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

A1.	PARTICIPANT ID: AFFIX ID LABEL OR ENTER NUMBER IF ID LABEL IS NOT AVAILABLE					ABLE	
			l <sub>-</sub>	_  -   _	-   _	_	
A2.	CKiD Post KRT	VISIT #:	_				
A3.	FORM VERSIO	DN:	_	0 6 /	<u>1</u> <u>5</u>	1 2 2	
A4.	SPECIMEN CO	DLLECTION	_	M M D	$\frac{1}{D} \frac{1}{Y}$	<u> </u>	
A5.	FORM COMPL	ETED BY (I	NITIALS): _		-		
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BATCHED SAMPLES SHOULD BE SHIPPED QUARTERLY (Jan, Apr, July or Oct)
OR MORE OFTEN IF DESIRED BY THE SITE COORDINATOR!

Samples should NOT be stored for more than six (6) months. For specific questions, contact your CCC prior to shipment.



### **SECTION B: PREGNANCY TEST**

B1.	ls p	Is participant a female of child-bearing potential?				
		S	1 (See PROMPT Below) 2 (Skip to C1)			
BLO	OD/L	•	PARTICIPANTS OF CHILD-BEARING POTENTIAL ONLY. MUST OCCUR ON DAY OF VISIT OR FALL WITHIN 72			
B2.	a.	Blood/Urine pregnancy test date:	$\overline{M} \overline{M} \overline{D} \overline{D} \overline{V} \overline{Y} \overline{Y} \overline{Y} \overline{Y}$			
	b.	Pregnancy results:  Positive  Negative	1 (END; COMPLETE TRANSITIONAL (TRS03) FORM) 2			

#### SECTION C: POST-DIALYSIS VISIT BLOOD DRAW

For Initial Blood Draw with <u>Syringe</u>, <u>Vacutainer</u> OR <u>Butterfly</u> Method: Select the type of consent obtained (options 1 through 2) that pertains to the CKiD participant:

If participant consented to BIOLOGICAL samples:

Collect 14-15 mL if participant is < 30 kg OR 20-21 mL if participant is  $\ge 30 \text{ kg}$ .

If < 30 kg, immediately transfer (using 18 gauge needle) or draw:

- 8.5 mL into (1) Tiger-Top SST for CBL & NIDDK BR
- 3 mL into (1) PST for NIDDK Biorepository
- 1 mL in lavender-top tube for local CBC (tube not provided in CBL kit)
- 1.5 mL in appropriate tube (not provided) for local Renal Panel
- 1 mL of additional blood in SST for CBL (if initial sample is GROSSLY HEMOLYZED)

If  $\geq 30$  kg, immediately transfer (using 18 gauge needle) or draw:

- 12.5 mL into (2) Tiger-Top SSTs for CBL & NIDDK BR
- 5 mL into (2) PST for NIDDK Biorepository
- 1 mL in lavender-top tube for local CBC (tube not provided in CBL kit)
- 1.5 mL in appropriate tube (not provided) for local Renal Panel
- 1 mL of additional blood in SST for CBL (if initial sample is GROSSLY HEMOLYZED)

#### 2 If participant did NOT consent to BIOLOGICAL samples:

Collect 5-6 mL from all participants (regardless of weight) as specified below.

Immediately transfer (using 18 gauge needle) or draw:

- 2.5 mL into (1) Tiger-Top SSTs for CBL
- 1 mL in lavender-top tube for local CBC (tube not provided in CBL kit)
- 1.5 mL in appropriate tube (not provided) for local Renal Panel
- 1 mL of additional blood in SST for CBL (if initial sample is GROSSLY HEMOLYZED)

#### SECTION C: POST-DIALYSIS VISIT BLOOD DRAW PROCESSING

Invert the SST 5 times & PST 8-10 times gently to mix. Stand SST upright to allow clotting at room temperature for 30 mins and not more than 1 hour (60 mins). Centrifuge SST & PST at MAX SPEED between 1100-1300g (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. If sample is moderately, slightly or NOT HEMOLYZED, proceed with CBL and NIDDK BR preparation. If sample is GROSSLY HEMOLYZED. You must send hemolyzed **CBL Studies** NIDDK BR (Serum) iPTH/hsCRP NIDDK BR (Plasma) sample to CBL. If participant Pipette 3mL (<30kg) or 5mL Pipette 1.5mL (<30kg) or Using the disposable Pipette 0.5 mL of serum has eaten, aliquot hemolyzed (≥30kg) serum into clear top pipette, pipette 0.75 mL 2.5mL (≥30kg) plasma into into red top cryovial tube sample for fasting lipid profile cryovial for NIDDK BR of serum into Orange Top cryovial with green cap insert for CBL iPTH &, hsCRP (use different pipettes for and send to CBL. Also, if the (use different pipettes for Transport Tube labeled serum and plasma). sample is **GROSSLY** "Serum CBL" for CBL serum and plasma). \*If there is any extra serum, then \*If there is any extra plasma, then renal/uric acid and lipids. HEMOLYZED (Dark Red). pipette the extra serum into the pipette the extra plasma into the Follow packaging Store sample in freezer at -70°C or lower, collect 1 mL of additional clear top cryovial marked green cap insert cryovial marked instructions and ship to blood in a SST. Centrifuge and batch up to 20 samples and ship on dry ice "NIDDK BR SERUM" "PLASMA (Extra)". CBL with accompanying then transfer serum into the quarterly (January, April, July and forms. No FRIDAY October) to the CBL. When shipper is extra Orange Top Transport shipments. Refrigerate needed, complete "CBL Dry Ice Shipper Tube provided. specimen and ship on Complete the SM01 form, store sample in freezer at -70°C or lower, batch next business day. Request Form" on the CKiD website: up to 40 samples and ship on dry ice quarterly (Jan, April, July and Oct) https://statepi.jhsph.edu/ckid/coordinatorto the NIDDK BR. No Thursday/Friday shipments. When shipper is resources/ Then, follow packaging needed, complete "NIDDK BR Dry Ice Shipper Request Form" on CKiD instructions and ship to CBL with website: https://statepi.jhsph.edu/ckid/coordinator-resources/ accompanying forms. No FRIDAY Then, follow packaging instructions. shipments. Refrigerate and ship on next business day. When pickup has been scheduled, complete "On-line Shipping Form" on CKiD website: https://statepi.jhsph.edu/ckid/coordinator-resources/

When pickup has been scheduled, complete "On-line Shipping Form" on CKiD website: <a href="https://statepi.jhsph.edu/ckid/coordinator-resources/">https://statepi.jhsph.edu/ckid/coordinator-resources/</a> to notify the appropriate personnel from the CBL and the NIDDK BR.

C1. ACTUAL TIME OF BLOOD DRAW : 1 = AM 2 =
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PROMPT: IF SUSPECTED BLOOD DRAW ADVERSE EVENT (i.e., infection), complete ADVERSE EVENT (ADVR) Form

Sample Type (Required Volume in Top Color Tube Type):		(a) Sample Obtained:		(b) If No, specify reason *SEE CODE LIST ABOVE	(c) Additional Requirements:	
C2.	Renal/Uric Acid Chemistries (1.0 mL in Tiger Top SST)	Yes  1 (skip to c→)	2	(skip to C3)	Indicate the appearance of the serum after centrifuging.  Grossly (Dark Red)	
C3.	Local CBC (1.0 mL in Lavender Top tube)	1 (skip to C4)	2	(skip to C4)	N/A	
C4.	Local Renal Panel (1.5 mL in Local SST)	1 (skip to C5)	2	(skip to C5)	N/A	
C5.	Serum for Fasting Lipid Panel (0.5 mL in Tiger Top SST)	1 (skip to c→)	2	(skip to C6)	Did the participant fast after midnight? Yes1 No2*	
C6.	Serum for ipth & hsCRP (1.0 mL in Tiger Top SST)	1 (skip to c→)	2	(skip to D1)	Date Frozen://	

<sup>\*</sup>If the participant did not fast, the Nephron Lipid Report will indicate that the participant did not fast.

Sites can obtain results for lab values that have been identified as "KEY VARIABLES". To obtain results, go the CKiD Nephron Website: <a href="https://statepiaps8.jhsph.edu/nephron/groups/aspproc/">https://statepiaps8.jhsph.edu/nephron/groups/aspproc/</a>, click on "Report Menu" and choose the appropriate lab report (i.e., Selected Renal Panel Lab Variables Report.)

#### SPECIMEN COLLECTION FORM for POST DIALYSIS VISIT (XX)

#### **SECTION D: NIDDK BIOREPOSITORY**

D1. Did the participant consent to have biological samples (i.e., serum and plasma samples) stored at the NIDDK Biorepository?

Reasons Code List\*: 1= Not required 4 = Red Blood Cell Contamination 7 = Exceed maximum allowable volume 5 = Inadvertently Destroyed 3 = Participant Refused 6 = Oversight

	Sample Type (Required Volume in Top Color Tube Type):		btained:	(b) If No, specify reason *SEE CODE LIST ABOVE	(c) Additional Requirements:
		<u>Yes</u>	<u>No</u>	SEE CODE EIST ABOVE	
D2.	Serum for NIDDK Biorepository (**6.0 mL or **10.0 mL of blood in Tiger Top SST)	1 (skip to c→)	2	(skip to D3)	Date Frozen://
D3.	Plasma for NIDDK Biorepository (***3.0 mL of blood in one Green Top or ***5.0 mL in two Green Top PSTs)	1 (skip to c→)	2	(skip to E1)	Date Frozen:  M M D D Y Y Y Y

<sup>\*\*</sup> Collect 6.0 mL of whole blood for participants < 30 kg and 10.0 mL for participants ≥ 30 kg

<sup>\*\*\*</sup> Collect 3.0 mL of whole blood for participants < 30 kg and 5.0 mL for participants ≥ 30 kg

#### **SECTION E: LOCAL Kt/V**

E1.	Are local Kt/V results available? Yes No, Specify reason below	1 <b>(Skip to E2)</b>			
	E1a. Reason:	(END Form)			
E2.	Date of most recent Kt/V results:  N	/ M D D Y Y Y Y			
E3.	Which modality is the participant receiving?  Hemodialysis  Peritoneal dialysis				
E4.	Hemodialysis ONLY:				
	a. URR (urea reduction ratio)	%			
	b. Total Kt/V	-    (END Form)			
E5.	Peritoneal dialysis (PD) ONLY:				
	a. PD Kt/V	•   _			
	b. 24-hour Urine Volume				
	c. Urine Kt/V	•			
	d. Total Kt/V (PD Kt/V plus urine Kt/V)	-			