

*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 INC/EXC Criteria.*

*Laboratory Collection at screening collected per standard of care **up to 60 days prior to consent***

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

Laboratory Assessment	Results	Clinically Significant
Hemoglobin		<input type="checkbox"/> Yes <input type="checkbox"/> No
Henatocrit		<input type="checkbox"/> Yes <input type="checkbox"/> No
WBC		<input type="checkbox"/> Yes <input type="checkbox"/> No
Platelet Count		<input type="checkbox"/> Yes <input type="checkbox"/> No

CHEMISTRY – SERUM CREATININE OR GFR eGFR

Date of serum Chemistry	____/____/____ (DD/MMM/YYYY)
	<input type="checkbox"/> Not Done (ENTER PD if neither Cr or GFR/eGFR were not obtained)

Laboratory Assessment	Results	Clinically Significant
Creatinine		<input type="checkbox"/> Yes <input type="checkbox"/> No
GFR or eGFR		<input type="checkbox"/> Yes <input type="checkbox"/> No

COAGULATION as Relevant

Was INR sample collected?	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Not applicable
Date of INR	____ / ____ / ____ (DD/MMM/YYYY)

<i>Laboratory Assessment</i>	<i>Results</i>	<i>Clinically Significant</i>
INR		<input type="checkbox"/> Yes <input type="checkbox"/> No

_____ Site Personnel Signature	____ / ____ / ____ Date (DD/MMM/YYYY)
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Reminder: Pre-procedure oral anticoagulation (Warfarin or DOAC) should be managed as per site protocol. Warfarin should be discontinued in accordance with site standard of care practices including the monitoring of INR levels on the day of the procedure.