For PDF Fillable Requisitions, the following applies:

- 1. The form shall be completed using a Digital Health assigned computer.
- 2. Absolutely no personal health information shall be electronically saved on a computer.
- 3. The completed form shall not be shared electronically. If you reasonably believe that e-mailing the information is the only available method of communication or the only way to send the information then you must adhere to the Privacy guideline titled "E-mailing Personal Health Information".
- 4. All forms must be completed in their entirety, e.g. if a staff member has only completed half of a form they cannot save their work and then come back to complete it at a later date.
- 5. Once the personal health information has been recorded onto the form, it is to be printed immediately, deleted (not saved) from the computer, and then stored securely inside the client (paper) health record or scanned into an electronic record.
- 6. Do not print unnecessary duplicate copies of the form.
- 7. Regular audits of the Digital Health assigned computer shall be undertaken to ensure that no personal health information is being duplicated and saved.

Ship samples to:

St/ Boniface Hospital Hematology Lab

L4006-409 Tache Ave Winnipeg, MB R2H 2A6 Phone: 204-237-2468 Fax: 204-237-2494

HIT Screen Heparin Induced Thrombocytopenia Screening Requisition

This space for lab use only Place DELPHIC Label here

Acceptance Policy 10-50-03: Requirements for Test Requisitions 2.1 - Fields marked with * are mandatory and must be clearly legible or can result in specimen rejection.					
ORDERING PROVIDER INFORMATION		PATIENT INFORMATION			
*Last & Full First Name:		*Last/First Name: (per MB Health Card)			
Billing Code:	Inpatient Location:	* Date of Birth (dd/mm/yyyy)			
*Facility Name/Address		*Sex: Female Male			
Phone No: Fax No:		*PHIN:			
Critical Results Phone Number:		*Specify Province or DND if different			
COPY REPORT TO: (if info missing, report may not be sent)		MRN:			
Last & Full First Name:	Fax No:	Encounter Number:			
Facility Name/Address: Phone No:		Patient Phone Number:			
Last & Full First Name: Fax No:		Patient Address:			
Facility Name/Address:	Phone No:	Demographics verified: ☐ Health Card ☐ Armband ☐ eChart/CR ☐ Other			
Collection Information Fields marked with "*" required by person collecting sample					
♦ Collector:	♦ Collection Date:	Collection: ☐ Venipuncture ☐ Capillary ☐ Indwelling Line			
♦ Collection Facility/Lab:	♦ Collection Time:	☐ Above shut off IV			
# Serum vial(s) # Plasma vials(p) Referring Lab: # of tube		s sent Samples shipped frozen 📮			
Last & Full First Name: Facility Name/Address: Collection Information Fields marked with Collection Facility/Lab: Collection Time:		Patient Address: Demographics verified: □ Health Card □ Armband □ eChart/CR □ Other with "*required by person collecting sample **Collection: □ Venipuncture □ Capillary □ Indwelling Line □ Above shut off IV			

**** Testing WIL	L NOT be i	nitiated if the section BELOV	V is not completed in ENTIRE	TY ****	
				Points Assigned	Points Given
Thrombocytopenia	Platelet count fall > 50% and platelet nadir ≥ 20 x 10 ⁹ /L		2		
	Platelet count fall 30-50% or platelet nadir 10-19 x 10 ⁹ /L			1	
	Platelet count fall < 30% or platelet nadir < 10 x 10 ⁹ /L			0	
Timing * of platelet count fall * 1 st day of heparin exposure = Day 0	Clear onset between days $5 - 14$ or platelet fall ≤ 1 day (prior heparin exposure within 30 days)			2	
	Consistent with days 5 – 14 fall, but not clear (e.g. missing platelet counts) or			1	
	onset after day 14 or fall ≤ 1 day (prior heparin exposure 30 – 100 days ago)				
	Platelet count fall ≤ 4 days without recent exposure			0	
Thrombosis or other sequelae (e.g. skin lesions)	New thrombosis (confirmed); skin necrosis at heparin injection sites; anaphylactoid reaction after IV heparin bolus; adrenal hemorrhage		2		
	Progressive or recurrent thrombosis; non-necrotizing (erythematous) skin lesions; suspected thrombosis (not confirmed)			ns; 1	
	None			0	
Other causes of	None apparent			2	
thrombocytopenia	Possible			1	
	Definite			0	
6 – 8 = High pre-test probability TOTAL PRE-TEST PROBABIL					
4 – 5 = Intermediate pre-test probability					
0 – 3 = Low pre-test probability				(0-8)	
TEST		SAMPLES REQUIRED		TEST COL	DE
☐ HIT Screen Note: Positive HIT Screened specimens will be referred out for confirmatory SRA (Serotonin Release Assay)		Serum for Heparin Induced Thrombocytopenia HIT Adult: 1 x 10 mL tube (serum) – red top 1 x 1.8 mL sodium citrate tubes (plasma) – light blue		НІТА	

