#### UNITED STATES DISTRICT COURT NORTHERN DISTRICT OF INDIANA SOUTH BEND DIVISION

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## IN RE: BIOMET M2a MAGNUM HIP IMPLANT PRODUCT LIABILITY LITIGATION

(MDL 2391)

This Document Relates to All Cases

Cause No. 3:12-MD-2391-RLM-CAN
Judge Robert L. Miller, Jr.

It is hereby ordered that any Plaintiff with a case pending in the above referenced litigation shall complete and serve a Preliminary Disclosure Form, attached as Exhibit A to this Order, within thirty (30) days of the entry of this Order. Service shall be made on Defendants' liaison counsel, Ryan C. Edwards, Taft Stettinius & Hollister LLP, by sending the form electronically by email to <u>BiometPFS@taftlaw.com</u> and a curtsey copy should be served on Plaintiffs' liaison counsel, Douglas A. Kreis, Aylstock, Witkin, Kreis & Overholtz, PLLC, by sending it electronically by email to <u>BiometPDF@awkolaw.com</u>.

For all cases filed after the date of this Order, the Preliminary Disclosure Form is to be served thirty (30) days from the date that the case is transferred into the MDL, or, if directly filed in the MDL, thirty (30) days from the date that the case is filed.

This Preliminary Disclosure Form must be completed and served regardless of whether a Plaintiff has previously served a Plaintiff Fact Sheet – it is a separate and independent form and must be completed. This Form is not a verified discovery response and is not evidence, but is designed to obtain information the Court finds necessary to assess the need for future discovery.

All Plaintiffs have a continuing duty to serve an updated form.

### IT IS SO ORDERED.

### ENTERED: January 7, 2014

/s/ Robert L. Miller, Jr. Robert L. Miller, Jr. Judge, United States District Court

# Exhibit A

### **BIOMET PRELIMINARY DISCLOSURE FORM** DO NOT FILE THIS FORM WITH THE COURT

## 1. Plaintiff Information

	(a)	Name:					
			First		Middle		Last
	(b)	DOB:					
	(0)	DOB	(Please form	at as MM/DD/YYY	<i>'</i> )		
	(c)	Address	:				
	( )		Street				
			City		<u></u>	ate (Abbreviation)	Zip Code
			/			,	P
	(d)	Venue:	(Please lis	t two letter state abb	reviation follow	wed by judicial distric	t and division )
			(i lease lis				
	(e)	Date File	ed:				
	( )			rmat as MM/DD/Y	(YY)		
	(f)	Attorne	y:				
2.	Im	plant In	<u>formation</u>				
	(a)	Locatior	n of Implant:	Right	Left	Bilateral	
	(-)		-	0			
(b) Implant date (Right):							
(Please format as MM/DD/YYYY) i. Hospital:							
		ii. Surg	geon:				
	(c)	Implant date (Left):					
(Please format as MM/DD/YYYY)							
		i. Hos	pital:				
			·				
		ii. Surg	geon:				

## 3. <u>Revision Information:</u>

(a)	Location of revision: Right Left Bilateral			
(b)	Revision date (Right): (Please format as MM/DD/YYYY)			
	i. Hospital:			
	ii. Surgeon:			
(c)	Revision date (Left): (Please format as MM/DD/YYYY)			
	i. Hospital:			
	ii. Surgeon:			
(d) If you have not had a revision yet, but one is scheduled, provide the date:(MM				
(e)	Has your doctor recommended a revision or re-revision but also advised you that this surgery is medically contraindicated and/or would be life threatening? Yes No			
	If so, identify the name and address of the doctor, date of the discussion, and the medical which prevents you from having the surgery and state whether you have been advised that conal will permanently prevent you from having revision surgery, as opposed to delaying a gery.			
	Date(s) of Discussion (MM/DD/YYY):			

Doctor:
Address:
Medical condition:

## 4. <u>Post-Revision Surgery</u>

	Do you claim that your revision surgery led to any of the followin	g: Yes No	
		<u>DATE</u> (MM/DD/YYYY)	
	(a) A second revision		
	(b) A third revision		
	(c) A forth revision		
	(d) Death		
	(e) Heart attack		
	(f) Stroke		
	(g) Pulmonary embolism		
	(h) Deep vein thrombosis/blood clot		
	(i) Fracture (femoral shaft or Trochanteric)		
	(j) Dislocation(s): Yes No Number of dislo	cations:	
	Please provide the DATE(S) of any dislocations (MM/DD/YYYY):		
	(k) Infection(s): Yes No		
	Please provide the DATE(S) of any infections (MM/DD/YYYY):		
lf you m	arked yes for infection, please check the box for any and all of th	e following treatment receiv	ved:
IV antibi	iotic treatment Antibiotic spacers Irrigation &	Debridement	
	<ul> <li>Permanent and full time use of a wheel chair or walker for ar revision surgery): Yes No</li> </ul>	nbulation (not used prior to	)
	(m) Foot drop: Yes No		
Note:	Only mark yes if this is documented in the medical records after r	revision surgery.	
	If you marked yes for foot drop, please identify the treatment red	ceived or recommended:	

(n)	Peripheral neuropathies or nerve damage (with objective EMG evidence):						
	DATE:						
	(MM/DD/YYYY)						

(o) Other Extreme Conditions/loss: Yes No

Explain:

#### 5. <u>Other Surger(ies)</u>

Have you had any other surgery post-revision (not already identified above) that you claim is related to the implant? Only answer yes if you have undergone surgery. Do not answer yes if you have only received injections. Yes No

Please state the condition treated:

Please provide the DATE(S) of any additional surgery(ies) (MM/DD/YYYY):

To the extent that you have not already provided authorizations with a previously submitted Plaintiff Fact Sheet (PFS), provide signed authorizations for any doctor or medical provider who has treated you for any condition identified in Question 4 above.