

PILOT PARTUM: End of Study

Site No.

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Subject No.

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## End of Study Case Report Form

A. Study Completion										
1. Date of study termination:	<table border="1"> <tr> <td>D</td><td>D</td><td>M</td><td>M</td><td>M</td><td>Y</td><td>Y</td><td>Y</td><td>Y</td> </tr> </table>	D	D	M	M	M	Y	Y	Y	Y
D	D	M	M	M	Y	Y	Y	Y		
2. Reason for study termination:	<input type="checkbox"/> Routine study termination, study protocol completed <input type="checkbox"/> Early study termination, due to: <ul style="list-style-type: none"> <li><input type="checkbox"/> Death*</li> <li><input type="checkbox"/> Withdrawal of subject's consent**:               <ul style="list-style-type: none"> <li><input type="checkbox"/> Subject allows data collection to continue</li> <li><input type="checkbox"/> Subject refuses further data collection</li> </ul> </li> <li><input type="checkbox"/> Other, please specify: _____</li> </ul> <p><b>*If selected, please complete Death Outcome Event form and SAE form</b></p> <p><b>**Reason(s) subject has withdrawn consent:</b></p>									

B. Suspected Secondary Outcome Events	
1. Did the subject have one or more of the outcome events listed below (check all that apply)*:	<input type="checkbox"/> None <input type="checkbox"/> Symptomatic venous thromboembolism <input type="checkbox"/> Bleeding <input type="checkbox"/> Death* <input type="checkbox"/> Symptomatic arterial thromboembolism <input type="checkbox"/> Postpartum pre-eclampsia <p><b>*If yes, please ensure corresponding Outcome Event and SAE form(s) are completed.</b></p>

Delegate's Name:									
Signature:									
I have reviewed all entries on the Case Report Forms. All information entered onto the Case Report Form for this subject is, to the best of my knowledge, correct.									
Investigator's Name:									
Signature:									
Date: <table border="1"> <tr> <td>D</td><td>D</td><td>M</td><td>M</td><td>M</td><td>Y</td><td>Y</td><td>Y</td><td>Y</td> </tr> </table>	D	D	M	M	M	Y	Y	Y	Y
D	D	M	M	M	Y	Y	Y	Y	