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

A1.	PARTICIPANTI	D: AFFIX ID LABEL OR E	NIER NUME	BER IF ID LABEL IS NOT AVAILABLE
			-   .	-
A2.	CKID VISIT #:			-
A3.	FORM VERSION	DN:	0 6	/ <u>0</u> <u>1</u> / <u>0</u> <u>8</u>
A4.	SPECIMEN CO	DLLECTION DATE:	/	D D Y Y Y Y
A5.	FORM COMPL	ETED BY (INITIALS):		<del></del>
A6.	Is this study vis	sit an accelerated visit?		1 2
<u>Sa</u> Se	e following sam mples: rum rum	ples should be collect Shipped to CBL CBL	Shipped: IMMEDIA Batched	TELY an, Apr, Jul or Oct)
loh	exol Blood	CBL	IMMEDIA	TELY
Uri	ne	CBL	IMMEDIA	TELY
Sa	mples:	ned for biological sam Shipped to: ) NIDDK Biosample F		Shipped:
Pla	sma (Biologica	l) NIDDK Biosample R	epository	Batched (Ship in Jul, Apr, Jul or Nov)
Hri	ne (Riological)	NIDDK Biosample R	onocitory	Ratched



(Ship in Jul, Apr, Jul or Nov)

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

B1.	ls p	articipant a female of child-bearing	potent	tial?
	Yes	S	1 (See	e PROMPT Below)
_		•		ICIPANTS OF CHILD-BEARING POTENTIAL ONLY. ITHIN 72 HOURS BEFORE GFR TESTING DATE.
B2.	a.	Urine pregnancy test date:		M D D Y Y Y
	b.	Urine pregnancy results: Positive Negative	1 <b>(EN</b>	ID; COMPLETE DISENROLLMENT FORM)
B3.	Yes	nis study visit a Make-Up GFR visit	1	ip to B5)
В3а.	Mak	ke-up GFR date:		M $D$ $D$ $Y$ $Y$ $Y$
MAKI	В	SEFORE THE IOHEXOL INFUSION	N.	L OF BLOOD IN THE PROVIDED SST LABELED B0 as not collected from a butterfly.)
B4.	Indi	cate reason(s) for a Make-Up GFR	visit: (	(Circle "Yes" or "No" for each):
	11.7.5	ofilius i o o	Yes	<u>No</u> 2
	Inat	nfiltrationbility to successfully draw 4 blood nples for lohexol	1	2
		er reasonSpecify:	1	2 (Skip to F1) (Skip to F1)

#### 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 central biochemistry laboratory (containers were shipped in batches to each site).



Pour 10 to 14.5 mL of urine into dark blue top urine collection tube and 5 to 14.5 mL into a second light blue top urine collection tube (provided by CBL).



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

(R	Sample Type equired 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>			
B5.	Urine Creatinine, Urine Protein, Urine Albumin (10.0 mL-14.5 mL in Dark Blue Top tube)	1 (skip to c→)	2	 (skip to B6)	i. Is this a first morning urine sample? Yes2 ii. Time of Collection:: 1 = am, 2 = pm	
B6.	Urine (Heavy Metals) (5.0-14.5 mL in Light Blue Top tube)	1 (skip to C1)	2	(skip to C1)	NA	

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



Complete Pre-Iohexol Infusion (B0) blood draw according to MOP instructions/flowchart on page 4. 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 on pre-iohexol infusion (B0) blood draw can be decreased by 1 ml.

#### SECTION C: PRE-IOHEXOL INFUSION (B0) 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 Pertain to the CKiD Participant:

## If participant consented to BIOLOGICAL samples:

Collect 20.5-23.0 mL if participant is < 30 kg OR 24.5-27.0 mL if participant is  $\ge 30 \text{ kg}$ .

If < 30 kg, immediately transfer (using 18-22 gauge needle) or draw:

- 12.5 mL into (2) Tiger-Top SSTs for CBL & NIDDK BR
- 3 mL into (1) PST for NIDDK Biosample Repository
- 1 mL into Tan-Top tube for heavy metals
- 1 mL in lavender-top tube for local CBC (tube not provided in CBL kit)
- 3 mL in appropriate tube (not provided) for local Renal Panel
- 2.5 mL of additional blood in SST for CBL (if initial sample is GROSSLY HEMOLYZED)

If ≥ 30 kg, immediately transfer (using 18-22 gauge needle) or draw:

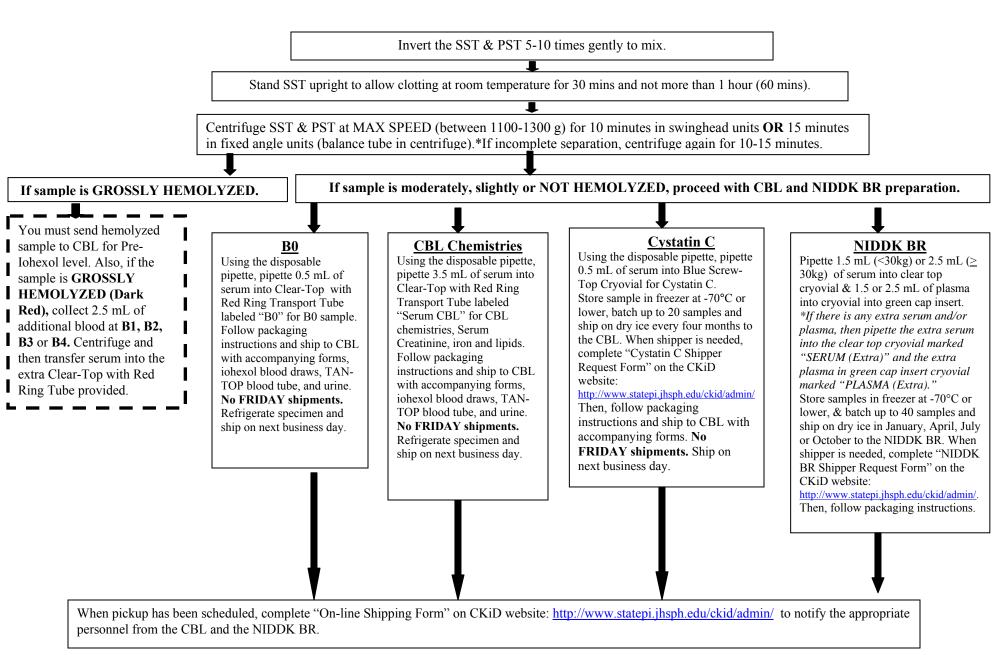
- 14.5 mL into (2) Tiger-Top SSTs for CBL & NIDDK BR
- 5 mL into (2) PST for NIDDK Biosample Repository
- 1 mL into Tan-Top tube for heavy metals
- 1 mL in lavender-top tube for local CBC (tube not provided in CBL kit)
- 3 mL in appropriate tube (not provided) for local Renal Panel
- 2.5 mL of additional blood in SST for CBL (if initial sample is GROSSLY HEMOLYZED)

2 If participant did NOT consent to BIOLOGICAL samples:
Collect 14.5-17.0 mL from all participants (regardless of weight) as specified below.

Immediately transfer (using 18 gauge needle) or draw:

- 9.5 mL into (2) Tiger-Top SSTs for CBL
- 1 mL into Tan-Top tube for heavy metals
- 1 mL in lavender-top tube for local CBC (tube not provided in CBL kit)
- 3 mL in another tube (not provided) for local Renal Panel
- 2.5 mL of additional blood in SST for CBL (if initial sample is GROSSLY HEMOLYZED)

#### PROCESSING OF PRE-IOHEXOL INUSION BLOOD FOR CBL & NIDDK BR



C1. ACTUAL TIME OF PRE-IOHEXOL INFUSION (B0) BLOOD DRAW \_\_\_\_ : \_\_\_\_ 1 = AM 2 = PM

#### PROMPT: IF SUSPECTED BLOOD DRAW ADVERSE EVENT (i.e., infection), complete ADVERSE EVENT (ADVR) Form

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

	2 = Diffi	cult Blood Dra	W	4 = Red Blood Cell Contamination 6 = Oversight		
(Re	Sample Type equired 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>			
C2.	Renal/Iron Chemistries (7.0 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 (3.0 mL in Local SST)	1 (skip to C5)	2	(skip to C5)	N/A	
C5.	Serum for Fasting Lipid Panel (1.5 mL in Tiger Top SST)	1 (skip to c→)	2	 (skip to C6)	Did the child fast after midnight?           Yes1           No2*	
C6.	Whole blood for Heavy Metals (1.0 mL in Tan-Top tube)	1 (skip to C7)	2	(skip to C7)	N/A	

<sup>\*</sup>If the child did not fast, the lipid results will be invalid and therefore will not be reported on the Nephron Lipid Report.

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://statepiaps.jhsph.edu/nephron/groups/aspproc/">https://statepiaps.jhsph.edu/nephron/groups/aspproc/</a>, click on "Report Menu" and choose the appropriate lab report (i.e., Selected Renal Panel Lab Variables Report.)

C7.	Did the participant consent to have biological sam Repository?	ples (i.e., serum, plasma and urine samples) stored at the NIDDK Biosample
	Yes	1
	No	2 (Skip to E2)

Reasons Code List*: 1= Not required		3 = Participant Refused	5 = Inadvertently Destroyed	
	2 = Difficult Blood Draw	4 = Red Blood Cell Contamination	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>				
C8.	Serum for NIDDK Biosample Repository (**3.0 mL or **5.0 mL of blood in Tiger Top SST)	1 (skip to c→)	2	(skip to C9)	Date Frozen:		
C9.	Plasma for NIDDK Biosample Repository (**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 D1)	Date Frozen:		

<sup>\*\*</sup> Collect 3.0 mL of whole blood for children < 30 kg and 5.0 mL for children ≥ 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 blue top 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.

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

Once the protease inhibitor tablets are completely dissolved, pour urine into 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) for 10 mins (swinghead units) – **OR** – 15 mins (fixed angle units).

Decant (pour off) the supernates (liquid reaction) into 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 up to 36 samples quarterly. When shipper 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 Heather Higgins, Sandra Ke and Alicia Wentz 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

Collection Refused Contamination Destroyed

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

## SECTION E: 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).

E2.	Was a 1 <sup>st</sup> morning urine protein to creatinine ratio assay performed at the clinical site's local laboratory?  Yes
	SECTION F: INFUSION SYRINGE WEIGHT
F1.	SCALE MUST FIRST BE ZEROED BEFORE WEIGHING. REMOVE ALUMINUM FOIL PRIOR TO WEIGHING THE SYRINGE. THE <u>SAME</u> SCALE MUST BE USED TO WEIGH THE SYRINGE <u>PRE AND POST</u> IOHEXOL 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.)
<u>PRI</u>	E AND POST SYRINGE WEIGHT MUST BE OBTAINED IN ORDER TO CALCULATE CHILD'S GFR.
	SECTION G: IOHEXOL – Refer to <u>Instructions for Iohexol Infusion and GFR Blood Draws Flow Chart on Page 12</u>
	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
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.
- > 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 10, 30, 120 & 300 MINUTES AFTER IOHEXOL INFUSION. FOR EXAMPLE, IF BLOOD IS DRAWN AT 33 MINS INSTEAD OF 30 MINS, DOCUMENT BLOOD DRAWN @ 33 MINS.
- > TIME SHOULD BE RECORDED IMMEDIATELY AFTER EACH BLOOD SAMPLE IS OBTAINED (i.e., B1, B2, B3, and B4).

ALL TIMES should be documented from the initial infusion time		(i) ACTUAL MINUTES on TIMER  (ii) ONLY if Timer malfunctions, record Clock Time using the same clock used for G1a		(iii) Difficult Blood Draw: Yes No	(iv) Blood Volume Collected (1 mL):	Centrifuged a	v) t Clinical Site: No
G2a.	<b>B1</b> 10 min:	minutes	: 1 = AM 2 = PM	1 (Skip to b) 2	mL	1 (Skip to G3a)	2 (Skip to G3a)
b.	<b>B1</b> 2 <sup>nd</sup> attempt:	minutes	: 1 = AM 2 = PM	1 2	mL	1	2

## INVERT TUBE 5-10 TIMES AFTER EACH BLOOD DRAW LET SST TUBE STAND 20-30 MINUTES (BUT NO LONGER THAN 1 HOUR)

CENTRIFUGE AT 1100-1300g (3000rpm with 10cm radius rotor) for 10 MINUTES IN SWING HEAD OR 15 MINUTES IN FIXED ANGLE

## POST VITALS SHOULD BE TAKEN IMMEDIATELY AFTER THE 10 MINUTE BLOOD DRAW 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:						
G3a.	Post- infusion blood pressure:	/					
b.	Post-infusion temperature:	1 = °C					
C.	Post-infusion number of heart beats per minute:						
d.	Post-infusion respirations per minute:						

	ALL TIMES should be documented from the initial infusion time	(i) ACTUAL HOURS/ MINUTES ON TIMER  (ii) ONLY if Timer malfunctions, record Clock Time using the same clock used for G1a		(iii) Difficult Blood Draw: Yes No	(iv) Blood Volume Collected (1 mL):	(v) Centrifuged at Clinical Site: Yes No	
G4a.	<b>B2</b> 30 min:	minutes	: 1 = AM 2 = PM	1 (Skip to b) 2	mL	1 (Skip to G5a)	2 (Skip to G5a)
b.	<b>B2</b> 2 <sup>nd</sup> attempt:	minutes	: 1 = AM 2 = PM	1 2	mL	1	2
G5a.	<b>B3 2 hrs</b> (120 min):	hr mins	: 1 = AM 2 = PM	1 (Skip to b) 2	mL	1 (Skip to G6a)	2 (Skip to G6a)
b.	<b>B3</b> 2 <sup>nd</sup> attempt:	hr mins	: 1 = AM 2 = PM	1 2	mL	1	2
G6a.	<b>B4 5 hrs</b> (300 min):	hr mins	: 1 = AM 2 = PM	1 (Skip to b) 2	mL	1 (Skip to H2)	2 (Skip to H2)
b.	<b>B4</b> 2 <sup>nd</sup> attempt:	hr mins	: 1 = AM 2 = PM	1 2	mL	1	2

IF FAMILY CONSENTED TO THE COLLECTION OF HAIR AND NAIL SAMPLES, BUT THE SAMPLES WERE NOT COLLECTED AT VISIT 1B, THEN PROCEED TO SECTION H (SEE QUESTIONS ON PAGE 13).





#### **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

Complete pre-iohexol infusion (B0) blood draw according to instructions

V2: after lipid panel is drawn, subject may resume drinking beverages containing sugar (if desired)

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 must occur within 3 months of initial visit

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

**V2:** remove 1 mL of blood and discard waste; draw 1 mL of blood into SST; record time of blood draw on Specimen Collection Form; flush used IV with at least 3 mL normal saline; CHECK POST-INFUSION VITALS

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

**V2:** Draw 1 mL of blood into SST at 10 mins (B1), 30 mins (B2), 120 mins (B3) and 300 mins (B4) post-infusion.

- \*Discard initial 1 mL blood waste at each blood draw
- \*Record the time each blood draw is completed

Following each blood draw:

gently invert tube 5-10 times

\*Flush after each blood draw with at least 3 mL normal saline

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.

Stand SST upright at room temp for 20-30 minutes, but no longer than 1 hour

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

Following packaging and shipping instructions; send urine, blood and copies of completed forms & 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



#### **SECTION H:**

H2.	Were nail clippings and hair samples previously co	ollected and shipped at V1b?
	Yes	1 (END)
	No	2
H3.	Did the participant consent to have biological sample	ples (i.e., nail clippings and hair samples) stored at NIDDK Biosample Repository?
	Yes	
	No	2 (END)

## **NAIL CLIPPING COLLECTION**

- Collection of fingernails is preferred. DO NOT collect fingernail clippings if the participant has acrylic nails, or nail fungus or discoloration causing pain or discomfort. If the participant cannot provide fingernail clippings, the Study Coordinator may clip the participant's toenails instead. FINGERNAILS AND TOENAILS SHOULD NOT BE COLLECTED IN THE SAME CRYOVIAL (collect one or the other).
- STAINLESS STEEL NAIL CLIPPERS MUST BE USED TO COLLECT NAIL CLIPPINGS. Use small (pediatric size) stainless
  steel nail clippers (see Figure A) for younger children 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) fingernails, 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.

Figure A







Figure B







Provide 10 nail clippings that are at least 1 mm tall

ı ı <del>-ı</del> .	Does the participant have acrylic nails?			
		Yes	1 (Skip to H6)	
		No	2	
H5.	We	re 10 fingernail clippings collected?		
		Yes	1 (Skip to I1)	
		No	2	
	a.	How many fingernail clippings were collected?		
	b.	b. Specify reason "10" fingernail clippings were not collected.		
		Nails not long enough	1 (Skip to H6)	
		Participant Refused	-7 (Skip to H6)	
		Other	2	
		i Chooify		
		i. Specify		
		i. Specify:		
H6.	We	re 10 toenail clippings collected?		
H6.	We		1 (Skip to I1)	
H6.	We	re 10 toenail clippings collected?		
H6.	We	re 10 toenail clippings collected? Yes	1 (Skip to I1)	
H6.		re 10 toenail clippings collected? Yes	1 (Skip to I1)	
H6.		re 10 toenail clippings collected? Yes	1 (Skip to I1)	
H6.	a.	re 10 toenail clippings collected? Yes No How many toenail clippings were collected? Specify reason "10" toenail clippings were not collected:	1 (Skip to I1)	
H6.	a.	re 10 toenail clippings collected? Yes	1 (Skip to I1) 2 (e.g., Nail fungus or discoloration causing pain or	
H6.	a.	re 10 toenail clippings collected? Yes No How many toenail clippings were collected? ——— Specify reason "10" toenail clippings were not collected: discomfort) Nail fungus or discoloration	1 (Skip to I1) 2 (e.g., Nail fungus or discoloration causing pain or 1 (Skip to I1)	
H6.	a.	re 10 toenail clippings collected? Yes No How many toenail clippings were collected? ——— Specify reason "10" toenail clippings were not collected: discomfort) Nail fungus or discoloration	1 (Skip to I1) 2 (e.g., Nail fungus or discoloration causing pain or 1 (Skip to I1) 2 (Skip to I1)	

#### **SECTION I: HAIR SAMPLE COLLECTION**

- STAINLESS STEEL SCISSORS MUST BE USED TO COLLECT HAIR SAMPLE. The scissors are included in the CKiD starter package.
- DO NOT collect hair sample if the participant has colored, straightened or chemically altered hair
- Clean blades of stainless steel scissors with SaniZide Plus prior to use.
- Use powder-free gloves.
- Refer to CKiD MOP Section 12 for further details.
  - Lift up the top layer of hair from the **occipital** region of the scalp (see Figure A). Isolate a small thatch of hair (at least 20 fibers) from this region (see Figure B).
  - Place the label with the participant's KID ID # tightly around all 20 strands of hair located at the distal end (furthest from the scalp) (see Figure C).
  - > Cut the hair sample off the participant's head as close to the scalp as possible (see Figure D).
  - Place cut thatch of hair inside aluminum foil (4 X 4) and fold the top of the foil to completely enclose the hair sample.
  - Place the aluminum foil inside a Ziplock bag (4 X 4) with the gel desiccant pellets in it and seal.
  - > Store sample at room temperature in a dark place prior to shipment.
  - After using the scissors, spray scissors with SaniZide Plus and wipe with clean cloth.

Figure A

**Occipital Region of Scalp** 

Figure B

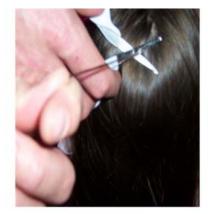


Figure C



Place the KID ID label tightly around all 20 strands.

Figure D



Cut the hair sample off the participant's head as close to the scalp as possible.

l1.	Does the participant have permed, dyed, colored, straig	htened or chemically altered hair?
	Yes	1 <b>(END)</b>
	No	2
l2.	Was the Study Coordinator able to cut at least 20 fibers	of hair from the occipital region?
	Yes	1 <b>(END)</b>
	No	2
	a. Specify reason "20" hair fibers were not collected:	
	Hair not long enough	1 (END)
	Participant Refused	7 (END)
	Other	2
	i. Specify:	