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

AI.	PARTICIPANT ID: AFFIX ID LABEL OR EI	CIPANT ID: AFFIX ID LABEL OR ENTER NUMBER IF ID LABEL IS NOT AVAILABLE		
		-   _  -   _		
A2.	CKiD VISIT #:			
A3.	FORM VERSION:	0 3 / 0 1 / 1 8		
A4.	SPECIMEN COLLECTION DATE:	$\frac{1}{M} \frac{1}{M} \frac{1}{D} \frac{1}{D} \frac{1}{Y} \frac{1}{Y} \frac{1}{Y} \frac{1}{Y}$		
A5.	FORM COMPLETED BY (INITIALS):			

The following samples should be collected.

Samples: Shipped to Shipped:

Serum CBL IMMEDIATELY

Serum CBL Batched

(Ship in Jan, Apr, Jul or Oct)

Urine CBL IMMEDIATELY Iohexol Blood\* CBL IMMEDIATELY

If consent is obtained for biological samples, collect the following:

Samples:Shipped to:Shipped:Serum (Biological)NIDDK BiorepositoryBatched

(Ship in Jan, Apr, Jul or Oct)

Plasma (Biological) NIDDK Biorepository Batched

(Ship in Jan, Apr, Jul or Oct)

Urine (Biological) NIDDK Biorepository Batched

(Ship in Jan, Apr, Jul or Oct)

Toenail Clippings NIDDK Biorepository IMMEDIATELY

(Biological)

### 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 one year. For specific questions, contact your CCC prior to shipment.



<sup>\*</sup> COLLECT IOHEXOL BLOOD DRAW: Only if Cohort 3 participants who consent to iohexol protocol or Cohorts 1 & 2 participants had previous iGFR>90

#### SECTION B: PREGNANCY TEST AND FIRST MORNING URINE COLLECTION

B1.	ls p	participant a female of child-bearing potential?				
		5				
		_(=====================================				
	PROMPT: QUESTION B2 IS FOR FEMALE PARTICIPANTS OF CHILD-BEARING POTENTIAL ONLY. URINE PREGNANCY TEST DATE MUST FALL WITHIN 72 HOURS BEFORE STUDY VISIT DATE.					
If per INITI		ning iohexol protocol, B2 MUST BE COMPLETED BEFORE IOHEXOL TESTING IS D.				
B2.	a.	Urine pregnancy test date:  M M D D Y Y Y Y				
	b.	Urine pregnancy results:				
		Positive				
		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 sample during CKiD visit.

(Refer to MOP Section 11 and/or CBL flowchart for additional information and directions)

Pour at least1mL 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 Obtained:		(b) If No, specify reason *SEE CODE LIST ABOVE	(c) Additional Requirements:	
		<u>Yes</u>	<u>No</u>	SEE CODE LIST ABOVE		
B3.	Urine Creatinine, Urine Protein, Urine Albumin (CBL)	1 (skip to c→)	2		i. Is this a first morning urine sample? Yes1  No2	
	(1 mL-10 mL)	(3Kip 10 0-7)		(skip to C1)	ii. Time of Collection: : 1 = am, 2 = pm	

#### **SECTION C: Visit 4 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 3) That Pertains to the CKiD Participant:

#### **1** 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 (2) Tiger-Top SSTs for CBL & NIDDK Biorepository
- 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 Biorepository
- 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)

#### For Participant Completing Iohexol Study Visit

Only Cohort 3 participants who consent to iohexol protocol or Cohorts 1 & 2 participants with previous iGFR>90

For IOHEXOL study visits:

• 1mL of additional blood into Tiger Top SST for CBL Iohexol Blank (B0) blood sample

Iohexol is infused at the time of initial blood draw.

Refer to page 10 for **Instructions for Iohexol Infusion and GFR Blood Draws.** 

#### PROCESSING OF BLOOD FOR CBL & NIDDK BR

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 1100-1300 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.

If sample is GROSSLY HEMOLYZED.

If sample is moderately, slightly or NOT HEMOLYZED, proceed with CBL and NIDDK BR preparation.

You must send hemolyzed sample to CBL. If participant has eaten, l aliquot hemolyzed sample for fasting lipid profile and send to CBL. Also, if the sample is GROSSLY **HEMOLYZED** (Dark

# Red), collect 1 mL of

additional blood in a SST. Centrifuge and then transfer serum into the extra Orange Top Transport Tube provided.

#### For IOHEXOL STUDY VISITS ONLY:

#### **B0** sample

Using the disposable pipette, pipette 0.5 mL of serum into Round-bottom Orange Top Transport Tube labeled "B0" for Iohexol Blank (B0) sample. Follow packaging instructions and ship to CBL with accompanying forms, iohexol I blood draws and urine. No FRIDAY shipments. Refrigerate specimen and ship

on next business day.

#### **CBL Studies**

Using the disposable pipette, pipette 0.75 mL of serum into Orange Top Transport Tube labeled "Serum CBL" for CBL renal/uric acid and lipid. Follow packaging instructions and ship to CBL with accompanying forms, iohexol blood draws, and urine. No FRIDAY shipments.

Refrigerate specimen and ship on next business day.

#### **Cystatin C**

Using the disposable pipette, pipette 0.5 mL of serum into Blue Screw-Top Cryovial for Cystatin C. Store sample in freezer at -70°C or lower, batch up to 20 samples and

ship on dry ice every four months to the CBL. When shipper is needed, complete "CBL Dry Ice Shipper Request Form" on the CKiD website:

http://www.statepi.jhsph.edu/ckid/admin/ Then, follow packaging instructions and ship to CBL with accompanying forms. No FRIDAY shipments. Ship on next business day.

#### **NIDDK BR** (Serum & Plasma)

Pipette 3 mL (<30kg) or 5 mL (>30kg) of serum into clear top cryovial & 1.5 (<30kg) or 2.5 mL(> 30kg) of plasma into cryovial into green cap insert (use different pipettes for serum and plasma).

\*If there is any extra serum and/or plasma, then pipette the extra serum into the clear top cryovial marked "NIDDK BR SERUM" and the extra plasma in green cap insert cryovial marked "PLASMA (Extra)." Store samples in freezer at -70°C or lower, & batch up to 40 samples and ship on dry ice in January, April, July or October to the NIDDK BR. When shipper is needed, complete "NIDDK BR Shipper Request Form" on the CKiD website:

http://www.statepi.jhsph.edu/ckid/admin/. Then, follow packaging instructions.

When pickup has been scheduled, complete "On-line Shipping Form" on CKiD website: http://www.statepi.jhsph.edu/ckid/admin/ to notify the appropriate personnel from the CBL and the NIDDK BR.

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

Reasons Code List\*: 1= Not required 4 = Red Blood Cell Contamination 7 = Exceed maximum allowable volume

2 = Difficult Blood Draw 5 = Inadvertently Destroyed

3 = Participant Refused 6 = Oversight

(Requ	Sample Type ired Volume in Top Color Tube Type):	(a) Sample Ob	tained:	(b) If No, specify reason *SEE CODE LIST ABOVE	(c) Additional Requirements:
		<u>Yes</u>	<u>No</u>	SEE CODE EIGT ABOVE	
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 C4a)	Date Frozen: /
C4a.	Local CBC (1.0 mL in Lavender Top tube)	1 (skip to C4b)	2	(skip to C4b)	N/A
C4b.	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*

<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.)

Yes	ples (i.e., serum, plasma and urine samples) stored at the NIDDK Biorepository?
No	

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

	Sample Type (Required Volume in Top Color Tube Type):	(a) Sample Obtained:		(b) If No, specify reason *SEE CODE LIST ABOVE	(c) Additional Requirements:
		<u>Yes</u>	<u>No</u>	"SEE CODE LIST ABOVE	
C7.	Serum for NIDDK Biorepository	1	2		Date Frozen:
	(**6.0 mL or **10.0 mL of blood in Tiger Top SST)	(skip to c→)		(skip to C8)	////
C8.	Plasma for NIDDK Biorepository	1	2		Date Frozen:
	(***3.0 mL of blood in one Green Top or	(skip to c→)		(skip to D1)	/
	***5.0 mL in two Green Top PSTs)				M M D D Y Y Y

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

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

#### SECTION D: URINE COLLECTION AND PROCESSING FOR REPOSITORY

Collect FRESH urine into an initial urine collection cup or hat (provided by the site).

Pour 15-60 mL (preferably 60 mL) of FRESH urine into 90 mL urine collection cup with 4 protease inhibitor tablets. Do not fill the urine past the 60 mL mark on the collection cup. One protease inhibitor tablet should be used for 10-15 mL of urine (see **Table A**). For example if 30 mL of urine is collected, ONLY 2 PI tablets are needed. (Like all unused supplies, unused protease inhibitor tablets should be returned to the CBL.)

Invert the urine cup gently 5 – 10 times.

TABLE A:

# of Protease
Inhibitor Tablets

10 - 15 mL
16 - 30 mL
31 - 45 mL
46 - 60 mL

4

The PROTEASE INHIBITOR TABLET(s) MUST BE COMPLETELY DISSOLVED in the urine.

Once the protease inhibitor tablet(s) are completely dissolved, pour urine into up to six (6) 10 mL urine centrifuge tubes. (For each tube: remove yellow top cap, pour urine into tube and SCREW cap back onto tube.) Place no more than 10 mL in each tube.

-- OR -

Sites may also substitute with tubes normally used to centrifuge urine at site.

Centrifuge urine tube(s) at MAX SPEED between 1100-1300g (3000rpm with 10cm radius rotor) for 10 mins (swinghead units) – **OR** – 15 mins (fixed angle units).

Decant (pour off) the supernates (liquid reaction) into up to seven (7) 10 mL urine cryovials. Pour no more than 9 mL of urine into each 10 mL cryovial to allow for expansion.

Check that all information is correct on the urine cryovials, promptly freeze and store sample(s) at -70°C or lower. Batch samples and ship at least quarterly (include maximum of 36 cryovials per shipper). When shipper(s) is needed, complete "NIDDK Shipper Request Form" on CKiD website: <a href="http://www.statepi.jhsph.edu/ckid/admin/">http://www.statepi.jhsph.edu/ckid/admin/</a>. Then, follow packaging instructions.

When pickup has been scheduled, complete "Online Shipping Form" on CKiD website to notify the NIDDK BR and KIDMAC that sample(s) have been shipped to NIDDK BR.

Reasons Code List\*: 1= Not required 2 = Difficult Urine 3 = Participant 4 = Collection 5 = Inadvertently 6 = Oversight 7 = Insufficient volume

Sample Type (Required Volume in Top Color Tube Type):	(a) Sample Obt	ained:	(b) If No, specify reason *SEE CODE LIST ABOVE	(c) Additional Requirements:
D1. Urine for NIDDK Biorepository (15.0 - 60.0 mL of urine in specimen container and transferred into collection cup with protease inhibitors)	1 (skip to c→)	2		i. Was supernate decanted into urine transport cryovials? Yes1 No2 ii. Date Frozen:
				M M D D Y Y Y

#### **SECTION E: OPTIONAL TESTS** 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).					
E2.	Was	a urine protein to creatinine ratio assay perfo	ormed at the clinical site	e's local laboratory?			
			Complete Local Urine CLINICALLY INDICAT	e Assay Results Form L06, ΟΝ ΓΕD	ILY if local labs are		
		ı	OHEXOL PROTOCO	DL			
E3.	Is the	e participant completing iohexol study visit?		ned 1 2 <b>→ (Sk</b>	ip to Section H)		
<u>O</u>	NLY	COMPLETE SECTIONS F & G IF	PARTICIPANT IS	S COMPLETING IOHEX	OL STUDY VISIT.		
On	ily Co	ohort 3 participants who consent to ioh should complete iohexol proto For an iohexol study visit, additional should be o	ocol. If you have add	ditional questions, contact of contact of the lonexol "B0" Black	CCC.		
	SECTION F: INFUSION SYRINGE WEIGHT						
F1.		ALE MUST FIRST BE ZEROED BEFORE WI E <u>SAME</u> SCALE MUST BE USED TO WEIGH					
	a.	Syringe Weight <b>Pre-Iohexol Infusion</b> :	(g)				
	b.	Syringe Weight <b>Post-Iohexol Infusion</b> :	(g)	(Post-Infusion Weight should I than Pre-Infusion Weight. If Po at least 6g less, please confirm	ost-Infusion Weight is no		
PRE	ANI	D POST SYRINGE WEIGHT MUST B	E OBTAINED IN O	RDER TO CALCULATE P	ARTICIPANT'S GFR		

#### <u>(.</u>

SECTION G: IOHEXOL – Refer to Instructions for Iohexol Infusion and GFR Blood Draws Flow Chart on Page 10

- 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

#### **Instructions for Iohexol Infusion and GFR Blood Draws for Make-up GFRs**

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 (or per device guidelines)

\*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.

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



G1.	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 (OR PER DEVICE GUIDELINES).
- > 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:						
G2a.	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	ACTUAL HOURS/ MINUTES on TIMER		(ii) ONLY if Timer malfunctions, record Clock Time using the same clock used for G1a	(iii) Difficult Blood Dra Yes	_	Blood D	iv) Orawn via uncture No	(v) Blood Volume Collected (1 mL):	(\ Centri at Clinio Yes	ifuged
G3a.	<b>B1 2 hrs</b> (120 min):	hr	mins	: 1 = AM 2 = PM	1 (Skip to b)	2	1	2	mL	1 (Skip to G4a)	2 (Skip to G4a)
b.	<b>B1</b> 2 <sup>nd</sup> attempt:	hr	mins	: 1 = AM 2 = PM	1	2	1	2	mL	1	2
G4a.	<b>B2 5 hrs</b> (300 min):	hr	mins	: 1 = AM 2 = PM	1 (Skip to b)	2	1	2	mL	1 (Skip to H1a)	2 (Skip to H1a)
b.	<b>B2</b> 2 <sup>nd</sup> attempt:	hr	mins	: 1 = AM 2 = PM	1	2	1	2	mL	1	2

# TOENAIL CLIPPINGS FOR THE REPOSITORY SHOULD BE SHIPPED ONLY IF THE SAMPLE WAS NOT COLLECTED AT V4

#### **SECTION H:**

lect at V6)								
Did the participant consent to have biological samples (i.e., nail clippings) stored at NIDDK Biorepository?								



#### **TOENAIL** CLIPPING COLLECTION

- The collection of TOENAILS is preferred. DO NOT collect fingernail clippings. Also DO NOT collect toenails if participant has nail
  fungus, or discoloration causing pain or discomfort.
- STAINLESS STEEL NAIL CLIPPERS MUST BE USED TO COLLECT NAIL CLIPPINGS. Use small (pediatric size) stainless steel nail clippers (see Figure A) for younger participants and large stainless steel nail clippers (see Figure B) for older children. Both sizes are included in the CKiD starter package.
- Clean the blades of the nail clippers with SaniZide Plus prior to use (provided by the CBL).
- Whenever possible, the Study Coordinator should clip all (10) toenails, removing approximately 1 millimeter from each nail (See Figure C). Be prepared to collect flyaway nails.
- ➤ (To use nail clippers, see Figures A D). Refer to CKiD MOP Section 12 for further details.
- Carefully place the nail clippings into the cryovial (see Figure D). After using the nail clipper,
- > spray the clipper with **SaniZide Plus** and wipe with clean cloth.

## ALTHOUGH PICTURES DEPICT THE CLIPPING OF FINGERNAILS, AT VISIT 4 TOENAIL CLIPPINGS ARE PREFERRED.

Figure A





Figure B





Figure C



Provide 10 nail clippings that are at least 1 mm tall



	Yes	1 (END)
	No	
a.	How many toenail clippings were collected?	
b.	——————————————————————————————————————	, Nail fungus or discoloration causing pain or discomfort
	Nail fungus or discoloration	1 (END)
	Nails not long enough	2 <b>(END)</b>
	Participant Refused	-7 (END)
	Other	3