

Original source should be obtained from a direct laboratory report from Subject Medical Record. Laboratory results to be reviewed by delegated investigator (either directly on lab report or as Source here) as relates to subject safety and general INC/EXC Criteria.

Pre-procedure oral anticoagulation should be managed as per site protocol. Warfarin should be discontinued in accordance with site standard of care practices including INR levels on the day of the procedure. We are not collecting day of procedure INR levels.

Laboratory Collection at Procedure required **within 48 hours of implant**.

Date of Hematology	____ / ____ / ____ (DD/MMM/YYYY)
	<input type="checkbox"/> Not Done (ENTER PD)

<i>Laboratory Assessment</i>	<i>Results Value/ unit</i>	<i>Clinically Significant</i>
Hemoglobin		<input type="checkbox"/> Yes <input type="checkbox"/> No
Hematocrit		<input type="checkbox"/> Yes <input type="checkbox"/> No
Platelet Count		<input type="checkbox"/> Yes <input type="checkbox"/> No

If utilizing this form as source (i.e., no other source exists), this form should be signed by Site Investigator.

_____ Site Personnel Signature	____ / ____ / ____ Date (DD/MMM/YYYY)
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