

## ORANGE CLAIM FORM FOR PART A BASE AWARD

The Claims Package and Required Submissions, including this Orange Claim Form, must be submitted no later than May 5, 2017, on behalf of all Enrolled Program Claimants, including Unrepresented (*pro se*) Enrolled Program Claimants in the U.S. Program outlined in the Settlement Agreement of March 2, 2015 (“the 2015 Agreement”) and the 2017 ASR Extension Agreement dated March 3, 2017, (“the 2017 Extension Agreement”) (generally and collectively referred to herein as the “U.S. Program”). Any references to the 2015 Agreement in this Orange Claim Form incorporate the terms detailed in the 2017 Extension Agreement in addition to those of the 2015 Agreement.

If a Claimant has had an ASR Revision Surgery in both hips, the Claimant must submit this Orange Claim Form and a Red Claim Form for Bilateral benefits. If a Claimant has had a revision of more than one ASR Hip Implant in a single hip (Re-Revision) or otherwise qualifies for compensation from the Extraordinary Injury Fund, the Claimant must submit this Orange Claim Form and also a Green Claim Form. All Claimants must also submit a Blue Claim Form for lien resolution to: (1) identify any liens, claims, interests or requests for reimbursement that are allegedly related to an ASR or ASR Revision Surgery, or (2) state that they are aware of no such liens or claims.

### INSTRUCTIONS

1. Counsel for Claimants, and all *pro se* Claimants, must complete this Claim Form.
2. A “Claimant” as referred to in this Claim Form means the individual submitting a claim in the U.S. Program, who is either the Product User or the Legal Representative, as defined in 1.2.43 of the 2015 Agreement.

### A. PERSONAL INFORMATION OF PRODUCT USER

<b>1. Name</b>	Last	First	Middle Initial
<b>2. Current Address</b>	Street		
	City	State	Zip Country
<b>3. Date Began Residing at this Address</b>	____/____/____ (MM/DD/YYYY)	<input type="checkbox"/> <b>Product User has resided at this same address since his/her ASR Index Surgery</b>	
<b>4. Telephone Number</b>	(____) ____-____	<b>5. Date of Birth</b>	____/____/____ (MM/DD/YYYY)
<b>6. Social Security Number</b>	____-____-____	<b>7. Gender</b>	<input type="checkbox"/> Male <input type="checkbox"/> Female
<b>8. Any Other Names Used or by which the Product User has been known, including but not limited to maiden name:</b>			
<b>9. Was the Product User a citizen or legal resident of the United States at the time of the Index Surgery to implant the ASR Hip Implant(s)?</b>		<input type="checkbox"/> Yes <input type="checkbox"/> No	

### B. LEGAL REPRESENTATIVE’S INFORMATION FOR DECEASED OR INCAPACITATED PRODUCT USERS (COURT APPROVAL OR OTHER AUTHORIZATION TO REPRESENT THE PRODUCT USER MUST BE ATTACHED)

<b>10. Does the Product User have a Legal Representative?</b>	<input type="checkbox"/> Yes <input type="checkbox"/> No	If Yes, complete Items 11-17. If No, skip to Section C.
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<b>11. Reason for Legal Representative</b>	<input type="checkbox"/> Product User is Deceased <input type="checkbox"/> Product User is Incompetent		
<b>12. Legal Representative's Relationship to Product User</b>	<input type="checkbox"/> Estate <input type="checkbox"/> Executor <input type="checkbox"/> Administrator <input type="checkbox"/> Guardian <input type="checkbox"/> Conservator <input type="checkbox"/> Other _____ <div style="text-align: right;">(specify)</div>		
<b>13. Legal Representative's Name</b>	Last	First	Middle Initial
<b>14. Legal Representative's Address</b>	Street		
	City	State	Zip                      Country
<b>15. Legal Representative's Social Security Number</b>	-         -		
<b>16. Date of Death of Product User (if applicable)</b>	____/____/____ (MM/DD/YYYY)		
<b>17. Do you claim the ASR Revision Surgery caused the death?</b>	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A		
<b>C. PRIMARY LAW FIRM INFORMATION (if represented by an attorney)</b>			
<b>18. Principal Responsible Attorney</b>	Last	First	Middle Initial
<b>19. Firm Name</b>			
<b>20. Current Address</b>	Street		
	City	State	Zip
<b>21. Telephone Number</b>	(       )           -		
<b>22. Fax Number</b>	(       )           -		
<b>23. Email Address</b>			
<b>24. Date of Retention Agreement with Claimant/Plaintiff</b>	____/____/____ (MM/DD/YYYY)		



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42. Select the reason the Product User is no longer married.

- Divorced     
  Death of Former Spouse     
  Death of Product User

### F. BASE AWARD CLAIM INFORMATION

**Check the ASR Hip Implants and Other Circumstances that apply to the Product User's use of ASR Hip Implants and indicate the date(s) of occurrence.**

If a Claimant has had a revision of an ASR in both hips and is submitting claims for both hips under the 2015 Agreement, the Claimant must submit this Orange Claim Form for the first hip revised and a Red Claim Form for Bilateral benefits. If a Claimant has had a revision of an ASR in both hips and submitted a claim for one of those under the 2013 ASR Master Settlement Agreement, the Claimant must submit this Orange Claim Form for the second hip revised. If a Claimant has had a revision of more than one ASR in a single hip, the Claimant must submit this Orange Claim Form for the first revision and a Green Claim Form for Re-Revision benefits.

#### LEFT HIP

43. Indicate the Product Implanted into the Product User

- Total Hip Replacement with ASR XL Hip Implant  
 ASR Hip Resurfacing Implant

44. Date of Index Surgery

\_\_\_\_/\_\_\_\_/\_\_\_\_  
(MM/DD/YYYY)

45. Location of Hospital Where Index Surgery Occurred

- Hospital Located in the U.S.  
 Military Hospital Located Outside of the U.S.  
 Non-Military Hospital Located Outside of the U.S.

46. Name of Hospital Where Index Surgery Occurred

47. Name of Index Surgery Surgeon

Last

First

Middle Initial

48. Did the Product User undergo a Revision Surgery involving the Left ASR Hip Implant?

- Yes       No

If Yes, complete Items 49 – 52.  
If No, skip to Item 53.

49. Date of Revision Surgery

\_\_\_\_/\_\_\_\_/\_\_\_\_  
(MM/DD/YYYY)

50. Location of Hospital Where Revision Surgery Occurred

- Hospital Located in the U.S.  
 Military Hospital Located Outside of the U.S.  
 Non-Military Hospital Located Outside of the U.S.

51. Name of Hospital Where Revision Surgery Occurred

52. Name of Revision Surgery Surgeon

Last

First

Middle Initial

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RIGHT HIP			
<b>53. Indicate the Product Implanted into the Product User</b>	<input type="checkbox"/> <b>Total Hip Replacement with ASR XL Hip Implant</b> <input type="checkbox"/> <b>ASR Hip Resurfacing Implant</b>		
<b>54. Date of Index Surgery</b>	_____ (MM/DD/YYYY)	<b>55. Location of Hospital Where Index Surgery Occurred</b>	<input type="checkbox"/> Hospital Located in the U.S. <input type="checkbox"/> Military Hospital Located Outside of the U.S. <input type="checkbox"/> Non-Military Hospital Located Outside of the U.S.
<b>56. Name of Hospital Where Index Surgery Occurred</b>	_____		
<b>57. Name of Index Surgery Surgeon</b>	<small>Last</small> _____	<small>First</small> _____	<small>Middle Initial</small> _____
<b>58. Did the Product User undergo a Revision Surgery involving the Right ASR Hip Implant?</b>		<input type="checkbox"/> Yes <input type="checkbox"/> No    If Yes, complete Items 59 – 62. If No, skip to Section G.	
<b>59. Date of Revision Surgery</b>	_____ (MM/DD/YYYY)	<b>60. Location of Hospital Where Revision Surgery Occurred</b>	<input type="checkbox"/> Hospital Located in the U.S. <input type="checkbox"/> Military Hospital Located Outside of the U.S. <input type="checkbox"/> Non-Military Hospital Located Outside of the U.S.
<b>61. Name of Hospital Where Revision Surgery Occurred</b>	_____		
<b>62. Name of Revision Surgery Surgeon</b>	<small>Last</small> _____	<small>First</small> _____	<small>Middle Initial</small> _____
G. BANKRUPTCY INFORMATION			
<b>63. Has the Product User at any time since the date of the ASR Index Surgery been party to a bankruptcy action in which he/she is seeking bankruptcy protection?</b>		<input type="checkbox"/> Yes <input type="checkbox"/> No    If Yes, complete Items 64 – 68. If No, skip to Section H.	
<b>64. Bankruptcy Court/ Jurisdiction</b>	_____		
<b>65. Case Number</b>	_____	<b>66. Date Filed</b>	_____ (MM/DD/YYYY)

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<b>67. Trust Name (If Trustee appointed.)</b>	
<b>68. Status of Bankruptcy Filing</b>	<input type="checkbox"/> Open <input type="checkbox"/> Closed (If closed, provide the date closed.) _____/_____/_____ <span style="font-size: small;">(MM/DD/YYYY)</span>

### H. REQUIRED SUBMISSIONS

You must submit all materials required by Section 4.1.3 of the 2015 Agreement:

- Enrollment Form.
- Release.
- Dismissal with Prejudice Stipulation (if applicable).
- This Orange Claim Form.
- Manufacturer/product stickers for the Qualifying ASR, identifying Product and Lot Codes for the device implanted into the Product User.
- A true and correct copy of all Contemporaneous Medical Records identifying the ASR XL Acetabular Hip System, ASR 300 Acetabular Cup System, or ASR Resurfacing System that was surgically implanted in the Product User in an ASR Index Surgery and removed during a Revision Surgery. This includes all records in your possession and obtained as a result of ordering the records.
- A true and correct copy of the Contemporaneous Medical Records, including Admission Records (including History and Physical Examination Records), Discharge Summaries, and Operative Reports pertaining to any ASR Index Surgery and ASR Revision Surgery.
- A medical record showing your weight and height at your ASR Index Surgery and your smoking status at the time of your ASR Revision Surgery.

### I. CERTIFICATION BY CLAIMANT

I declare under penalty of perjury under 28 U.S.C. §1746 that all of the information provided in and with this Claim Form is true and correct to the best of my knowledge, information and belief.

I further certify that by participating in this U.S. Program, I agree to abide by the terms of the 2015 Agreement, and further understand that by enrolling in the Settlement Program, I agree to be bound by the terms of MDL Case Management Order 13, as amended, which permits a holdback of 5% fees and 1% costs to be deducted from any final award/gross recovery to me from the U.S. Program which shall be used, in part, for the funding of the administration of the U.S. Program. I further agree to comply with any Orders entered by the United States District Court for the Northern District of Ohio (MDL Docket No. 1:10-md-2197) in the furtherance of Case Management Order 13, and consent to the jurisdiction of that MDL Court for that purpose. I further grant and convey to the Settlement Oversight Committee for MDL 2197 a lien upon and/or security interest for such holdback amounts in any recovery by me from the U.S. Program. If I qualify for a settlement award payment pursuant to the terms of the 2015 Agreement, I authorize such settlement payment to be made to my Counsel identified as my Primary Law Firm in trust for me in accordance with the 2015 Agreement.

<b>Claimant's Signature</b>		<b>Date</b>	_____/_____/_____ <span style="font-size: small;">(MM/DD/YYYY)</span>
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<b>Printed Name</b>	First	Middle Initial	Last
<b>J. COUNSEL SIGNATURE</b>			
<b>Counsel's Signature</b>			<b>Date</b> ____/____/____ (MM/DD/YYYY)
<b>Printed Name</b>	First	Middle Initial	Last

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