Site No.

Subject No.

End of Study Case Report Form

1.	dy Comp	etion												
	Date of stu	dy term	ination:	D D M M	D D M M M Y Y Y									
2.	Reason for	study to	ermination:											
	D Routin	ne study	termination, study protoc	ol completed										
	□ Early	study ter	rmination, due to:											
		Lost to	follow up											
		Death*												
		Withdra	awal of subject's consent*	·*·										
			Subject allows data collec	tion to continue										
			Subject refuses further day	a collection										
		Other, J	please specify:											
:	*If selected, please complete Death Outcome Form and SAE form													
:	**Reason(s	s) subjec	et has withdrawn consen	t:										
D C			D-4											
B. Suspected Secondary Outcome Events														
			ve one or more suspected ck all that apply)*:	l outcome events liste	ed b	elow that will undergo								
	- Jan		en an enac appig)											
	None		Symptomatic venous thr	omboembolism		Bleeding/Hematoma								
_	None Death*		Symptomatic venous thr Symptomatic arterial thr			Bleeding/Hematoma Postpartum pre-eclampsia								
	Death*		•	omboembolism		Postpartum pre-eclampsia								
	Death*		Symptomatic arterial thr	omboembolism		Postpartum pre-eclampsia								
□ □ *If	Death*		Symptomatic arterial thr	omboembolism		Postpartum pre-eclampsia								
□ □ *If	Death* yes, please te's Name:		Symptomatic arterial thr	omboembolism		Postpartum pre-eclampsia								
□ ×If Delegat Signatu I have r	Death* yes, please te's Name: are: reviewed all	ensure	Symptomatic arterial thr corresponding Outcome	omboembolism Event and SAE forn	□ n(s)	Postpartum pre-eclampsia								

Signature:

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Date:	D	D	M	M	M	Y	Y	Y	Y