

## TRAINING UPDATE

<b>Lab Location:</b>	GEC, SGMC & WAH	<b>Date Distributed:</b>	8/3/2015
<b>Department:</b>	Core & Processing	<b>Due Date:</b>	8/31/2015
		<b>Implementation:</b>	<b>9/1/2015</b>

### DESCRIPTION OF PROCEDURE REVISION

**Name of procedure:**

**Hemolysis, Icteria and Lipemia Interference  
GEC.L04, SGAH / WAH.L05 v3**

**HIL Index Alert Values AG.F247.1**

**Description of change(s):**

Section 5: refer to related documents in step 7

Section 6: updated SOP list, moved HIL Index from section 9 and updated

This revised SOP will be implemented on September 1, 2015

Document your compliance with this training update by taking the quiz in the MTS system.

**Approved draft for training (version 3)**

Non-Technical SOP

Title	<b>Hemolysis, Icteria and Lipemia Interference</b>	
<b>Prepared by</b>	Hannah Tran	Date: 12/8/2008
<b>Owner</b>	Robert SanLuis	Date: 4/1/2013

<b>Laboratory Approval</b>		
<b>Print Name and Title</b>	<b>Signature</b>	<b>Date</b>
<i>Refer to the electronic signature page for approval and approval dates.</i>		
<b>Local Issue Date:</b>	<b>Local Effective Date:</b>	

<b>Review:</b>		
<b>Print Name</b>	<b>Signature</b>	<b>Date</b>

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### **1. PURPOSE**

The presence of hemolysis, icteria and lipemia in serum or plasma may lead to erroneous laboratory test results. This procedure outlines how specimens, which have visible hemolysis, icteria or lipemia or which have been measured for hemolysis, icteria or lipemia by the Siemens Dimension System, should be handled in this laboratory.

### **2. SCOPE**

Applies to all test procedures.

### **3. RESPONSIBILITY**

All specimen processing and technical staff must be competent in identifying hemolyzed, icteric and lipemic samples and follow appropriate procedures.

### **4. DEFINITIONS**

Hemolysis – alteration or destruction of red blood cells in such a manner that hemoglobin is liberated into the medium in which the cells are suspended

#### Hemolysis Grading System (see Appendix A)

Slight:           ≈ 100 mg/dL hemoglobin  
Moderate:       ≈ 200 mg/dL hemoglobin  
Gross:           ≈ or > 500 mg/dL hemoglobin

Icteria – The yellow greenish color of the serum or plasma cause by bilirubin, a byproduct of old red cells.

Lipemia – Is manifested by a milky appearance of the serum or plasma caused by an excess of lipids in the blood.

## 5. PROCEDURE

Handling visible hemolysis:

1. Remove the specimen from the centrifuge. Make a visual assessment of the specimen's integrity, before testing.
2. If the plasma or serum portion of the specimen has an obvious pink to reddish appearance, compare it to the Hemolysis chart (Appendix A) to estimate the degree of hemolysis. Borderline readings should be reported to the next higher category.
3. If the hemolysis is slight (~100-200 mg/dL) the specimen should be given to the technical staff for testing in all the departments.
4. If moderate or gross hemolysis is observed, the specimen should be handled in the following manner  
Moderate - All coagulation specimens must be recollected  
                  - Chemistry specimens should be given to the department for testing.  
Gross         - All specimens should be recollected

**Note:** The Dimension chemistry system will calculate the HIL and Auto-verification rules in Data Innovation (Instrument Manager will flag appropriate action).

5. The staff member handling the sample will query the computer system to determine if other blood specimens have been drawn on the patient at the same time. Each specimen drawn will be located and examined for hemolysis. If moderate (for coagulation only) or gross hemolysis exists in any of the primary tubes, these must also be recollected.

*Note: Staff should exercise due diligence in determining if an alternate, acceptable specimen, might be available for testing before requesting recollection. For example: a green top tube might have been drawn in addition to a red top tube and could be an acceptable alternative for various chemistry assays.*

For nurse-collected specimens:

- a) Notify the nurse that the specimen is being canceled due to hemolysis.
- b) Nursing needs to reorder for a re-draw.

For Lab-collected specimens:

- a) Notify the nurse that a specimen is being canceled due to hemolysis.
- b) The Lab will order for a re-draw and a phlebotomist will be notified to re-collect the specimen.

### 6. Specific specimens

- a) For coagulation specimens, analyze the slightly hemolyzed specimen and report with the comment @HMS (Slight Hemolysis). Lipemia and icteria do not affect coagulation testing.

- b) For chemistry specimens, analyze the specimen and report according to section 8 below.
- 7. Grossly hemolyzed specimens (500 mg/dL or greater) should not be analyzed without the express permission of the Medical Director, or pathologist on call (example: post-mortem analysis) except when hemolysis does not effect a specific analyte. *For chemistry values refer to related document HIL Index Alert Values appendix B.*
- 8. The Siemens Dimension clinical chemistry analyzer is set to measures HIL with every analyte tested, which is based on the spectral characteristics of a serum or plasma sample. The HIL provides an index to alert the user to potential interference from hemolysis, icterus, and lipemia in the sample, where:

H = hemoglobin resulting from lysis of red blood cells  
I = icterus resulting from endogenous bilirubin  
L = lipemia or turbidity caused by insoluble lipids

The HIL Index appears on the report slip as a three-digit value where:

1st digit - H index  
2nd digit - I index  
3rd digit - L index

- a) When instrument reports are printed the technologists must check the instrument printouts for “HIL interference” codes. Under normal operation the Data Innovation (Instrument Manager) will flag HIL interference and guide the technical staff with appropriate result commenting or remedial action instruction.
- b) Append the canned comment HIR (Results may be inaccurate due to specimen hemolysis) to all analytes that have the “HIL interference” code for hemolysis.
- c) Append the canned comment IIR (Results may be inaccurate due to specimen icteria) to all analytes that have the “HIL interference” code for icteria.
- d) If the “HIL interference” code for lipemia is displayed, repeat the test(s) after ultra-centrifuging the specimen for all tests except AMON ( $\text{NH}_3$ ).

**Note:** At GEC, if the “HIL interference” code for lipemia is displayed, the specimen is referred to SGMC Laboratory for testing.

- e) For LDH and K results with the “HIL interference” code for hemolysis, remove the numeric results and result with the canned comment HLK (unable to analyze due to hemolysis).
- f) For  $\text{NH}_3$  results with the “HIL interference” code for lipemia, remove the numeric results and result with the canned comment LLK (unable to analyze due to lipemia), then send out the specimen for testing.

- g) The ordering doctor should be notified every time a NH<sub>3</sub> result is removed.
9. For the processing of lipemic and icteric hematology specimens see the procedure Coulter LH750 Operation for Complete Blood Count and Reticulocyte Automated Tests ~~Coulter HmX Operation for Complete Blood Count as applicable.~~

## 6. RELATED DOCUMENTS

Coulter LH750 Operation for Complete Blood Count and Reticulocyte Automated Tests

~~Coulter HmX Operation for Complete Blood Count~~

Airfuge Ultracentrifuge (Beckman Coulter<sup>TM</sup>)

Data Innovations Instrument Manager

HIL Index Alert Values (AG.F247)

## 7. REFERENCES

None

## 8. REVISION HISTORY

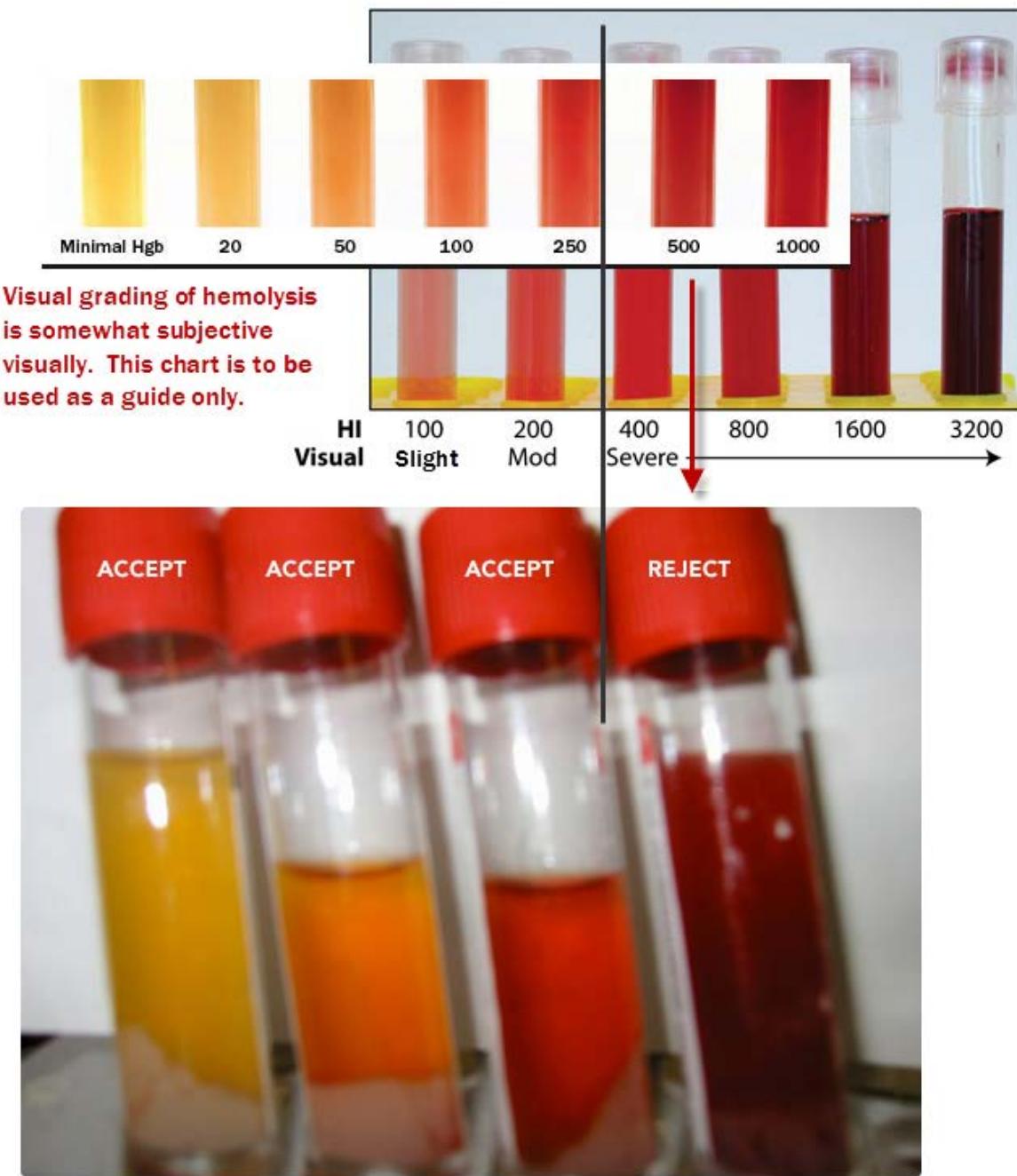
Version	Date	Reason for Revision	Revised By	Approved By
		Supersedes SOP L010.002		
000	4/16/2010	Updated owner	L. Barrett	L. Loffredo
001	4/1/2013	Updated owners Section 4: revise hemoglobin levels of slight and moderate hemolysis Section 6: updated documents Section 9: add App A, update App B	R. SanLuis	L. Loffredo
002	7/21/2015	Section 5: refer to related documents in step 7 Section 6: updated SOP list, moved HIL Index from section 9 and updated	L. Barrett	R. SanLuis

## 9. APPENDICES

A: Hemolysis, Icteria and Lipemia (HIL) Interference Chart

## APPENDIX A

### Hemolysis, Icteria, and Lipemia Interference



Note: It can be difficult to establish hemolysis when in combination with icteria and/or lipemia. In addition, special consideration/care should be taken when evaluating bullet tubes.

Form revised 3/31/00

TestCode	Hemolysis Hold	Icteria Hold	Lipemia Hold	Hemolysis Comment	Icteria Comment	Lipemia Comment	Add Comment	Replace Result	Internal Comment Instruction
ALC						8	LIP		
ALC					8		IIR		
ALC				8			HIR		
ALP					8		IIR		
ALP						8	LIP		
ALP				8			HIR		
ALT	7			7			HIR		
ALT	8			8			HIR		
ALT						8	LIP		
ALT						7	LIP		
ALT						5	LIP		
ALT						6	LIP		
AMON	7			7			HIR		
AMON	8			8			HIR		
AMON		7			7		IIR		
AMON		8			8		IIR		
AMON			7			7	*	LLK	LLK - Notify RN
AMON			8			8	*	LLK	LLK - Notify RN
AST				3			HIR		
AST					7		IIR		
AST				4			HIR		
AST				6			HIR		
AST				5			HIR		
AST					8		IIR		
AST						6	LIP		
AST						8	LIP		
AST						7	LIP		
AST					9		IIR		
AST						4	LIP		
AST						5	LIP		
AST					5		IIR		
AST					6		IIR		
AST					4		IIR		
AST				7			HIR		
AST				8			HIR		
CA					8		IIR		
CA						7	LIP		
CA						8	LIP		
CA						6	LIP		
CA						5	LIP		
CA				8			HIR		
CHOL				8			HIR		
CHOL					6		IIR		
CHOL					7		IIR		
CHOL					8		IIR		
CHOL					3		IIR		
CHOL					4		IIR		
CHOL					5		IIR		
CHOL						8	LIP		
CKI	5			5			HIR		
CKI	6			6			HIR		
CKI	7			7			HIR		
CKI	8			8			HIR		
CKI				4			HIR		
CRBM			8			8	LIP		Ultra-Centrifuge if Required
CRBM				8			HIR		
CRBM					8		IIR		
CREA					7		IIR		
CREA					5		IIR		
CREA					6		IIR		
CREA					8		IIR		
CREA						8	LIP		

TestCode	Hemolysis Hold	Icteria Hold	Lipemia Hold	Hemolysis Comment	Icteria Comment	Lipemia Comment	Add Comment	Replace Result	Internal Comment Instruction
CREA				7			HIR		
CREA				8			HIR		
DBIL	8			8			HIR		
DBIL						8	LIP		
DBIL					8		IIR		
DBILN	8			8			HIR		
DBILN			8			8	LIP		
DBILN				8			IIR		
ECO2				8			HIR		
ECO2					8		IIR		
ECO2						8	LIP		
FE	4			4			HIR		
FE	5			5			*	HLK	Recollect
FE	6			6			*	HLK	Recollect
FE	7			7			*	HLK	Recollect
FE	8			8			*	HLK	Recollect
FE			4			4	LIP		
FE			5			5	LIP		
FE			6			6	LIP		
FE			7			7	LIP		
FE			8			8	LIP		
FE				3			HMS		
FERR	7			7			*	HLK	Recollect
FERR	8			8			*	HLK	Recollect
FERR		6			6		IIR		
FERR		7			7		IIR		
FERR		8			7		IIR		
FERR			8			8	LIP		
FOLA	4			4			HIR		
FOLA	5			5			*	HLK	Recollect
FOLA	6			6			*	HLK	Recollect
FOLA	7			7			*	HLK	Recollect
FOLA	8			8			*	HLK	Recollect
FOLA		8			8		IIR		
FOLA			8			8	LIP		
FOLA				3			HMS		
GENP			8			8	LIP		Ultra-Centrifuge if Required
GENP					8		IIR		
GENP				8			HIR		
GENT			8			8	LIP		Ultra-Centrifuge if Required
GENT				8			HIR		
GENT					8		IIR		
GLUC	8			8			HIR		
GLUC						6	LIP		
GLUC						5	LIP		
GLUC						8	LIP		
GLUC						7	LIP		
GLUC					6		IIR		
GLUC						4	LIP		
GLUC					8		IIR		
GLUC					7		IIR		
IBCT	4			4			HIR		
IBCT	5			5			*	HLK	Recollect
IBCT	6			6			*	HLK	Recollect
IBCT	7			7			*	HLK	Recollect
IBCT	8			8			*	HLK	Recollect
IBCT						8	LIP		
IBCT						7	LIP		
IBCT						6	LIP		
IRON	4			4			HIR		
IRON	5			5			*	HLK	Recollect
IRON	6			6			*	HLK	Recollect

TestCode	Hemolysis Hold	Icteria Hold	Lipemia Hold	Hemolysis Comment	Icteria Comment	Lipemia Comment	Add Comment	Replace Result	Internal Comment Instruction
IRON	7			7			*	HLK	Recollect
IRON	8			8			*	HLK	Recollect
IRON						5	LIP		
IRON						6	LIP		
IRON						4	LIP		
IRON						7	LIP		
IRON						8	LIP		
IRON				3			HMS		
K	5						*	HLK	Hemolyzed
K	6						*	HLK	Hemolyzed
K	7						*	HLK	Hemolyzed
K	8						*	HLK	Hemolyzed
K				3			HMS		
K				4			HIR		
LA					7		IIR		
LA					8		IIR		
LA						8	LIP		
LA						7	LIP		
LA				8			HIR		
LDHPT	4						*	HLK	
LDHPT	5						*	HLK	
LDHPT	6						*	HLK	
LDHPT	7						*	HLK	
LDHPT	8						*	HLK	
LDHPT						6	LIP		Lipemic Specimen
LDHPT						7	LIP		Lipemic Specimen
LDHPT						8	LIP		Lipemic Specimen
LDHPT					8		IIR		
LDHPT				3			HMS		
LDI	4						*	HLK	
LDI	5						*	HLK	
LDI	6						*	HLK	
LDI	7						*	HLK	
LDI	8						*	HLK	
LDI						8	LIP		Lipemic specimen
LDI						6	LIP		Lipemic specimen
LDI				3			HMS		
LDI						7	LIP		Lipemic specimen
LDI					8		IIR		
LIPL	8			8			HIR		
LIPL					8		IIR		
LIPL						8	LIP		
MG	5			5			HIR		
MG	6			6			HIR		
MG	7			7			HIR		
MG	8			8			HIR		
MG						8	LIP		
MG							HIR		
MG				4					
MG						8	IIR		
PAL	8			8			HIR		
PAL		8				8	IIR		
PAL			8				8	LIP	
PFLDH	4						*	HLK	
PFLDH	5						*	HLK	
PFLDH	6						*	HLK	
PFLDH	7						*	HLK	
PFLDH	8						*	HLK	
PFLDH						6	LIP		Lipemic Specimen
PFLDH						7	LIP		Lipemic Specimen
PFLDH						8	LIP		Lipemic Specimen
PFLDH						8	8 LIP		Lipemic Specimen
PFLDH						8	IIR		

TestCode	Hemolysis Hold	Icteria Hold	Lipemia Hold	Hemolysis Comment	Icteria Comment	Lipemia Comment	Add Comment	Replace Result	Internal Comment Instruction
PFLDH				3			HMS		
PHOS	7			7			HIR		
PHOS	8			8			HIR		
PHOS							7 LIP		
PHOS							8 LIP		
PHOS					8		IIR		
PHOS						7	IIR		
PLTP	8			8			HIR		
PLTP						6	IIR		
PLTP						7	IIR		
PLTP						8	IIR		
PSA	8			8			HIR		
PSA		8				8	IIR		
PSA			8				8 LIP		
TBIL	8			8			HIR		Hemolyzed
TBIL						8	IIR		
TBIL							8 LIP		
TBILN	8			8			HIR		Hemolyzed
TBILN							8 LIP		
TBILN						8	IIR		
THEO			8				8 LIP		Ultra-Centrifuge if Required
TIBC	4			4			*	HLK	
TIBC	5			5			*	HLK	
TIBC	6			6			*	HLK	
TIBC	7			7			*	HLK	
TIBC	8			8			*	HLK	
TIBC			8				8 LIP		
TIBC							7 LIP		
TIBC							6 LIP		
TOBP							8 LIP		Ultra-Centrifuge if Required
TOBR							8 LIP		Ultra-Centrifuge if Required
TOBT							8 LIP		Ultra-Centrifuge if Required
TP	8			8			HIR		
TP						6	IIR		
TP						7	IIR		
TP						8	IIR		
TP							8 LIP		
TPPL	8			8			HIR		
TPPL						6	IIR		
TPPL						7	IIR		
TPPL						8	IIR		
URCA	8			8			HIR		
URCA							8 LIP		
URCA						8	IIR		
VANC			8				8 LIP		Ultra-Centrifuge if Required
VANC						8	IIR		
VANC						8	HIR		
VANP			8				8 LIP		Ultra-Centrifuge if Required
VANP						8	IIR		
VANP						8	HIR		
VANT			8				8 LIP		Ultra-Centrifuge if Required
VANT						8	IIR		
VANT						8	HIR		
VITB12	6			6			HIR		
VITB12	7			7			HIR		
VITB12	8			8			HIR		
VITB12		8				8	IIR		
VITB12			8				8 LIP		

Hold = Hold for Manual Test Review in Data Innovation (Instrument Manager).