## CKiD Chronic Kidney Disease in Children Cohort Study (CKiD) SECTION A: GENERAL INFORMATION

A1.		PARTICIPANT ID: AFFIX ID LABEL	OR ENTER NUMBER IF ID LABEL IS NOT AVAILABLE							
			-    -							
A2.		CKID VISIT #:	<u>0 1 a</u>							
A3.		FORM VERSION:	<u>1</u> <u>0</u> / <u>0</u> <u>1</u> / <u>1</u> <u>2 a</u>							
A4.		DATE OF VISIT:	$\frac{1}{M} \frac{1}{M} \frac{1}{D} \frac{1}{D} \frac{1}{Y} \frac{1}{Y} \frac{1}{Y} \frac{1}{Y}$							
A5.		FORM COMPLETED BY (INITIA	LS):							
The follo	owi	ng samples should be collecte	d.							
Sample	<u>s:</u>	Shipped to:	Shipped:							
Serum		CBL	IMMEDIATELY							
Serum		CBL	Batched							
			(Ship in Jan, Apr, Jul or Oct)							
Iohexol	Blo	ood CBL	IMMEDIATELY							
Urine		CBL	IMMEDIATELY							
	В		E SHIPPED QUARTERLY (Jan, Apr, July or Oct) SIRED BY THE SITE COORDINATOR!							
	Samples should NOT be stored for more than one year.									
		For specific questions,	contact your CCC prior to shipment.							
			T AND FIRST MORNING URINE COLLECTION							
B1.	•	participant a female of child-bearing								
		S	•							
	No		2 (Skip to B3)							
	PROMPT: QUESTION B2 IS FOR FEMALE PARTICIPANTS OF CHILD-BEARING POTENTIAL ONLY. URINE PREGNANCY TEST DATE MUST FALL WITHIN 72 HOURS BEFORE GFR TESTING DATE.									
B2.	a.	Urine pregnancy test date:	$\frac{1}{M} \frac{1}{M} \frac{1}{D} \frac{1}{D} \frac{1}{Y} \frac{1}{Y} \frac{1}{Y} \frac{1}{Y}$							
	b.	Urine pregnancy results:	4 (END. COMPLETE TRANSITIONAL (TDCC4) FORTY							
			1 (END; COMPLETE TRANSITIONAL (TRS01) FORM)							
		Negative	2							



#### FIRST MORNING URINE COLLECTION

Obtain urine collected at home in the specimen container that was shipped to the family before the visit. If URINE WAS NOT COLLECTED at home, collect FRESH urine into a specimen container provided by the central biochemistry laboratory.

Pour at least 1 mL of urine into the CBL transport tube.



Check that all information is correct on the urine collection tube and follow packaging instructions and ship to CBL.

Reasons Code List\*: 1= Not required 3 = Participant Refused 5 = Inadvertently Destroyed 7=Insufficient Volume

2 = Difficult Urine Collection 4 = Collection Contamination 6 = Oversight

	Sample Type (Required Volume):	(a) Sample Obta	ained:	(b) If No, specify reason *SEE CODE LIST ABOVE	(c) Additional Requirements:		
B3.	Urine Creatinine, Urine Protein, Urine Albumin (1 mL–10 mL)	1 (skip to c→)	2	 (skip to C1)	i. Is this a first morning urine sample? Yes2 ii. Time of Collection:: 1 = am, 2 = pm		

Encourage fluids throughout the visit.



Place two IV lines (18-22 gauge polyethylene catheters); using two separate vascular access sites --OR---

Place one butterfly and one IV line (18-22 gauge polyethylene catheter); using two separate vascular access sites; use tape to stabilize butterfly for Iohexol infusion

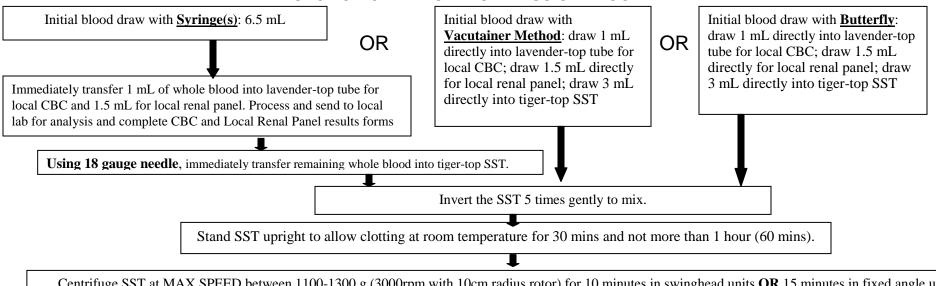
--OR---

Use butterfly for pre-iohexol and individual blood draw sticks; use tape to stabilize butterfly for Iohexol infusion

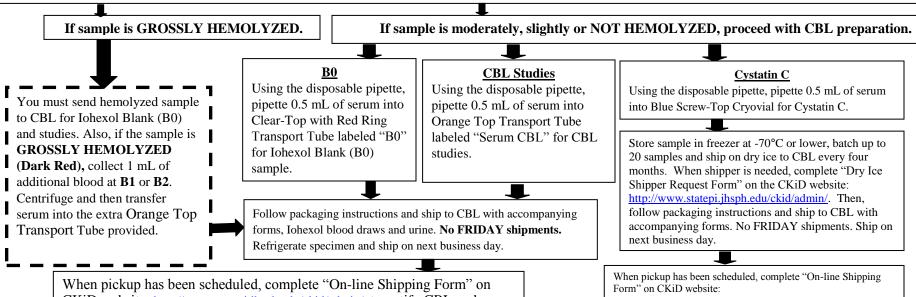


Complete Time=0 (Pre-Iohexol Infusion) blood draw according to MOP instructions/flowchart on page 3. NOTE: If patient has had a local CBC drawn within the past 30 days, those CBC results may be used instead of drawing another CBC and blood draw amounts can be decreased by 1 mL.

#### SECTION C: PRE-IOHEXOL INFUSION BLOOD DRAW



Centrifuge SST at MAX SPEED between 1100-1300 g (3000rpm with 10cm radius rotor) for 10 minutes in swinghead units OR 15 minutes in fixed angle units (balance tube in centrifuge). \*If incomplete separation, centrifuge again for 10-15 minutes.



CKiD website: http://www.statepi.jhsph.edu/ckid/admin/ to notify CBL and KIDMAC that sample(s) have been shipped to CBL.

http://www.statepi.jhsph.edu/ckid/admin/ to CBL and KIDMAC that sample(s) have been shipped.

C1. ACTUAL TIME OF PRE-IOHEXOL INFUSION BLOOD DRA		_	_	_	_			<b>.</b>		. ~				$\overline{}$			_			_	_						_
	4 V/V	$)$ R $^{\prime}$	1)	( )	( )(	. (	ΙR	)N	1(	1.5	-1	Νŀ	ı	()	- X	Н١	( )	⊢-I	РΚ	)⊢	⊢ ≀	I IIVA	IAI	: 11	Α(	1	

	:	1 = AM	2 = PM
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#### PROMPT: IF SUSPECTED BLOOD DRAW ADVERSE EVENT (i.e., infection), complete Adverse Event (ADVR) Form

Reasons Code List\*:1= Not required3 = Participant Refused5 = Inadvertently Destroyed2 = Difficult Blood Draw4 = Red Blood Cell Contamination6 = Oversight

(R	Sample Type equired Volume in Top Color Tube Type):	(a) Sample Obta	ined:	(b) If No, specify reason *SEE CODE LIST ABOVE	(c) Additional Requirements:				
C2.	Renal/ Uric Acid Chemistries (2.0 mL in Tiger Top SST)	1 (skip to c→)	2	(skip to C3)	Indicate the appearance of the serum after centrifuging.  Grossly (Dark Red)				
C3.	Cystatin C (1.0 mL in Tiger Top SST)	1 (skip to c→)	2	 (skip to C4)	Date Frozen: /				
C4.	Local CBC (1.0 mL in Lavender Top tube)	1 (skip to C5)	2	(skip to C5)	N/A				
C5.	Local Renal Panel (1.5 mL in Local SST)	1 (skip to D2)	2	(skip to D2)	N/A				

#### SECTION D: OPTIONAL LOCAL LAB TEST (IF CLINICALLY INDICATED)

Check with the PI at your clinical site to determine whether or not it is **CLINICALLY INDICATED** to obtain urine for local lab. These are instances when the PI needs results immediately and/or the participant needs additional local labs performed (i.e., local Urine Creatinine and Urine Protein).

	QUESTION D1 HAS BEEN DELETED.
D2.	Was a urine protein to creatinine ratio assay performed at the clinical site's local laboratory?  Yes
	SECTION E: INFUSION SYRINGE WEIGHT
E1.	SCALE MUST BE FIRST ZEROED BEFORE WEIGHING. REMOVE ALUMINUM FOIL PRIOR TO WEIGHING THE SYRINGE. THE <u>SAME</u> SCAL MUST BE USED TO WEIGH THE SYRINGE <u>PRE AND POST</u> IOXEHOL INFUSION.
	a. Syringe Weight <b>Pre- lohexol Infusion</b> :(g)
	b. Syringe Weight <b>Post- Iohexol Infusion</b> : (g) (Post-Infusion Weight should be <b>at least 6.0g</b> less than Pre Infusion Weight. If Post-Infusion Weight is not at least 6g less, please confirm.)
	PRE AND POST SYRINGE WEIGHT MUST BE OBTAINED IN ORDER TO CALCULATE CHILD'S GFR.
	SECTION F: IOHEXOL - Refer to Instructions for Iohexol Infusion and GFR Blood Draws Flow Chart on Page 8
>	BEFORE INFUSING 5 mL of IOHEXOL, SET TIMER = 0. SIMULTANEOUSLY START TIMER AND BEGIN IOHEXOL INFUSION COMPLETE INFUSION BETWEEN 1 TO 2 MINS.  LEAVE TIMER RUNNING THROUGHOUT IOHEXOL INFUSION AND SUBSEQUENT BLOOD DRAWS
F1.	IOHEXOL INFUSION a. INFUSION START TIME: : 1 = AM 2 = PM

- > DO NOT DRAW BLOOD FROM THE IV SITE WHERE IOHEXOL WAS INFUSED. ANOTHER IV SITE MUST BE USED.
- WASTE 1 mL OF BLOOD IF DRAWING FROM A SALINE/HEPARIN LOCK.
- COLLECT 1 mL OF BLOOD FOR EACH IOHEXOL BLOOD DRAW IN THE PROVIDED SST.
- > RECORDING THE EXACT NUMBER OF MINUTES ON THE TIMER IS MORE IMPORTANT THAN DRAWING THE BLOOD EXACTLY AT 120 & 300 MINUTES AFTER IOHEXOL INFUSION. FOR EXAMPLE, IF BLOOD IS DRAWN AT 133 MINS INSTEAD OF 120 MINS, DOCUMENT BLOOD DRAWN @ 133 MINS.
- > TIME SHOULD BE RECORDED IMMEDIATELY <u>AFTER</u> EACH BLOOD SAMPLE IS OBTAINED (i.e., B1, B2).

### POST VITALS SHOULD BE TAKEN 10 MINUTES AFTER INFUSION USING LOCAL BLOOD PRESSURE MEASUREMENT (i.e. DINAMAP)

- If rash develops after lohexol Infusion, consider it a reaction to lohexol and notify PI immediately. Consider administration of 1 mg/kg Benadryl IV (maximum dose: 50 mg Benadryl IV).
- In the rare event that systolic BP decreases more than 25 mm Hg, diastolic BP decreases more than 20 mmHg, or pulse increases more than 20 beats per min, notify PI immediately to evaluate reaction and complete the Adverse Event (ADVR) Form. Consider the possibility of an anaphylactic reaction to lohexol. Consider administration of 1 mg/kg Benadryl IV (maximum dose: 50 mg Benadryl IV). Draw up to 0.1 mL 1:1000 Epinephrine for SQ injection and 2 mg/kg Solumedrol IV for administration as ordered by physician.

	(i) Post Vitals:	
F2a.	Post- infusion blood pressure:	/
b.	Post-infusion temperature:	1 = °C Typical range: <b>36.1 – 38.3</b> 2 = °F Typical range: <b>94.5 – 100.6</b>
C.	Post-infusion number of heart beats per minute:	
d.	Post-infusion respirations per minute:	

# INVERT TUBE 5-10 TIMES AFTER EACH BLOOD DRAW LET SST TUBE STAND 30 MINUTES (BUT NO LONGER THAN 1 HOUR) CENTRIFUGE AT MAX SPEED BETWEEN 1100-1300g (3000rpm with 10cm radius rotor) for 10 MINUTES IN SWING HEAD OR 15 MINUTES IN FIXED ANGLE (BALANCE TUBES IN CENTRIFUGE)

	ALL TIMES should be documented from the initial infusion time	HOURS/	i) UAL MINUTES IMER	(ii) ONLY if Timer malfunctions, record Clock Time using the same clock used for G1a	(iii) Difficult Blood Draw: Yes No		(iv) Blood Drawn via Venipuncture Yes No		(v) Blood Volume Collected (1 mL):	(v Centri at Clinio Yes	fuged
F3a.	<b>B1 2 hrs</b> (120 min):	hr	mins	: 1 = AM 2 = PM	1 (Skip to b)	2	1	2	mL	1 (Skip to F4a)	2 (Skip to F4a)
b.	<b>B1</b> 2 <sup>nd</sup> attempt:	hr	mins	: 1 = AM 2 = PM	1	2	1	2	mL	1	2
F4a.	<b>B2 5 hrs</b> (300 min):	hr	mins	: 1 = AM 2 = PM	1 (Skip to b)	2	1	2	mL	1 (END FORM)	2 (END FORM)
b.	<b>B2</b> 2 <sup>nd</sup> attempt:	hr	mins	: 1 = AM 2 = PM	1	2	1	2	mL	1	2

#### **Instructions for Iohexol Infusion and GFR Blood Draws**

Place two IV lines (18-22 gauge polyethylene catheters) using two separate vascular access sites ---OR---

Place one butterfly and one IV line (18-22 gauge polyethylene catheter) using two separate vascular access sites; use tape to stabilize butterfly for iohexol infusion

--OR---

Use butterfly for pre-iohexol and individual blood draw sticks; use tape to stabilize butterfly for Iohexol infusion

Complete pre-iohexol infusion blood draw according to instructions

Start timer and infuse iohexol over 1-2 minutes

If infusion site for iohexol IV or butterfly infiltrates, study MUST BE DISCONTINUED Must wait at least 48 hours to repeat study, and repeat visit should occur within 3 months of initial

Flush with 10 mL normal saline to ensure infusion of all of the iohexol; IOHEXOL IV OR BUTTERFLY MAY NOW BE REMOVED

10 minutes post-infusion CHECK POST-INFUSION VITALS DO NOT DRAW BLOOD AT 10 MINUTES

Obtain post-infusion weight of syringe on the same scale as prior to infusion. Record weight on the Specimen Collection Form prior to shipment to the CCC

Draw 1 ml of blood into SST at 120 mins (B1), and 300 mins (B2).

\*For IV ONLY: Discard initial 1 mL blood waste at each blood draw

\*For IV ONLY: Flush after each blood draw with at least 3 mL normal saline

\*Record the time each blood draw is completed

If rash develops, consider it a reaction to iohexol and **notify PI immediately**. Consider administering 1mg/kg Benadryl IV (maximum dose 50mg).

If systolic BP decreases > 25 mm Hg, diastolic BP decreases > 20 mm Hg or pulse increases > 20 beats per min, consider this an anaphylactic reaction to iohexol and **notify PI immediately**; consider administering 1 mg/kg Benadryl IV (maximum dose 50 mg) and draw up to 0.1 mL 1:1000 Epinephrine for SQ injection and 2 mg/kg Solumedrol IV if necessary.

COMPLETE ADVERSE EVENT FORM and send to CCC for data entry

If blood draw is difficult with poor volume delivery, repeat draw 5 minutes later; record time on Specimen Collection Form and tube. If blood draw is impossible, repeat with new venipuncture.

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Following each blood draw: gently invert tube 5-10 times

Stand SST upright at room temp for 30 minutes, but no longer than 1 hour

Centrifuge at MAX SPEED between 1100-1300 g for 10 minutes in swinghead or 15 minutes in fixed angle\*

Transfer serum into the cryovial labeled "Iohexol Serum."

Following packaging and shipping instructions; send urine, blood and copies of completed shipping form(s) & confirmation of written consent to CBL

Physician should be immediately available (in person or by phone) during Iohexol Infusion Encourage fluids throughout the visit.

\*1100-1300 g = 3000 rpm with 10 cm radius rotor

