserves the right to supplen additional information.	nent any and all responses upon the receipt of
I declare under penalty of perjury that	the information in this Plaintiff Profile Form is correct:
Date	Signature of Plaintiff
Date	Signature of Plaintiff's Spouse (signature necessary only if loss of consortium is alleged)

THIS PROFILE FORM AND THE RECORDS SHOULD BE UPLOADED TO WWW.MDLCENTRALITY.COM/BARDPORT PURSUANT TO CMO NO. 8.