

BABY HUG FOLLOW-UP STUDY II

**ENROLLMENT**

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ 2. Current Clinic: \_\_\_\_\_  
SUBJECT\_ID SITE\_ID
3. Patient's Letter Code: \_\_\_\_\_ LETTER\_CD
4. Form Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ VISIT\_DT  
Month Day Year

**PART II: ENROLLMENT INFORMATION**

1. Did this child complete at least 24 mos of follow-up in BABY HUG Follow-up I? TREAT\_COMPLET\_1 Yes (1) No (2)  
(inel)

If No, Skip to Part III.1.

2. Has this child had a stem cell transplant since December 31, 2011? TRANSPLANT Yes (1) No (2)  
(inel)

If Yes, Skip to Part III.1 and complete Form 26.

3. Has informed consent been obtained? FOLLOWUP\_STUDY (1) (2)

If No, Skip to Part III.1.

4. Consent Information:

- A. Consent Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ CONSENT\_DT  
Month Day Year
- B. Consent for data file to include child's information? DATA\_CONSENT Yes (1) No (2) N/A
- C. Consent for blood specimens to be saved indefinitely? BLOOD\_SAVE\_CONSENT (1) (2)
- D. Consent for urine to be saved indefinitely? URINE\_SAVE\_CONSENT (1) (2)
- E. Consent for blood samples to be used for future research on sickle cell disease and related disorders? BLOOD\_FUTURE\_CONSENT (1) (2)
- F. Consent for urine samples to be used for future research on sickle cell disease and related disorders? URINE\_FUTURE\_CONSENT (1) (2)
- G. Consent for DNA testing to be performed on blood samples? DNA\_CONSENT (1) (2)
- H. Was Assent signed? ASSENT (1) (2) (3)

ID Number                      Visit                      Seq

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5. Follow-up Group: FOLLOWUP\_GROUP Active Passive  
(1) (2)

**PART III: COORDINATION**

1. Checked for completeness and accuracy:

- A. Certification number:      \_\_\_ \_\_\_ - \_\_\_ \_\_\_      CERT\_NO
  - B. Signature:                \_\_\_\_\_ CERT\_SIG
  - C. General Comments:       \_\_\_\_\_ GEN\_CMNT
- 

ID Number		Visit		
		-		

BABY HUG FOLLOW-UP STUDY II

**CENTRAL LAB COLLECTION  
ENTRY/Q12 MONTHS/EXIT**

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ SUBJECT\_ID                      2. Current Clinic: \_\_\_\_\_ SITE\_ID
3. Patient's Letter Code: \_\_\_\_\_ LETTER\_CD                      4. Visit: \_\_\_\_\_ VISIT\_NBR
5. Visit Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ VISIT\_DT  
                                    Month                      Day                      Year

**PART II: SPECIMEN COLLECTION**

**Please refer to Appendices A and B of the BHFUII Protocol for Lab Collection Requirements.**

1. Urine for Storage: (8-10 ml) (Entry/Exit Only)
- A. Label Number: \_\_\_\_\_ URINE\_STORED\_LABEL (1) Not Done URINE\_STORED\_ND
- B. Date Collected: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
URINE\_STORED\_DT                      Month                      Day                      Year
- C. Time Collected: \_\_\_\_\_ : \_\_\_\_\_ (24-hr clock)  
                                    H H                      M M  
URINE\_STORED\_HR      URINE\_STORED\_MN
2. Urine for Microalbumin: Creatinine (1-2 Cryovial ml): (Entry/Exit Only)
- A. Label Number: \_\_\_\_\_ URINE\_LABEL (1) Not Done URINE\_ND
- B. Date Collected: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
URINE\_DT                      Month                      Day                      Year
- C. Time Collected: \_\_\_\_\_ : \_\_\_\_\_ (24-hr clock)  
                                    H H                      M M  
URINE\_COL\_HR              URINE\_COL\_MN
3. Stored Blood Sample (5 ml EDTA lavender top) Entry/Exit Only:
- A. Label Number: \_\_\_\_\_ BLOOD\_LABEL (1) Not Done BLOOD\_ND
- B. Date Collected: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
BLOOD\_DT                      Month                      Day                      Year
- C. Time Collected: \_\_\_\_\_ : \_\_\_\_\_ (24-hr clock)  
                                    H H                      M M  
BLOOD\_COL\_HR              BLOOD\_COL\_MN

4. Cystatin C (1 ml red top):

A. Label Number:                      CYSTATIN\_LABEL ( 1 ) Not Done CYSTATIN\_ND

B. Date Collected:      -      -       
CYSTATIN\_DT      Month          Day          Year

C. Time Collected:      :      (24-hr clock)  
                                    H   H          M   M  
CYSTATIN\_COL\_HR      CYSTATIN\_COL\_MN

5. HbF (0.5 ml EDTA lavender top):

A. Label Number:                      HBF\_LABEL ( 1 ) Not Done HBF\_ND

B. Date Collected:      -      -       
HBF\_DT      Month          Day          Year

C. Time Collected:      :      (24-hr clock)  
                                    H   H          M   M  
HBF\_COL\_HR      HBF\_COL\_MN

6. Creatinine/BUN (4.0 ml red top at 12 mos) (1.0 ml red top at entry/exit):

A. Label Number:                      CREATININE\_LABEL ( 1 ) Not Done CREATININE\_ND

B. Date Collected:      -      -       
CREATININE\_DT      Month          Day          Year

C. Time Collected:      :      (24-hr clock)  
                                    H   H          M   M  
CREATININE\_COL\_HR      CREATININE\_COL\_MN

7. Pitted Cells (0.1 ml EDTA lavender top w/glutaraldehyde):

A. Label Number:                      CELL\_LABEL ( 1 ) Not Done CELL\_ND

B. Date Collected:      -      -       
CELL\_DT      Month          Day          Year

C. Time Prepared:      :      (24-hr clock)  
                                    H   H          M   M  
CELL\_COL\_HR      CELL\_COL\_MN



PEDIATRIC HYDROXYUREA CLINICAL TRIAL

LOCAL LABORATORY RESULTS

**Active – Entry, Q12 Months, Exit**

**Passive – Entry, Exit**

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ **SUBJECT\_ID**      2. Current Clinic: \_\_\_\_\_ **SITE\_ID**  
sequence # **VISIT\_NBR**
3. Patient's Letter Code: \_\_\_\_\_ **LETTER\_CD**      4. Visit: \_\_\_\_\_ - 0 0
5. Visit Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ **VISIT\_DT**  
Month                      Day                      Year

**PART II: LAB RESULTS**

1. A. White Blood Cell Count (WBC)      \_\_\_\_\_ . \_\_\_\_\_ K/mm<sup>3</sup>      **WBC**
- B. Red Blood Cell Count (RBC)      \_\_\_\_\_ . \_\_\_\_\_ M/mm<sup>3</sup>      **RBC**
- C. Hemoglobin      \_\_\_\_\_ . \_\_\_\_\_ g/dL      **HB**
- D. Hematocrit      \_\_\_\_\_ . \_\_\_\_\_ %      **PCV**
- E. Platelet Count      \_\_\_\_\_ . \_\_\_\_\_ K/mm<sup>3</sup>      **PLAT**
2. A. Differential Type:      (1) Manual      (2) Automated      **DIFFTYPE**
- B. Absolute Neutrophil Count      \_\_\_\_\_ . \_\_\_\_\_ K/mm<sup>3</sup>      **NEUT\_CT**
- C. Neutrophils (% of WBC)      \_\_\_\_\_ %      **NEUT\_PT**
- D. Lymphocytes (% of WBC)      \_\_\_\_\_ %      **LYMPH\_PT**
- E. Monocytes (% of WBC)      \_\_\_\_\_ %      **MONO\_PT**
- F. Nucleated Red Blood Cells (nRBC)\*      \_\_\_\_\_ **NRBC**
- \*1. If not 0, corrected WBC Count†      \_\_\_\_\_ . \_\_\_\_\_ K/mm<sup>3</sup>      **CWBC**

- G. Reticulocytes (% of RBC)      \_\_\_\_ . \_\_\_\_ %      **RETIC\_PT**
- H. Absolute Reticulocyte count      \_\_\_\_ . \_\_\_\_ K/mm<sup>3</sup>      **RETIC\_CT**
- I. MCV      \_\_\_\_ . \_\_\_\_ fL      **MCV**
3. A. LDH      \_\_\_\_\_ U/L      **LDH**
- B. Bilirubin, Total      \_\_\_\_ . \_\_\_\_ Mg/DL      **T\_BILI**
- C. Bilirubin, Direct      \_\_\_\_ . \_\_\_\_ Mg/DL      **D\_BILI**
- D. ALT      \_\_\_\_\_ U/L      **ALT**
4. A. Urine Osmolality (Active-Entry/Exit Only)      \_\_\_\_\_ mOsm/kg      **U\_OSMO**
1. Hours NPO      \_\_\_\_      **URN\_NPO**

**PART III: COORDINATION**

1. Checked for completeness and accuracy:

- A. Certification number:      \_\_\_\_ - \_\_\_\_      **CERT\_NO**
- B. Signature:      \_\_\_\_\_      **CERT\_SIG**
- C. General Comments:      \_\_\_\_\_      **GEN\_CMNT**

ID Number

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Visit

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Seq

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BABY HUG FOLLOW-UP STUDY II  
CLINICAL DATA REPORT

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ 2. Current Clinic: \_\_\_\_\_  
SUBJECT\_ID SITE\_ID
3. Patient's Letter Code: \_\_\_\_\_ LETTER\_CD
4. Abstraction Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ VISIT\_DT  
Month Day Year

**PART II: INTERVAL INFORMATION**

1. Visit: \_\_\_\_\_ M VISIT
2. Interval Start Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ INTERVAL\_START\_DT  
Month Day Year
3. Interval End Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ INTERVAL\_END\_DT  
Month Day Year
4. Any patient contact during this interval? PATIENT\_CONTACT Yes No\*  
(1) (2)
- \*A. If no, reason \_\_\_\_\_ PATIENT\_CONTACT\_RSN

\*If No, Skip to Part IX.

**PART III: HU USE**

1. Was the patient prescribed HU at any time during this interval? HU\_PRESCRIBED Yes\*\* No\*  
(1) (2)

\*If No, Skip to Part IV.

\*\*A. If yes, what was the:

1. Dose at the first time it was prescribed this interval: \_\_\_\_\_ . \_\_\_\_\_ mg/kg  
HU\_DOSE\_WEIGHT





- F. Absolute Neutrophil Count LAST\_NEUTROPHIL\_CNT  
 \_\_\_\_\_ . \_\_\_\_\_ K/mm<sup>3</sup> (1 ) Not Done LAST\_NEUTROPHIL\_NOT\_DONE
- G. Platelet Count LAST\_PLATELETS\_CNT  
 \_\_\_\_\_ . \_\_\_\_\_ K/mm<sup>3</sup>
- H. Red Blood Cell Count LAST\_RBC  
 \_\_\_\_\_ . \_\_\_\_\_ M/mm<sup>3</sup>

4. Were any of the following laboratory values obtained during this interval? LAB\_VALUES Yes No\*  
 (1) (2)

- \*A. If No, reason: NO\_LAB\_REASON
1. Not a routine part of care (1)
  2. Other (2)
    - a. If other, Specify: \_\_\_\_\_ NOLAB\_REASON\_SP

\*If No, Skip to Part V.

B. Creatinine:

1. Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ (1 ) Not Done  
CREATININE\_DT Month Day Year CREATININE\_NOT\_DONE

2. Value: \_\_\_\_\_ . \_\_\_\_\_ mg/dL CREATININE\_VALUE

C. ALT:

1. Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ (1 ) Not Done  
ALT\_DT Month Day Year ALT\_NOT\_DONE

2. Value: \_\_\_\_\_ IU/L ALT\_VALUE

D. GGT:

1. Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ (1 ) Not Done  
GGT\_DT Month Day Year GGT\_NOT\_DONE

2. Value: \_\_\_\_\_ u/L GGT\_VALUE

ID Number	Visit	Seq





2. MRI Date\*      -      -      (1) Not Done  
           MRI\_DT      Month              Day              Year              MRI\_NOT\_DONE

A0. Performed per protocol, results unknown      Yes      No  
    (1)      (2)      MRI\_UNKNOWN

If Yes, skip to 3.

\*A. If MRI done, result:

**CHECK THE MOST SEVERE RESULT**      MRI\_RESULTS

- Normal (1)
- Silent Infarct(s) (gliosis) (2)
- Stroke (CVA or thrombosis) (3)
- Hemorrhage (subarachnoid or subdural) (4)
- Other (5)^

1. ^If Other, Specify: \_\_\_\_\_ MRI\_RESULTS\_SP

**PASSIVE SUBJECTS: If DONE, MRI Performed closest to age 10 should be sent for central review. Complete Form 33 and submit CD to DCC**

3. MRA Date\*      -      -      (1) Not Done  
           MRA\_DT      Month              Day              Year              MRA\_NOT\_DONE

A0. Performed per protocol, results unknown      Yes      No  
    (1)      (2)      MRA\_UNKNOWN

If Yes, skip to 4.

\*A. If MRA done, any result abnormal?      Yes      No  
    (1)      (2)      MRA\_ABNORMAL

**PASSIVE SUBJECTS: If DONE, MRA Performed closest to age 10 should be sent for central review. Complete Form 33 and submit CD to DCC**

4. PFTs Date\*      -      -      (1) Not Done  
           PFT\_DT      Month              Day              Year              PFT\_NOT\_DONE

\*A. If Pulmonary Function Tests done, any result abnormal?      Yes      No  
    (1)      (2)      PFT\_ABNORMAL

**PASSIVE SUBJECTS: Complete Form 31**

ID Number      Visit      Seq  
    -



**PART VI: OTHER PROCEDURES**

1. EEG Date\* \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ (1 ) Not Done  
EEG\_DT                      Month                      Day                      Year                      EEG\_NOT\_DONE

Yes      No

\*A. If EEG done, any result abnormal?                      EEG\_ABNORMAL                      (1 )                      (2 )

2. CT Date\* \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ (1 ) Not Done  
CT\_DT                      Month                      Day                      Year                      CT\_NOT\_DONE

Yes      No

\*A. If CT done, any result abnormal?:                      CT\_ABNORMAL                      (1 )                      (2 )

3. Neuropsych Date\* \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ (1 ) Not Done  
NEUROPSYCH\_DT                      Month                      Day                      Year                      NEUROPSYCH\_NOT\_DONE

NEUROPSYCH\_ABNORMAL      Yes^      No

\*A. If neuropsychology testing done, any results abnormal?                      (1 )                      (2 )

^1. Specify test: \_\_\_\_\_ NEUROPSYCH\_SP

Yes\*      No

4. Other clinical tests done:                      OTHER\_TEST                      (1 )                      (2 )

\*A. If Yes, specify: \_\_\_\_\_ OTHER\_TEST\_SP

ID Number	Visit	Seq									
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4. Pain

A. Has the child experienced pain (defined as pain lasting four hours or more without other obvious cause for which medication such as ibuprofen, acetaminophen, or acetaminophen with opioid was taken for relief) even if not seen by a medical professional during the interval? PAIN2 Yes\* No  
(1) (2)

\*1. If yes, how many episodes of pain has the patient experienced during this interval? PAIN\_EPISODES  
\_\_\_\_\_

5. Surgery

A. Did the patient have at least one surgery during this interval? SURGERY Yes\* No  
(1) (2)

\*1. If yes, identify the type of each surgery and give date:

a. Tonsillectomy, Adenoidectomy or both TONSILLECTOMY\_ND  
(1) Not Done  
 Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ TONSILLECTOMY\_DT  
                   Month                  Day                  Year

b. Splenectomy (open or aparoscopic) SPLENECTOMY\_ND  
(1) Not Done  
 Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ SPLENECTOMY\_DT  
                   Month                  Day                  Year

c. Cholecystectomy and/or ERCP CHOLECYSTECTOMY\_ND  
(1) Not Done  
 Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ CHOLECYSTECTOMY\_DT  
                   Month                  Day                  Year

d. Ear tubes, hernia repair, dental rehabilitation EAR\_NOT\_DONE  
(1) Not Done  
 Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ EAR\_DT  
                   Month                  Day                  Year

e. Other Yes^ No  
(1) (2)  
SURGERY\_OTHER

^1. If other, specify: SURGERY\_OTHER\_SP  
\_\_\_\_\_

ID Number		Visit		-	Seq



C. Was iron overload assessed during this interval? Yes\* No  
(1) (2)  
IRONOVL

\*If yes,

1. Ferritin (highest value in interval) \_\_\_\_\_ ng/ml FERRITIN\_HIGH  
Not done

2. Ferriscan or MRI \_\_\_\_\_ . \_\_\_\_\_ gm/gm dn weight of liver FERRISCAN\_MRI (1)  
FERRISCAN\_MRI\_ND

3. Liver Bx \_\_\_\_\_ . \_\_\_\_\_ gm/gm dn weight of liver LIVER\_BX (1)  
LIVER\_BX\_ND

D. Was iron chelation therapy prescribed during this interval? Yes\* No  
(1) (2)  
IRONTHPY

\*If yes,

Desferal (Deferioxamine) (1)

Ex Jade (Deferrisirox) (2)

L1 (Deferitronine) (3) IRON\_MED

ID Number	Visit	Seq



3. A. Was the spleen reported to be palpable below the costal margin at any time during this interval? SPLEEN\_PALPABLE Yes No  
(1) (2)

If No, Skip to Part IX.

- B. On what date was it the largest (most centimeters below costal margin): SPLEEN\_LARGEST\_DT  
 \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
 Month Day Year

Write the largest value below:

1. Mid-clavicular line: MID\_CLAVICULAR MID\_CLA\_NOTDONE  
 \_\_\_\_\_ . \_\_\_\_\_ cm below costal margin (1) Not Done
2. Anterior axillary line: ANTEROR\_AXILLARY ANT\_AXI\_NOTDONE  
 \_\_\_\_\_ . \_\_\_\_\_ cm below costal margin (1) Not Done

- C. Was the child diagnosed with acute splenic sequestration during this interval? DIAG\_SPLENIC\_SEQU Yes No  
(1) (2)

**PART IX: COORDINATION**

1. Checked for completeness and accuracy:

- A. Certification number: \_\_\_\_\_ - \_\_\_\_\_ CERT\_NO
- B. Signature: \_\_\_\_\_ CERT\_SIG
- C. General Comments: \_\_\_\_\_ GEN\_CMNT
- \_\_\_\_\_

ID Number	Visit	Seq

BABY HUG FOLLOW-UP STUDY II

**EXIT FORM**

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ SUBJECT\_ID      2. Current Clinic: \_\_\_\_\_ SITE\_ID
3. Patient's Letter Code: \_\_\_\_\_ LETTER\_CD
4. Date of Form Submission: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
VISIT\_DT      Month      Day      Year

**PART II: END OF TREATMENT**

1. End of Follow-up Participation      Yes      No
- A. Completed Follow-up Study II      END\_PARTICIPATION      (1)      (2)

**IF YES, GO TO PART III. IF NO, ANSWER IIB 1-6 AND II.2.**

- B. Reason for study exit:
- |                   |   |      |  |     |
|-------------------|---|------|--|-----|
| <b>INACTIVE</b>   | 1. Inactive follow-up status                            | (1)  |  | (2) |
| <b>RELOCATION</b> | 2. Permanent relocation to area with no BABY HUG Clinic | (1)  |  | (2) |
| <b>WITHDRAW</b>   | 3. Withdrew consent                                     | (1)  |  | (2) |
| <b>TRANSPLANT</b> | 4. Stem Cell Transplant                                 | (1)  |  | (2) |
| <b>DEATH</b>      | 5. Death  | (1)  |  | (2) |
| <b>OTHER</b>      | 6. Other condition requiring end of participation       | (1)* |  | (2) |
- a. \*Specify: OTHER\_SP \_\_\_\_\_

**If Yes, complete Form 26.**

2. Date of last contact with family: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
LAST\_CONTACT\_DT      Month      Day      Year

**If participant had stem cell transplant, record the date started conditioning for the transplant as the last contact date.**

ID Number	Visit	Seq
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BABY HUG FOLLOW-UP STUDY II

**TRANSCRANIAL DOPPLER (TCD) EXAM**

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ **SUBJECT\_ID**      2. Current Clinic: \_\_\_\_\_ **SITE\_ID**
3. Patient's Letter Code: \_\_\_\_\_ **LETTER\_CD**
4. Procedure Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ **VISIT\_DT**  
   Month                      Day                      Year

**PART II: EQUIPMENT**

1. TCD examiner's last name:  
 \_\_\_\_\_ **RDR46**

2. Patient's position during exam **PTNTPOS**
- Sitting (1)
  - Lying on exam table (2)
  - Other (3)\*
  - No information available (4)

\*A. Specify: \_\_\_\_\_ **POS\_SP**

**PART III: EXAMINATION PERFORMANCE**

1. Completeness of exam **COMPEXAM**
- Attempted, but no data collected (1)\*
  - Started, but aborted with some data (2)^\*
  - Complete exam given (3)^
  - No information available (4)

ID Number

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- \*A. Reason for incomplete exam INCEXAM
- Patient uncooperative (1)
- Other (2)\*\*

\*\*1. Specify INCEX\_SP \_\_\_\_\_  
\_\_\_\_\_

^B. TCD Label TCD\_LBL \_\_\_\_\_

**PART IV: COORDINATION**

1. Checked for completeness and accuracy:

A. Certification number: \_\_\_\_\_ - \_\_\_\_\_ CERT\_NO

B. Signature: \_\_\_\_\_ CERT\_SIG

C. General Comments: GEN\_CMNT

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

ID Number

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- F. Neck NECK
- No adenopathy (1)
  - Small shotty cervical nodes (< 1 cm) (2)
  - Enlarged nodes (3)\*
  - \*1. If enlarged, describe largest \_\_\_\_\_ cm NECK\_NDLGST
  - \*2. Site:
    - Right (1) NECK\_RT
    - Left (2)
- G. Chest (check all that apply)
- 1. Clear to auscultation (normal) (1) CHEST\_CLEAR
  - 2. Retractions (1) CHEST\_RETRACTIONS
  - 3. Transmitted upper airway sounds (1) CHEST\_TRANSAIRWAY
  - 4. Ronchi or Rales (1) CHEST\_RONCHI
  - 5. Wheezing (1) CHEST\_WHEEZING
  - 6. Other (1)\* CHEST\_OTHER
  - \*a. Specify: \_\_\_\_\_ CHEST\_SPECIFY
- H. Cardiac CARDIAC
- S1S2 with no murmur (normal) (1)
  - S1S2 with systolic ejection murmur (flow murmur) (2)
  - Other abnormal heart sound or murmur (3)\*
  - \*1. Describe: \_\_\_\_\_ CARDIAC\_OTHER
- I. Abdomen ABDOMEN
- Soft (non-tender) (1)
  - Tender (2)
  - Rebound and/or Guarding (3)
- J. Liver LIVER
- Not enlarged (1)
  - Enlarged (2)\*
  - \*1. \_\_\_\_\_cm below right costal margin in midclavicular line LIVRCM

ID Number	Visit	Seq







BABY HUG FOLLOW-UP STUDY II  
**QUESTIONNAIRE**

**PART I: IDENTIFYING INFORMATION**

1. Patient’s ID Number: \_\_\_\_\_ SUBJECT\_ID                      2. Current Clinic: \_\_\_\_\_ SITE\_ID
3. Patient’s Letter Code: \_\_\_\_\_ LETTER\_CD    4. Visit: \_\_\_\_\_ VISIT\_NBR
5. Form Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
VISIT\_DT                      Month                      Day                      Year

This is a composite questionnaire to be done annually on patients. Please ask the questions as they are written and record the responses on this form (this is your source document). Please read ALL of the possible responses before accepting a patient’s answer. Ensure that there is privacy during the conversation.

**PART II: SLEEP**

Please answer these questions regarding the behavior of your child during sleep and wakefulness. The questions apply to how your child acts in general during the past month, not necessarily during the past few days since these may not have been typical if your child has not been well. A “Y” means “yes,” “N” means “no,” and “DK” means “don’t know.”

- |  | Yes     | No      | Don't Know<br>(DK) |               |
|--|---------|---------|--------------------|---------------|
| 1. While sleeping, does your child:  |         |         |                    |               |
| A. snore more than half the time?  | (1)     | (2)     | (3)                | SNR_HLFT2     |
| B. always snore?   | (1)     | (2)     | (3)                | SNORE2        |
| C. snore loudly?   | (1)     | (2)     | (3)                | SNR_LOUD2     |
| D. have heavy or loud breathing?   | (1)     | (2)     | (3)                | BRTHLOUD2     |
| E. have trouble breathing, or struggle to breathe?   | (1)     | (2)     | (3)                | BRTHTRBL2     |
| <br>2. Have you ever seen your child stop breathing during the night?                                    | <br>(1) | <br>(2) | <br>(3)            | <br>BRTHSTP2  |
| 3. Does your child:  |         |         |                    |               |
| A. tend to breathe through the mouth during the day?   | (1)     | (2)     | (3)                | BRTHMTH2      |
| B. have a dry mouth on waking up in the morning?   | (1)     | (2)     | (3)                | DRYMTH2       |
| C. occasionally wet the bed?   | (1)     | (2)     | (3)                | WETBED2       |
| D. wake up feeling unrefreshed in the morning?   | (1)     | (2)     | (3)                | WKPRFRSH2     |
| E. have a problem with sleepiness during the day?  | (1)     | (2)     | (3)                | SLP_PROB2     |
| <br>4. Has a teacher, supervisor or other adult commented that your child appears sleepy during the day? | <br>(1) | <br>(2) | <br>(3)            | <br>SLPY_DAY2 |

ID Number		Visit		Seq



- |  | Yes        | No        |          |
|--|------------|-----------|----------|
| D. Bell alarm  |            |           |          |
| 1. Tried   | (1)*       | (2)       | ALARM_T  |
| *a. Successful   | (1)        | (2)       | ALARM_S  |
| E. Medication DDAVP (desmopressin)   |            |           |          |
| 1. Tried   | (1)*       | (2)       | DDAVP_T  |
| *a. Successful   | (1)        | (2)       | DDAVP_S  |
| F. Medication Tofranil (imipramine)  |            |           |          |
| 1. Tried   | (1)*       | (2)       | TOFR_T   |
| *a. Successful   | (1)        | (2)       | TOFR_S   |
| G. Therapy or counseling?  |            |           |          |
| 1. Tried   | (1)*       | (2)       | THRPY_T  |
| *a. Successful   | (1)        | (2)       | THRPY_S  |
| 4. After going to bed, has your child ever woken up at night to urinate in the bathroom during the last 3 months?    | (1)        | (2)       | NT_URI   |
| <div style="border: 1px solid black; padding: 2px; display: inline-block;">If No, skip to 6.</div>                   |            |           |          |
| 5. If your child urinates in the bathroom during the night, how often?   |            |           | NTURIFRQ |
| Rarely   |            | (1)       |          |
| Several times month (but less than once a week)  |            | (2)       |          |
| 1-2 times per week   |            | (3)       |          |
| 3-5 times per week   |            | (4)       |          |
| Every night or almost every night  |            | (5)       |          |
| 6. Does your child urinate in his or her clothes during the day?   | Yes<br>(1) | No<br>(2) | WET_DAY  |
| 7. A. Does your child have an immediate family member (parent, sibling) with history of bedwetting when a child?     | (1)        | (2)       | WTBD_FM  |
| <div style="border: 1px solid black; padding: 2px; display: inline-block;">If No, skip to 8.</div>                   |            |           |          |
| B. Does the family member with bed wetting have sickle cell disease?   | Yes<br>(1) | No<br>(2) | FM_SCD   |
| 8. Does your child drink any caffeinated beverages (coffee, tea, energy drinks, soda with caffeine) after 4:00 p.m.? | (1)        | (2)       | DRK_CAFF |
| 9. Is your child a deep sleeper?   | (1)        | (2)       | DP_SLPER |
| 10. Does your child have constipation?   | (1)        | (2)       | CONST    |

ID Number                      Visit                      Seq  
            -



**PART V: PRIAPISM – ask of parents/guardians of MALE patients only (regardless of age)**

1. Is the child male? Yes (1) No (2) **MALE**

If No, skip to Part VI.

A. Have you ever heard the word priapism before? (1) (2) **PRIAPWRD**

If No, skip to “Read to Patient and Parent.”

B. Where have you heard the word priapism before? (Check all that apply)

1. Doctor or nurse	(1)	<b>SRC_DR</b>
2. Friend or relative	(1)	<b>SRC_FR</b>
3. Written information	(1)	<b>SRC_INFO</b>
4. Other	(1)*	<b>SRC_OTH</b>
*a. Specify: _____		
		<b>SRCOTHSP</b>

**READ TO PATIENT AND PARENT:** *Priapism is a painful erection of the penis. It may last minutes to hours. It is more common in boys and men with sickle cell disease.*

C. Has your son ever had a painful unwanted erection of the penis that lasted 30 minutes or more? Yes (1) No (2) **PRIAP30M**

If No, skip to Part VI.

D. Has your son ever had a painful unwanted erection of the penis that lasted 4 hours or more? (1) (2) **PRIAP4HR**

For the next two questions, read all of the answers, then ask for one best answer.

E. How many episodes of priapism has your son had in the last year? **PRIAPEP1**

None	(1)	
One	(2)	
2 to 5	(3)	
6 to 20	(4)	
More than 20	(5)	
Do not know	(6)	

F. How many episodes of priapism did your son have before the last year? **PRIAPEP2**

None	(1)	
One	(2)	
2 to 5	(3)	
6 to 20	(4)	
More than 20	(5)	
Do not know	(6)	

G. How old was your son when the first episode happened? \_\_\_\_\_ years old **PRIAPAGE**

ID Number	Visit	-	Seq



BABY HUG FOLLOW-UP STUDY II

**SPECIAL TESTS (AGE 10)**  
**(All Active Subjects)**

**PART I: IDENTIFYING INFORMATION**

1. Subject ID Number: \_\_\_\_\_ **SUBJECT\_ID**      2. Current Clinic: \_\_\_\_\_ **SITE\_ID**
3. Subject Letter Code: \_\_\_\_\_ **LETTER\_CD**
4. Visit Start Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
**VISIT\_DT**      Month      Day      Year

**PART II: SPECIAL TESTS AND PROCEDURES**

1. Liver/Spleen Scan Performed? \_\_\_\_\_ **LIVER\_SCAN**      Yes (1)      No (2)

**IF YES, RECORD DATE PERFORMED AND COMPLETE FORM 21.**

- A. Date Liver/Spleen Scan Performed: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
**LIVER\_SCAN\_DT**      Month      Day      Year

2. Abdominal Sonogram Performed? \_\_\_\_\_ **ABDOMINAL\_SONO**      Yes (1)      No (2)

**IF YES, RECORD DATE PERFORMED AND COMPLETE FORM 23.**

- A. Date Abdominal Sonogram Performed: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
**ABDOMINAL\_SONO\_DT**      Month      Day      Year

3. TCD Performed? \_\_\_\_\_ **TCD**      Yes (1)      No (2)

**IF YES, RECORD DATE PERFORMED AND COMPLETE FORM 13.**

- A. Date TCD Performed: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
**TCD\_DT**      Month      Day      Year

4. PFT Performed? PFT Yes (1) No (2)

**IF YES, RECORD DATE PERFORMED AND COMPLETE FORM 31.**

A. Date PFT Performed: PFT\_DT Month - Day - Year

5. Cardiac Echocardiogram Performed? CARDIAC Yes (1) No (2)

**IF YES, RECORD DATE PERFORMED AND COMPLETE FORM 32.**

A. Date Echocardiogram Performed: CARDIAC\_DT Month - Day - Year

6. MRI/MRA Performed? MRIMRA Yes (1) No (2)

**IF YES, RECORD DATE PERFORMED AND COMPLETE FORM 33.**

A. Date MRI/MRA Performed: MRIMRA\_DT Month - Day - Year

7. Vineland Performed? VINELAND Yes (1) No (2)

**IF YES, RECORD DATE PERFORMED AND COMPLETE FORM 27.**

A. Date Vineland Performed: VINELAND\_DT Month - Day - Year

8. Peds QOL Performed? PEDSQOL Yes (1) No (2)

**IF YES, RECORD DATE PERFORMED AND COMPLETE FORM 29.**

A. Date Peds QOL Performed: PEDSQOL\_DT Month - Day - Year

ID Number		Visit		-	Seq

9. Connor CPT II Performed? CONNORCPT2 Yes No  
(1) (2)

**IF YES, RECORD DATE PERFORMED AND COMPLETE FORM 30.**

A. Date Connor CPT II Performed: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
CONNORCPT2\_DT Month Day Year

10. WISC IV Performed? WISC4 Yes No  
(1) (2)

**IF YES, RECORD DATE PERFORMED AND COMPLETE FORM 28.**

A. Date WISC IV Performed: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
WISC4\_DT Month Day Year

**PART III: COORDINATION**

1. Checked for completeness and accuracy:

- A. Certification number: \_\_\_\_\_ - \_\_\_\_\_ CERT\_NO
- B. Signature: \_\_\_\_\_ CERT\_SIG
- C. General Comments: GEN\_CMNT \_\_\_\_\_

ID Number	Visit	Seq

BABY HUG FOLLOW-UP STUDY II

LIVER-SPLEEN SCAN PERFORMANCE

PART I: IDENTIFYING INFORMATION

1. Patient's ID Number: **SUBJECT\_ID** \_\_\_\_\_ 2. Current Clinic: **SITE\_ID** \_\_\_\_\_
3. Patient's Letter Code: \_\_\_\_\_ **LETTER\_CD**
4. Visit Date **VISIT\_DT** \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
Month Day Year

PART II: SCAN SPECIFICS

1. Camera Manufacturer: **CAMTYPE**
2. Camera Model: **CAMMODEL**
3. Collimator: **COLLIMAT**
4. Supplier of TC-Sulfur Colloid: **SUPCOLLD**
5. Dose Injected: **DOSINJ44**  .  mCi
6. Time of Injection (24-hour clock): **INJ44HR**  :  **INJ44MN**
7. Time Imaging Started: **IMSTRHR**  :  **IMSTRMN**
8. Time Imaging Completed: **IMCOMHR**  :  **IMCOMMN**
9. Camera Angle: **CAMANGLE**  °
10. True Posterior Imaging Time (min:sec): **ANTPOSMN**  :  **ANTPOSSC**
11. Right Posterior Oblique Image Counts: **OBLIMCNT**
12. Film Label: **LSSCNLBL**
13. Adequacy of Imaging (Answer both questions):
- |    |                                       | Yes | No  |
|----|---------------------------------------|-----|-----|
| A. | 400 K Image adequate: <b>AOI400K</b>  | (1) | (2) |
| B. | Timed Image adequate: <b>AOITIMED</b> | (1) | (2) |

**PART III: QUANTITATIVE ASSESSMENT**

1. 400K Image

A. Anterior View

1. Spleen

a. Total Counts: **KASPLTOT**

b. # Pixels in ROI: **KASPLPIX**

c. Counts/Pixel: **KASPLCNT**

2. Liver

a. Total Counts: **KALIVTOT**

b. # Pixels in ROI: **KALIVPIX**

c. Counts/Pixel: **KALIVCNT**

B. Posterior View

1. Spleen

a. Total Counts: **KPSPLTOT**

b. # Pixels in ROI: **KPSPLPIX**

c. Counts/Pixel: **KPSPLCNT**

2. Liver

a. Total Counts: **KPLIVTOT**

b. # Pixels in ROI: **KPLIVPIX**

c. Counts/Pixel: **KPLIVCNT**

C. Spleen/Liver Ratio

1. Total Counts: **KSLRTTOT**  .

2. Counts/Pixel: **KSLRTCNT**  .

ID Number                      Visit                      Seq

                      -

2 Timed Image

A. Left Anterior Oblique View

1. Spleen

a. Total Counts: **TASPLTOT**

b. # Pixels in ROI: **TASPLPIX**

c. Counts/Pixel: **TASPLCNT**

2. Liver

a. Total Counts: **TALIVTOT**

b. # Pixels in ROI: **TALIVPIX**

c. Counts/Pixel: **TALIVCNT**

B. Right Posterior Oblique View

1. Spleen

a. Total Counts: **TPSPLTOT**

b. # Pixels in ROI: **TPSPLPIX**

c. Counts/Pixel: **TPSPLCNT**

2. Liver

a. Total Counts: **TPLIVTOT**

b. # pixels in ROI: **TPLIVPIX**

c. Counts/Pixel: **TPLIVCNT**

C. Spleen/Liver Ratio

1. Total Counts: **TSLRRTOT**  .

2. Counts/Pixel: **TSLRTCNT**  .

ID Number                      Visit                      Seq

                      -







**BABY HUG FOLLOW-UP STUDY II  
 SERIOUS ADVERSE EVENT  
 (ACTIVE GROUP ONLY)**

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ 2. Current Clinic: \_\_\_\_\_  
SUBJECT\_ID SITE\_ID
3. Patient's Letter Code: \_\_\_\_\_ LETTER\_CD
4. Reporting Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
VISIT\_DT Month Day Year

**PART II: EVENT PERIOD**

1. Date of Event
- A. Event Start Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ START\_DT  
Month Day Year
- B. Event End Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ E\_END\_DT  
Month Day Year
2. Qualifying Procedure (Event must have occurred during the 5 days following an "Active" assessment procedure.) Please note all that apply:
- A. Liver/Spleen Scan \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
LIVER\_SPLEEN\_DT Month Day Year LIVER\_SPLEEN\_NA
- B. Abdominal Sonogram \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
ABD\_SONO\_DT Month Day Year ABD\_SONO\_NA
- C. WISC IV \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
WISC4 Month Day Year WISC4\_NA
- D. Blood Specimens \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
BLOOD\_SPEC\_DT Month Day Year BLOOD\_SPEC\_NA
- E. TCD \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
TCD\_DT Month Day Year TCD\_NA
- F. PFT \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
PFT\_DT Month Day Year PFT\_NA
- G. Cardiac Echocardiogram \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
CARDIAC\_DT Month Day Year CARDIAC\_NA

- H. MRI MRI\_DT \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
 Month Day Year MRI\_NA
- I. MRA MRA\_DT \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
 Month Day Year MRA\_NA
- J. Vineland VINELAND\_DT \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
 Month Day Year VINELAND\_NA
- K. Connor CPT II CONNORCPT2\_DT \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
 Month Day Year CONNORCPT2\_NA
- L. Peds QOL PEDSQOL\_DT \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ ( 1 ) N/A  
 Month Day Year PEDSQOL\_NA

**PART III: SAE**

1. Please indicate all diagnoses:		YES	NO
A. Acute Chest Syndrome	HX_ACS	( 1 )	( 2 )
B. Splenic Sequestration Crisis	HXSPLSEQ	( 1 )	( 2 )
C. Prolonged Hospitalization (greater than 7 days)	LONGHOSP	( 1 )	( 2 )
D. Stroke or TIA	HX_STROKE_TIA	( 1 )	( 2 )
E. Life Threatening Event	LIFE_THREAT_EVT	( 1 )	( 2 )
1. Specify:	LIFE_THREAT_EVT_SP		
F. Death	HX_DEATH	( 1 )	( 2 )
G. ICU Admission	ICU	( 1 )	( 2 )

**PART IV: ADDITIONAL DIAGNOSIS INFORMATION**

If PART III, Item 1A is YES, answer 1. Otherwise, skip to 2.

1. Acute Chest Syndrome	None	1 Lobe	>1 Lobe	N/A	
A. New Infiltrate	( 1 )	( 2 )	( 3 )	( 4 )	ACSNINF
B. O <sub>2</sub> % Saturation on Room Air at Presentation	_____	_____	. _____%		ACSSRAP
C. Oxygen Administered	_____	_____	. _____L		ACSOXADM
D. Mechanical Ventilation	Yes ( 1 )		No ( 2 )		ACSMVENT

If PART III, Item 1B is YES, answer 2. Otherwise, skip to 3.

ID Number

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2. Splenic Sequestration

A. Spleen size below LCM **prior** to SAE

**SPLNSIZE\_PRIOR**

<2 cm	2-4 cm	4-6 cm	6-8 cm	>8 cm
(1)	(2)	(3)	(4)	(5)

B. Spleen size below LCM **during** SAE

**SPLNSIZE\_DURING**

<2 cm	2-4 cm	4-6 cm	6-8 cm	>8 cm
(1)	(2)	(3)	(4)	(5)

C. Nadir hemoglobin

\_\_\_\_ . \_\_\_\_ gm/dL

**SPLNHMGL**

D. Platelet count at time of nadir hemoglobin

\_\_\_\_ k/ $\mu$ L

**SPLPTCNT**

If PART III, Item 1C is YES, answer 3. Otherwise, skip to 4.

3. Prolonged Hospitalization

A. Reason:

**LONGHOSP\_SP**

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If PART III, Item 1D is YES, answer 4-5. Otherwise, skip to Part V.

4. (Stroke or TIA) Findings of

A. Loss of consciousness

**LOS\_CONS**

YES

NO

N/A

(1)

(2)

(3)

B. Change in mental status

**CHG\_MENT**

(1)

(2)

(3)

C. Loss of or difficulty with speech or vocalization

**SPEECH**

(1)

(2)

(3)

D. Paralysis or weakness

**PARALYS**

(1)

(2)

(3)

E. Difficulty with swallowing

**DIFFSWAL**

(1)

(2)

(3)

F. Difficulty with vision

**DIFF\_SEE**

(1)

(2)

(3)

G. Loss of balance or dizziness

**BALANCE**

(1)

(2)

(3)

H. Seizures

**SEIZURE**

(1)

(2)

(3)

I. Headache

**HEADACHE**

(1)

(2)

(3)

ID Number

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5. Results of Imaging Tests		Normal	Abnormal	Not Done
A. MRI of brain	F50MRI	(1)	(2)	(3)
B. CT scan of brain	F50CTBR	(1)	(2)	(3)
C. PET scan of brain	F50PTBR	(1)	(2)	(3)
D. MRA cerebral vasculature	F50MRA	(1)	(2)	(3)
E. Transcranial Doppler	F50TCD	(1)	(2)	(3)
F. Arteriogram	F50ARTGR	(1)	(2)	(3)

**PART V: DIAGNOSIS/PROBLEM SEVERITY AND ATTRIBUTION**

Complete PART V for each item in PART III checked YES.

PROBLEM	ONSET_DT	NUMDAYS	SEVERITY	ATTR_TRT	DIAGUNXP
1. Diagnosis/ Problem	2. Date of Onset	3. Number of Days	4. <sup>1</sup> Severity	5. <sup>2</sup> Attribution to Study Treatment	6. <sup>3</sup> Diagnosis Unexpected

<u><sup>1</sup>Severity</u>	<u><sup>2</sup>Attribution to Study Test</u>	<u><sup>3</sup>Diagnosis Unexpected</u>
1. Mild	1. Definite (clearly related)	1. Yes
2. Moderate	2. Probably (likely related)	2. No
3. Severe	3. Possible (may be related)	3. N/A
4. Life threatening	4. Unlikely (doubtfully related)	
5. Disabling	5. Unrelated (definitely not related)	
6. FATAL		
7. Unknown		

ID Number

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**PART VI: REPORTABLE TREATMENTS**

1. Answer each item YES NO N/A
- A. Transfusion **TRANSFUS** (1) (2) (3)
1. If yes, complete a. – d. Otherwise, skip to B.
- a. Transfusion Type: **TR\_TYPE**  
(1) Simple (2) Exchange
- b. Volume, answer b 1 or 2.
1. Whole Blood **TRVOLWBL** \_\_\_\_\_ \_\_\_\_\_ \_\_\_\_\_ cc
- OR
2. Packed Red Cells **TRVOLPR2** \_\_\_\_\_ \_\_\_\_\_ \_\_\_\_\_ cc
- c. Start Date: **TSTRT\_DT** \_\_\_\_\_ \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
Month Day Year
- d. Stop Date: **TSTOP\_DT** \_\_\_\_\_ \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
Month Day Year
- YES NO N/A
- B. Placement on chronic transfusion therapy **CHRTRAN** (1) (2) (3)
- C. Splenectomy **SPLCTMY** (1) (2) (3)
- D. Parenteral antibiotics **PAR\_ANTI** (1) (2) (3)
- E. Dialysis, limited course **DIALYS\_L** (1) (2) (3)

**PART VII: HOSPITALIZATION**

1. Hospital Name: \_\_\_\_\_ **HOSPNAME**
2. City: \_\_\_\_\_ **HOSPCITY**
3. State: \_\_\_\_\_ **HOSP\_ST**      4. Zip Code: \_\_\_\_\_ **HOSP\_ZIP**
5. Admission Date: **ADM\_DT** \_\_\_\_\_ \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
Month Day Year
6. Discharge Date: **DISCH\_DT** \_\_\_\_\_ \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
Month Day Year

ID Number

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**PART VIII: OUTCOMES**

		YES	NO
1. Significant new disability	<b>SNEWDISA</b>	(1)	(2)
2. Persistent new disability	<b>PNEWDISA</b>	(1)	(2)
3. Permanent new disability	<b>PERMDISA</b>	(1)	(2)
4. <b>DEATH</b>	<b>DEATH</b>	(1)	(2)

A.. Date of Death: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
**DEATH\_DT**      Month                  Day                  Year

B. Location:

**DTH\_LOC**

1. Inpatient (1)

2. In-Community (2)

**PART IX: COORDINATION**

1. Checked for completeness and accuracy:

A. Certification number: **CERT\_NO** \_\_\_\_\_ - \_\_\_\_\_

B. Signature: \_\_\_\_\_ **CERT\_SIG**

C. General Comments: **GEN\_CMNT**

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

ID Number

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**PART IV: COORDINATOR**

1. Checked for completeness and accuracy:

A. Certification number:        \_\_\_\_ - \_\_\_\_        **CERT\_NO**

B. Signature: \_\_\_\_\_ **CERT\_SIG**

C. General Comments: \_\_\_\_\_ **GEN\_CMNT**

\_\_\_\_\_

\_\_\_\_\_

ID Number				Visit			Seq		
							-		

BABY HUG FOLLOW-UP STUDY II

VINELAND SUMMARY

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ 2. Current Clinic: \_\_\_\_\_  
**SUBJECT\_ID** **SITE\_ID**
3. Patient's Letter Code: \_\_\_\_\_ **LETTER\_CD**
4. Testing Date:: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
**VISIT\_DT** Month Day Year

**PART II: CAREGIVER CODES**

1. Chronological Age: **CHRAGEYR** **CHRAGEMN** **CHRAGEDS**  
\_\_\_\_\_                        
Years Months Days
2. Caregiver's Relationship to Child: **CARE41**
- Mother (1)
  - Father (2)
  - Grandparent (3)
  - Aunt or Uncle (4)
  - Foster Parent (5)
  - Other (6)





BABY HUG FOLLOW-UP STUDY II

WISC-IV

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number:      \_\_\_\_\_      **SUBJECT\_ID**           2. Current Clinic:      \_\_\_\_\_      **SITE\_ID**
3. Patient's Letter Code:      \_\_\_\_\_      **LETTER\_CD**
4. Visit Date:      \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_      **VISIT\_DT**  
                                     Month      Day      Year

**PART II: TESTING RESULTS**

1. Full scale IQ:      \_\_\_\_\_      **FAIQ**
2. Verbal comprehension composite score:      \_\_\_\_\_      **VCI**
3. Perceptual reasoning composite score:      \_\_\_\_\_      **PRI**
4. Working memory composite score:      \_\_\_\_\_      **WMI**
5. Processing speed composite score:      \_\_\_\_\_      **PSI**
6. Block design scaled score:      \_\_\_\_\_      **BD**
7. Similarities scaled score:      \_\_\_\_\_      **SI**
8. Digit span scaled score:      \_\_\_\_\_      **DS**
9. Picture concepts scaled score:      \_\_\_\_\_      **PC**
10. Coding scaled score:      \_\_\_\_\_      **CD**
11. Vocabulary scaled score:      \_\_\_\_\_      **VC**
12. Letter-number sequence scaled score:      \_\_\_\_\_      **LN**
13. Matrix reasoning scaled score:      \_\_\_\_\_      **MR**
14. Comprehension scaled score:      \_\_\_\_\_      **CO**
15. Symbol search scaled score:      \_\_\_\_\_      **SS**

ID Number	Visit	-	Seq

**PART III: EXAMINER'S INFORMATION**

1. Name: \_\_\_\_\_ EXAMINER\_NM

**PART IV: COORDINATION**

1. Checked for completeness and accuracy:

- A. Certification number:      \_\_\_ \_\_\_ - \_\_\_ \_\_\_      CERT\_NO
  - B. Signature: \_\_\_\_\_      CERT\_SIG
  - C. General Comments: \_\_\_\_\_      GEN\_CMNT
- 

ID Number	Visit	-	Seq



















		Never	Almost Never	Sometimes	Often	Almost Always	
I.	About communication II (problems with..)						
	1. It is hard for me when others do not understand about my sickle cell disease	(0)	(1)	(2)	(3)	(4)	SCRPT_I1
	2. It is hard for me when others do not understand how much pain I feel	(0)	(1)	(2)	(3)	(4)	SCRPT_I2
	3. It is hard for me to tell others I have sickle cell disease	(0)	(1)	(2)	(3)	(4)	SCRPT_I3

2. **Parent** Report (Ages 8-12)

In the past one month, how much of a problem has your child had with ...

		Never	Almost Never	Sometimes	Often	Almost Always	
A.	Pain and hurt (problems with..)						
	1. Hurting a lot	(0)	(1)	(2)	(3)	(4)	SPRPT_A1
	2. Hurting all over his/her body	(0)	(1)	(2)	(3)	(4)	SPRPT_A2
	3. Hurting in his/her arms	(0)	(1)	(2)	(3)	(4)	SPRPT_A3
	4. Hurting in his/her legs	(0)	(1)	(2)	(3)	(4)	SPRPT_A4
	5. Hurting in his/her stomach	(0)	(1)	(2)	(3)	(4)	SPRPT_A5
	6. Hurting in his/her chest	(0)	(1)	(2)	(3)	(4)	SPRPT_A6
	7. Hurting in his/her back	(0)	(1)	(2)	(3)	(4)	SPRPT_A7
	8. Having pain every day	(0)	(1)	(2)	(3)	(4)	SPRPT_A8
	9. Having so much pain that he/she has to take medicine	(0)	(1)	(2)	(3)	(4)	SPRPT_A9

B.	Pain impact (problems with..)						
	1. It is hard for him/her to do things because he/she might get pain	(0)	(1)	(2)	(3)	(4)	SPRPT_B1
	2. Missing school when he/she has pain	(0)	(1)	(2)	(3)	(4)	SPRPT_B2
	3. It is hard for him/her to run when he/she has pain	(0)	(1)	(2)	(3)	(4)	SPRPT_B3
	4. It is hard for him/her to have fun when having pain	(0)	(1)	(2)	(3)	(4)	SPRPT_B4
	5. Having trouble moving around when he/she has pain	(0)	(1)	(2)	(3)	(4)	SPRPT_B5
	6. It is hard for him/her to stay standing when he/she has pain	(0)	(1)	(2)	(3)	(4)	SPRPT_B6
	7. It is hard for him/her to take care of himself/herself when he/she has pain	(0)	(1)	(2)	(3)	(4)	SPRPT_B7
	8. It is hard for him/her to do what others can do because he/she might get pain	(0)	(1)	(2)	(3)	(4)	SPRPT_B8
	9. Waking up at night when he/she has pain	(0)	(1)	(2)	(3)	(4)	SPRPT_B9
	10. Getting tired when he/she has pain	(0)	(1)	(2)	(3)	(4)	SPRPT_B10

ID Number

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Visit

--	--	--

Seq

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-



In the past one month, how much of a problem has your child had with ...

		Never	Almost Never	Some- times	Often	Almost Always		
H.	Communication I (problems with..)							
	1.	It is hard for him/her to tell others when he/she is in pain	(0)	(1)	(2)	(3)	(4)	SPRPT_H1
	2.	It is hard for him/her to tell the doctors and nurses how he/she feels	(0)	(1)	(2)	(3)	(4)	SPRPT_H2
	3.	It is hard for him/her to ask the doctors and nurses questions	(0)	(1)	(2)	(3)	(4)	SPRPT_H3
I.	Communication II (problems with..)							
	1.	It is hard for him/her when other people do not understand about his/her sickle cell disease	(0)	(1)	(2)	(3)	(4)	SPRPT_I1
	2.	It is hard for him/her when others do not understand how much pain he/she feels	(0)	(1)	(2)	(3)	(4)	SPRPT_I2
	3.	It is hard for him/her to tell others that he/she has sickle cell disease	(0)	(1)	(2)	(3)	(4)	SPRPT_I3

**PART VI: COORDINATION**

1. Checked for completeness and accuracy:

- A. Certification number: \_\_\_\_\_ - \_\_\_\_\_ CERT\_NO
  - B. Signature: \_\_\_\_\_ CERT\_SIG
  - C. General Comments: \_\_\_\_\_ GEN\_CMNT
- 

ID Number                      Visit                      Seq  

								-		
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BABY HUG FOLLOW-UP STUDY II

CONNERS CONTINUOUS PERFORMANCE TEST-II (CPT-II)

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ **SUBJECT\_ID**      2. Current Clinic: \_\_\_\_\_ **SITE\_ID**

3. Patient's Letter Code: \_\_\_\_\_ **LETTER\_CD**      4. Visit: \_\_\_\_\_

5. Visit Date:      \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ **VISIT\_DT**  
    Month                                   Day                                   Year

**PART II: CPT II RESULTS**

	Value	T-Score	Percentile	<u>Guideline</u>
1. Omissions	_____ . _____ <b>OMI_V</b>	_____ . _____ <b>OMI_T</b>	_____ . _____ <b>OMI_P</b>	( 1 ) ( 2 ) ( 3 ) <b>OMI_G</b>
2. Commissions	_____ . _____ <b>COMMI_V</b>	_____ . _____ <b>COMMI_T</b>	_____ . _____ <b>COMMI_P</b>	( 1 ) ( 2 ) ( 3 ) <b>COMMI_G</b>
3. Hit RT	_____ . _____ <b>RT_V</b>	_____ . _____ <b>RT_T</b>	_____ . _____ <b>RT_P</b>	( 1 ) ( 2 ) ( 3 ) <b>RT_G</b>
4. Hit RT Std Error	_____ . _____ <b>RTSE_V</b>	_____ . _____ <b>RTSE_T</b>	_____ . _____ <b>RTSE_P</b>	( 1 ) ( 2 ) ( 3 ) <b>RTSE_G</b>
5. Variability	_____ . _____ <b>VARI_V</b>	_____ . _____ <b>VARI_T</b>	_____ . _____ <b>VARI_P</b>	( 1 ) ( 2 ) ( 3 ) <b>VARI_G</b>
6. Detectability (d')	_____ . _____ <b>DETECT_V</b>	_____ . _____ <b>DETECT_T</b>	_____ . _____ <b>DETECT_P</b>	( 1 ) ( 2 ) ( 3 ) <b>DETECT_G</b>
7. Response Style	_____ . _____ <b>RESP_V</b>	_____ . _____ <b>RESP_T</b>	_____ . _____ <b>RESP_P</b>	( 1 ) ( 2 ) ( 3 ) <b>RESP_G</b>
8. Perservations	_____ . _____ <b>PERS_V</b>	_____ . _____ <b>PERS_T</b>	_____ . _____ <b>PERS_P</b>	( 1 ) ( 2 ) ( 3 ) <b>PERS_G</b>
9. Hit RT Block Change	_____ . _____ <b>RTBCHG_V</b>	_____ . _____ <b>RTBCHG_T</b>	_____ . _____ <b>RTBCHG_P</b>	( 1 ) ( 2 ) ( 3 ) <b>RTBCHG_G</b>
10. Hit SE Block Change	_____ . _____ <b>SEBCHG_V</b>	_____ . _____ <b>SEBCHG_T</b>	_____ . _____ <b>SEBCHG_P</b>	( 1 ) ( 2 ) ( 3 ) <b>SEBCHG_G</b>
11. Hit RT ISI Change	_____ . _____ <b>RTICHG_V</b>	_____ . _____ <b>RTICHG_T</b>	_____ . _____ <b>RTICHG_P</b>	( 1 ) ( 2 ) ( 3 ) <b>RTICHG_G</b>
12. Hit SE ISI Change	_____ . _____ <b>SEICHG_V</b>	_____ . _____ <b>SEICHG_T</b>	_____ . _____ <b>SEICHG_P</b>	( 1 ) ( 2 ) ( 3 ) <b>SEICHG_G</b>

**PART III: COORDINATION**

1. Checked for completeness and accuracy:

A. Certification number: \_\_\_\_\_ - \_\_\_\_\_ **CERT\_NO**

B. Signature: \_\_\_\_\_ **CERT\_SIG**

C. General Comments: \_\_\_\_\_ **GEN\_CMNT**

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BABY HUG FOLLOW-UP STUDY II  
PULMONARY FUNCTION TESTING

Do hemoglobin and pulse oximetry along with PFT

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ 2. Current Clinic: \_\_\_\_\_  
SUBJECT\_ID SITE\_ID
3. Patient's Letter Code: \_\_\_\_\_ LETTER\_CD
4. Testing Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
VISIT\_DT Month Day Year

**PART II: DEMOGRAPHIC INFORMATION**

1. Height: \_\_\_\_\_ . \_\_\_\_\_ (1) inches HT\_UNIT  
(2) centimeters
- A. Height is measured by:
- Standing height (1)  
Arm span (2) HT\_METHOD
2. Weight: \_\_\_\_\_ (1) pounds WT\_UNIT  
(2) kilograms
3. With which primary race or ethnicity does the patient identify? PFTRACE  
**(Check only one)**
- White (Caucasian) (1)  
Hispanic (2)  
African-American (3)  
Asian or Pacific Islander (4)  
Other or none of the above (5)  
*Unknown / undetermined* (6)
4. Does the patient identify with more than one race or ethnicity? MORETHAN\_ONERACE  
Yes (1)  
No (2)

**PART III: SPIROMETRY**

1. Date of spirometry: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ SPIROMETRY\_DT  
Not done  
( 1 )  
Month Day Year

***If spirometry 'Not done', skip to Part V.***

**NOTE: The PFT tech should try to obtain an exhalation effort of  $\geq$  6 seconds**

2. Pre-bronchodilator spirometry: \_\_\_\_\_ PRE\_BRONCH\_SPIROM\_ND  
Not done  
( 1 )

***If pre-bronchodilator spirometry 'Not done', skip to 3.***

A. Pre-bronchodilator Results:

0. Was the participant's effort acceptable and reproducible according to ATS guidelines?

Yes ( 1 ) PRE\_EFFORT  
 No ( 2 ) \*  
 Questionable ( 3 ) \*

\*a. If no or questionable, why was effort unacceptable, unreproducible, or questionable?

\_\_\_\_\_ PRE\_EFFORT\_SP

\_\_\_\_\_

				Not done		
1.	FEV <sub>1</sub>	PREFEV1	_____ . _____	<i>L (largest)</i>	( 1 )	PREFEV1_ND
2.	FVC	PREFVC	_____ . _____	<i>L (largest)</i>	( 1 )	PREFVC_ND
3.	PEFR (FEF <sub>max</sub> )	PREPEFR	_____ . _____	<i>L/second (largest)</i>	( 1 )	PREPEFR_ND
4.	FEF 25-75	PREFEF25	_____ . _____	<i>L/second (from largest FEV<sub>1</sub>+FVC)</i>	( 1 )	PREFEF25_ND
5.	Ratio (FEV <sub>1</sub> / FVC)	PRE_RATIO	_____ . _____		( 1 )	PRE_RATIO_ND

ID Number
Visit
Seq  
[ ] [ ] [ ] [ ]
[ ] [ ] [ ] [ ]
-
[ ] [ ] [ ] [ ]



**PART IV: LUNG VOLUME**

**Record actual Pre-Bronchodilator Measurements**

1. Lung volume: **LUNG\_VOL2**  
 Not done (*skip to Part V*) (1)  
 Performed in conformance with BHFUII requirements (i.e., meets ATS guidelines for acceptability) (2)  
 Technique was acceptable with good effort (3)  
 Technique was acceptable with questionable effort (4)  
 Results not interpretable (5)
- A. Was the participant able to perform 3 acceptable maneuvers? **LUNG\_3MANEU**  
 Yes (1)  
 No (2)
2. Date lung volume performed: **LUNG\_VOL\_DT**  
 \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
Month Day Year
3. Technique: **LUNGV\_TECH**  
 Plethysmography (preferred) Helium dilution Nitrogen washout  
(1) (2) (3)  
Not done
4. TLC \_\_\_\_\_ . \_\_\_\_\_ *L (mean FRC+MAX IC)* **TLC** (1) **TLC\_ND**
5. Maximum SVC \_\_\_\_\_ . \_\_\_\_\_ *L* **MAX\_SVC** (1) **MAX\_SVC\_ND**
6. RV \_\_\_\_\_ . \_\_\_\_\_ *L (TLC-highest VC)* **RV** (1) **RV\_ND**
7. Mean FRC (TGV) \_\_\_\_\_ . \_\_\_\_\_ *L (mean from 3 maneuvers)* **MEAN\_FRC** (1) **MEAN\_FRC\_ND**

**PART V: DIFFUSING CAPACITY**

1. DLCO: **DLCO**  
 Not done (*skip to Part VI*) (1)  
 Performed in conformance with BHFUII requirements (2)  
 Not in conformance with BHFUII requirements, but results are clinically interpretable (3)  
 Results not interpretable (4)
- A. Was the participant's effort acceptable and reproducible according to ATS guidelines?  
 Yes (1) **DLCO\_EFFORT**  
 No (2)\*  
 Questionable (3)\*

\*1. If no or questionable, why was effort unacceptable, unreproducible, or questionable?

**DLCO\_EFFORT\_SP**

---

ID Number	Visit	-	Seq

2. Date D<sub>L</sub>CO performed: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
 Month Day Year
- Not  
done
3. Mean D<sub>L</sub>CO  
 (uncorrected for hemoglobin) \_\_\_\_\_ . \_\_\_\_\_ *mL/min/mmHg* ( 1 ) MEAN\_DLCO\_ND
4. Hemoglobin \_\_\_\_\_ . \_\_\_\_\_ *g/dL* HEMOGLOBIN ( 1 ) HEMOGLOBIN\_ND
5. Alveolar Volume \_\_\_\_\_ . \_\_\_\_\_ *L (largest)* VA ( 1 ) VA\_ND

**PART VI: PULSE OXIMETRY**

1. Oxygen saturation (room air): \_\_\_\_\_ % O<sub>2</sub>SAT ( 1 ) O<sub>2</sub>S\_ND
- Not  
done

**PART VII: COORDINATION**

1. Checked for completeness and accuracy:
- A. Certification number: \_\_\_\_\_ - \_\_\_\_\_ CERT\_NO
- B. Signature: \_\_\_\_\_ CERT\_SIG
- C. General Comments: \_\_\_\_\_  
 \_\_\_\_\_  
 \_\_\_\_\_

ID Number	Visit	-	Seq

BABY HUG FOLLOW-UP STUDY II  
 ECHOCARDIOGRAM PERFORMANCE

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ 2. Current Clinic: \_\_\_\_\_  
SUBJECT\_ID SITE\_ID
3. Patient's Letter Code: \_\_\_\_\_ LETTER\_CD
4. Testing Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
VISIT\_DT Month Day Year

**PART II: GENERAL INFORMATION**

**INSTRUCTIONS**

The following information **MUST BE** collected on the day echocardiogram is completed, if an echocardiogram was ever performed for this visit.

1. Date of echocardiogram visit: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
ECHO Month Day Year
2. Child's date of birth: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
BIRTH\_DT Month Day Year
3. Source indication: SOURCE
- Routine BABY HUG FU II visit (1)  
 Abstract from non-BHFU II visit (2)
4. Patient state: PT\_STATE
- Relaxed (1)  
 Tense (2)  
 Unmanageable (3)  
 N/A (4)
5. Label Number: \_\_\_\_\_ LABEL

ID Number	Visit	Seq										
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6. Height: \_\_\_\_\_ . \_\_\_\_\_ cm **HEIGHT**

7. Weight: \_\_\_\_\_ . \_\_\_\_\_ kg **WEIGHT**

8. a. Temperature

i. **TEMPF** \_\_\_\_\_ . \_\_\_\_\_ °F **OR** ii. **TEMPC** \_\_\_\_\_ . \_\_\_\_\_ °C

b. Thermometer placement:

- |          |                    |
|----------|--------------------|
|          | <b>THERM_PLACE</b> |
| Axillary | (1)                |
| Oral     | (2)                |
| Rectal   | (3)                |
| Tympanic | (4)                |
| N/A      | (5)                |

9. Heart rate: \_\_\_\_\_ beat/min **HEARTRATE**

10. Respiratory rate: \_\_\_\_\_ breath/min **RESP**

11. Blood Pressure:

a. Systolic: \_\_\_\_\_ mm Hg **BP\_SYSTOLIC**

b. Diastolic: \_\_\_\_\_ mm Hg **BP\_DIASTOLIC**

c. Method: **BP\_METHOD**

- |              |     |
|--------------|-----|
| Dinamap      | (1) |
| Doppler      | (2) |
| Auscultation | (3) |
| Palpation    | (4) |
| N/A          | (5) |

ID Number	Visit	Seq									
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**PART III: ECHOCARDIOGRAM STATUS**

- |  |     |     |  |
|--|-----|-----|--|
|  | Yes | No  |  |
|  | (1) | (2) |  |
1. Was the echocardiogram completed? ECHO\_COMP
  
  2. Reason(s) that the echocardiogram not completed: **(CHECK ALL THAT APPLY)**
    - a. Equipment availability (1) EQUIP\_AVAIL
    - b. Caretaker objects to participation (1) CARETAKER\_OBJ
    - c. Illness of subject, test cancelled by physician (1) ILLNESS\_SUB
    - d. Subject developed complications during procedure (1) COMPLIC-DURING
    - e. Other (specify) (1) INCOM\_OTHER
      - i. Specify: INCOMP\_RSN\_SP  
 \_\_\_\_\_
    - f. Unknown (1) INCOM\_UNK

**IF NOT COMPLETED, GO TO SECTION V**

- |  |      |     |
|--|------|-----|
|  | Yes  | No  |
|  | (1)* | (2) |
3. Were there any complications from the echocardiogram? COMPLICATIONS
    - a. What were the complications?  
 \*(Specify): COMPLICATION\_SP  
 \_\_\_\_\_  
 \_\_\_\_\_

4. Blood Pressure at the time closest to M-mode:

	i. First Reading	ii. Second Reading	iii. Third Reading
a. Systolic blood pressure	BP_SYSTOLIC_1ST ____ mm Hg	BP_SYSTOLIC_2ND ____ mm Hg	BP_SYSTOLIC_3RD ____ mm Hg
b. Diastolic blood pressure	BP_DIASTOLIC_1ST ____ mm Hg	BP_DIASTOLIC_2ND ____ mm Hg	BP_DIASTOLIC_3RD ____ mm Hg
c. Mean blood pressure	BP_MEAN_1ST ____ mm Hg	BP_MEAN_2ND ____ mm Hg	BP_MEAN_3RD ____ mm Hg

ID Number	Visit	Seq







BABY HUG FOLLOW-UP STUDY II

**MRI/MRA PERFORMANCE**

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ 2. Current Clinic: \_\_\_\_\_  
**SUBJECT\_ID** **SITE\_ID**
3. Patient's Letter Code: \_\_\_\_\_ **LETTER\_CD**
4. Visit Date:: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
**VISIT\_DT** Month Day Year

**PART II: EQUIPMENT AND QUALITY**

1. Equipment: \_\_\_\_\_ **MRIMRA\_EQPT**

2. MRI Film Label **MRI\_LBL**

3. MRA Film Label **MRA\_LBL**

4. Scan Quality **MRIMRA\_QUALITY**
- |                                    |     |
|------------------------------------|-----|
| Excellent                          | (1) |
| Slight Artifact/Motion, Adequate   | (2) |
| Severe Artifact/Motion, Inadequate | (3) |

**PART III: TECHNICIAN INFORMATION**

1. Technician Name: \_\_\_\_\_ **TECH\_NM**

2. Signature: \_\_\_\_\_ **SIGNATURE**

ID Number				Visit			-	Seq	
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	-	<input type="text"/>	<input type="text"/>

**PART IV: COORDINATION**

1. Checked for completeness and accuracy:

A. Certification number: \_\_\_\_\_ - \_\_\_\_\_ CERT\_NO

B. Signature: \_\_\_\_\_ CERT\_SIG

C. General Comments: GEN\_CMNT

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

ID Number				Visit			Seq			
								-		

BABY HUG FOLLOW-UP STUDY  
LIVER-SPLEEN CENTRAL READING

**PART I: IDENTIFYING INFORMATION**

1. Film Label BH2 \_\_\_\_\_ SPEC\_ID
2. Date read: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ VISIT\_DT  
Month Day Year
3. Visit: Y 1 0 VISIT\_NBR

**PART II: LIVER-SPLEEN SCAN QUALITY**

1. Reader's Last Name: LSRDRNM \_\_\_\_\_
2. Reader Signature: LSRDRSIG \_\_\_\_\_
3. Reader Number: \_\_\_\_\_ LSRDRNBR
4. Current Status of this Reading: LSSCN\_QLTY

- Quality adequate and reading complete (1)  
Quality inadequate for reading (2)\*

\*A. If inadequate, explain: QLTY\_SP \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

If Item 4 is 2, skip to Part IV.

Film Label 

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BABY HUG FOLLOW-UP STUDY II

**ABDOMINAL SONOGRAM (ULTRASOUND) CENTRAL READING**

**PART I: IDENTIFYING INFORMATION**

1. Film Label BH2 \_\_\_\_\_ **SPEC\_ID**
2. Date read: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ **VISIT\_DT**  
Month Day Year
3. Visit: Y 1 0 **VISIT\_NBR**

**PART II: EQUIPMENT**

1. Reader's Last Name: \_\_\_\_\_ **ABDRDRNM**
2. Reader Signature: **ABDRDRSIG** \_\_\_\_\_
3. Reader Number: \_\_\_\_\_ **ABDRDRNBR**
4. Current Status of this Reading: **ABD\_QLTY**
- |   |        |
|---|--------|
| Quality adequate and reading complete                     | (1 )   |
| Returned for reprocessing                                 | (2 )*  |
| Quality inadequate for reading after reprocessing (final) | (3 )** |
- \*A. If returned for reprocessing, explain: \_\_\_\_\_ **QLTY1\_SP**
- \_\_\_\_\_
- \*\*B. If inadequate, explain: \_\_\_\_\_ **QLTY3\_SP**
- \_\_\_\_\_

If 2 or 3, Skip to Part IV.

Film Label 

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**PART III: RESULTS**

	Present (1)	Absent (2)	N/A (3)	<b>GALBLA</b>
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If Absent or N/A , Skip to Item 2.

A. If Present				<b>GBWALL</b>
Normal thin wall	(1)			
Thick walled or edema	(2)			
Not able to assess	(3)			

**GBCDV**

B. Color Doppler Vascularity	Minimal (1)	Moderate (2)	Marked (3)	N/D (4)
------------------------------	----------------	-----------------	---------------	------------

C. If gallbladder present, answer C1 or C2:

1. Number of Stones			<b>GBNSTN</b>
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If no gallbladder stones, record 00 in C1 and N/A in D and E.

OR

2. Multiple stones not countable	Yes (1)	<b>GBMSTN</b>
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D. Largest stone			mm	N/A (1)	<b>GBLGSTNA</b>
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E. Stones Freely Mobile?	Yes (1)	No (2)	N/A (3)	<b>GBSFM</b>
--------------------------	------------	-----------	------------	--------------

F. Dilation	Dilated	Normal	N/A	
1. Common bile duct	(1)	(2)	(3)	<b>GBCBD</b>
2. Pancreatic duct	(1)	(2)	(3)	<b>GBPAND</b>
3. Intrahepatic ducts	(1)	(2)	(3)	<b>GBIHEP</b>

G. Sludge	Present (1)	Absent (2)	N/A (3)	<b>GBSLDG</b>
H. Pericholecystic fluid	(1)	(2)	(3)	<b>GBPRFL</b>

Film Label

2. Spleen	Present (1)	Absent (2)	N/A (3)	<b>SPLEEN</b>
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If Absent or N/A, Skip to Item 3.

A. Accessory spleen(s)	(1)	(2)	(3)	<b>ACCSPL</b>
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B. Cephalocaudad length	<input style="width: 30px; height: 20px;" type="text"/> <input style="width: 30px; height: 20px;" type="text"/> . <input style="width: 30px; height: 20px;" type="text"/> cm	<b>SPLCLN</b>
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C. Transverse	<input style="width: 30px; height: 20px;" type="text"/> <input style="width: 30px; height: 20px;" type="text"/> . <input style="width: 30px; height: 20px;" type="text"/> cm	<b>SPLTRN</b>
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D. Anterior – Posterior	<input style="width: 30px; height: 20px;" type="text"/> <input style="width: 30px; height: 20px;" type="text"/> . <input style="width: 30px; height: 20px;" type="text"/> cm	<b>SPLANP</b>
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E. Estimated total spleen volume	<input style="width: 30px; height: 20px;" type="text"/> <input style="width: 30px; height: 20px;" type="text"/> <input style="width: 30px; height: 20px;" type="text"/> cu cm	<b>SPLVOL</b>	(1) N/D <b>SPLVOLND</b>
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F. Homogeneity	<b>SPLHOM</b>
Homogeneous	(1)
Inhomogeneous	(2)*
N/A	(3)

\*1. If inhomogeneous, explain: **INHOM\_SP**

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3. Right Kidney	Present (1)	Absent (2)	N/A (3)	<b>RKID</b>
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If Absent or N/A, Skip to Item 4.

A. Estimated volume	<input style="width: 30px; height: 20px;" type="text"/> <input style="width: 30px; height: 20px;" type="text"/> <input style="width: 30px; height: 20px;" type="text"/> cu cm	<b>RKVOL</b>
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B. Renal parenchyma	Normal (1)	Abnormal (2)*	N/A (3)	<b>RKRPAR</b>
---------------------	---------------	------------------	------------	---------------

\*1. If abnormal, explain: **RKRPEX**

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C. Echogenicity	(1)	(2)*	(3)	<b>RKECHO</b>
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\*1. If abnormal, explain: **RKECEX**

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Film Label

4. Left Kidney	Present ( 1 )	Absent ( 2 )	N/A ( 3 )	<b>LKID</b>
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If Absent or N/A, Skip to Item 5.

A. Estimated volume	<table border="1" style="display: inline-table; border-collapse: collapse;"> <tr> <td style="width: 20px; height: 20px;"></td> <td style="width: 20px; height: 20px;"></td> <td style="width: 20px; height: 20px;"></td> </tr> </table>				cu cm	<b>LKVOL</b>

B. Renal parenchyma	Normal ( 1 )	Abnormal ( 2 )*	N/A ( 3 )	<b>LKRPAR</b>
---------------------	-----------------	--------------------	--------------	---------------

\*1. If abnormal, explain: \_\_\_\_\_ **LKRPEX**

C. Echogenicity	( 1 )	( 2 )*	( 3 )	<b>LKECHO</b>
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\*1. If abnormal, explain: \_\_\_\_\_ **LKECEX**

5. Liver enlarged	Yes ( 1 )	No ( 2 )	N/A ( 3 )	<b>LVRENL</b>
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6. Any other abnormalities	( 1 )*	( 2 )	( 3 )	<b>ABOABN</b>
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\*A. If yes, explain: \_\_\_\_\_ **ABOABNEX**

**PART IV: COORDINATION**

1. Checked for completeness and accuracy:

A. Certification number:	<table border="1" style="display: inline-table; border-collapse: collapse;"> <tr> <td style="width: 20px; height: 20px;"></td> <td style="width: 20px; height: 20px;"></td> </tr> </table> - <table border="1" style="display: inline-table; border-collapse: collapse;"> <tr> <td style="width: 20px; height: 20px;"></td> <td style="width: 20px; height: 20px;"></td> </tr> </table>					<b>CERT_NO</b>

B. Signature:	_____	<b>CERT_SIG</b>
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C. General Comments:	_____	<b>GEN_CMNT</b>
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Film Label 

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BABY HUG FOLLOW-UP STUDY II

**MRA READING**

**PART I: IDENTIFYING INFORMATION**

- 1. Film Label BH2 \_\_\_\_\_ **SPEC\_ID**
  
- 2. Date read: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ **VISIT\_DT**  
                                Month                        Day                        Year
  
- 3. Visit: Y 1 0 **VISIT\_NBR**

**PART II: TO BE COMPLETED BY READER**

- 1. Reader [Last Name]: \_\_\_\_\_ **MRARDRNM**
  
- 2. Reader # \_\_\_\_\_ **MRARDRNO**
  
- 3. SCAN QUALITY (MARK ONE):
  - Excellent (1) **MRA\_QUAL**
  - Slight Artifact/Motion, Adequate (2)
  - Severe Artifact/Motion, Inadequate (3)
  
- 4. CURRENT STATUS OF THIS READING
  - Adequate scan, reading complete (1) **MRA\_STAT**
  - Returned for reprocessing (2)
  - Inadequate, reading not completed (3)

**If (2) or (3), SKIP to Part IV.**

Film Label 

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**PART III: CENTRAL REVIEW INTERPRETATION** (Answer items 1-8 using the codes below.)

a. OVERALL RATING		b. DESCRIPTION OF ABNORMALITY		d. INVOLVED SEGMENTS	
1	= Normal	1	= Stenosis, Mild (25% to 50% narrowing)	1	= Supraclinoid
2	= Equivocal	2	= Stenosis, Moderate (50% to 75% narrowing)	2	= Pre- or Juxtaseilar
3	= Abnormal	3	= Stenosis, Severe (75% to 99% narrowing)	3	= Petrous
		4	= Occlusion	4	= Distal cervical

	OVERALL RATING	DESCRIPTION OF ABNORMALITY (IF OVERALL RATING =3)	LENGTH OF STENOTIC SEGMENT (mm)	INVOLVED SEGMENT (INDICATE ALL THAT APPLY IF RATING =3)			
1. Right ICA	a _____ <b>ORRICA</b>	b _____ <b>ABRICA</b>	c _____ <b>LSSRICA</b>	d1 _____ <b>INVSEGR1</b>	d2 _____ <b>INVSEGR2</b>	d3 _____ <b>INVSEGR3</b>	d4 _____ <b>INVSEGR4</b>
2. Right MCA	a _____ <b>ORRMCA</b>	b _____ <b>ABRMCA</b>	c _____ <b>LSSRMCA</b>				
3. Right ACA	a _____ <b>ORRACA</b>	b _____ <b>ABRACA</b>	c _____ <b>LSSRACA</b>				
4. Left ICA	a _____ <b>ORLICA</b>	b _____ <b>ABLICA</b>	c _____ <b>LSSLICA</b>	d1 _____ <b>INVSEGL1</b>	d2 _____ <b>INVSEGL2</b>	d3 _____ <b>INVSEGL3</b>	d4 _____ <b>INVSEGL4</b>
5. Left MCA	a _____ <b>ORLMCA</b>	b _____ <b>ABLMCA</b>	c _____ <b>LSSLMCA</b>				
6. Left ACA	a _____ <b>ORLACA</b>	b _____ <b>ABLACA</b>	c _____ <b>LSSLACA</b>				
7. Basilar	a _____ <b>ORBASIL</b>	b _____ <b>ABBASIL</b>	c _____ <b>LSSBASIL</b>				
8. Overall MRA	a _____ <b>ORMRA</b>	b _____ <b>ABMRA</b>	c _____ <b>LSSMRA</b>	d1 _____ <b>INVSEG1</b>	d2 _____ <b>INVSEG2</b>	d3 _____ <b>INVSEG3</b>	d4 _____ <b>INVSEG4</b>
9. Collateral Blood Vessels (Mark One):			<b>BLDVLSL</b>	Right (1)	Left (2)	Both (3)	Not Present (4)

Film Label 

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**PART IV: COORDINATION**

1. To be Completed by Radiology Technician:

- A. Certification number:      \_\_\_ \_\_\_ - \_\_\_ \_\_\_      **CERT\_NO**
  - B. Signature: \_\_\_\_\_      **CERT\_SIG**
  - C. General Comments:      **GEN\_CMNT** \_\_\_\_\_
- 

Film Label 

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**BABY HUG FOLLOW-UP STUDY II  
 MAJOR EVENT**

**PART I: IDENTIFYING INFORMATION**

1. Patient's ID Number: \_\_\_\_\_ 2. Current Clinic: \_\_\_\_\_  
SUBJECT\_ID SITE\_ID
3. Patient's Letter Code: \_\_\_\_\_ LETTER\_CD
4. Reporting Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
VISIT\_DT Month Day Year

**PART II: EVENT PERIOD**

1. Date of Event
- A. Event Start Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ START\_DT  
Month Day Year
- B. Event End Date: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_ E\_END\_DT  
Month Day Year

**PART III: MAJOR EVENT**

- | 1. Please indicate all diagnoses:   | YES | NO  |
|---|-----|-----|
| A. Acute Chest Syndrome <span style="float: right;">HX_ACS</span>                   | (1) | (2) |
| B. Splenic Sequestration Crisis <span style="float: right;">HXSPLSEQ</span>         | (1) | (2) |
| C. Initial or prolonged hospitalization <span style="float: right;">LONGHOSP</span> | (1) | (2) |
| D. Stroke or TIA <span style="float: right;">HX_STROKE_TIA</span>                   | (1) | (2) |
| E. Emergency Room Visit   | (1) | (2) |
| F. Life Threatening <span style="float: right;">LIFE_THREAT_EVT</span>              | (1) | (2) |
| G. Disability or Permanent Damage   | (1) | (2) |
| H. Death <span style="float: right;">HX_DEATH</span>                                | (1) | (2) |
| I. ICU Admission <span style="float: right;">ICU</span>                             | (1) | (2) |
| J. Pain crisis  | (1) | (2) |
| K. Other  | (1) | (2) |
| 1. Specify: _____   |     |     |

**PART IV: ADDITIONAL DIAGNOSIS INFORMATION**

If PART III, Item 1A is YES, answer 1. Otherwise, skip to 2.

- |    |  |         |        |          |     |          |
|----|--|---------|--------|----------|-----|----------|
| 1. | Acute Chest Syndrome                                       | None    | 1 Lobe | >1 Lobe  | N/A |          |
|    | A. New Infiltrate  | (1)     | (2)    | (3)      | (4) | ACSNINF  |
|    | B. O <sub>2</sub> % Saturation on Room Air at Presentation | _____   | _____  | . _____% |     | ACSSRAP  |
|    | C. Oxygen Administered                                     | _____   | _____  | . _____L |     | ACSOXADM |
|    | D. Mechanical Ventilation                                  | Yes (1) |        | No (2)   |     | ACSMVENT |

If PART III, Item 1B is YES, answer 2. Otherwise, skip to 3.

2. Splenic Sequestration
- |    |  |                 |        |        |        |            |          |
|----|--|-----------------|--------|--------|--------|------------|----------|
| A. | Spleen size below LCM <b>prior</b> to Major Event  | SPLNSIZE_PRIOR  |        |        |        |            |          |
|    |  | <2 cm           | 2-4 cm | 4-6 cm | 6-8 cm | >8 cm      |          |
|    |  | (1)             | (2)    | (3)    | (4)    | (5)        |          |
|    | B. Spleen size below LCM <b>during</b> Major Event | SPLNSIZE_DURING |        |        |        |            |          |
|    |  | <2 cm           | 2-4 cm | 4-6 cm | 6-8 cm | >8 cm      |          |
|    |  | (1)             | (2)    | (3)    | (4)    | (5)        |          |
|    | C. Nadir hemoglobin                                | _____ . _____   |        |        |        | gm/dL      | SPLNHMGL |
|    | D. Platelet count at time of nadir hemoglobin      | _____           |        |        |        | k/ $\mu$ L | SPLPTCNT |

If PART III, Item 1C is YES, answer 3. Otherwise, skip to 4.

3. Prolonged Hospitalization
- A. Reason: LONGHOSP\_SP
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If PART III, Item 1D is YES, answer 4-5. Otherwise, skip to Part V.

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4.	(Stroke or TIA) Findings of		YES	NO	N/A
A.	Loss of consciousness	LOS_CONS	(1)	(2)	(3)
B.	Change in mental status	CHG_MENT	(1)	(2)	(3)
C.	Loss of or difficulty with speech or vocalization	SPEECH	(1)	(2)	(3)
D.	Paralysis or weakness	PARALYS	(1)	(2)	(3)
E.	Difficulty with swallowing	DIFFSWAL	(1)	(2)	(3)
F.	Difficulty with vision	DIFF_SEE	(1)	(2)	(3)
G.	Loss of balance or dizziness	BALANCE	(1)	(2)	(3)
H.	Seizures	SEIZURE	(1)	(2)	(3)
I.	Headache	HEADACHE	(1)	(2)	(3)
5.	Results of Imaging Tests		Normal	Abnormal	Not Done
A.	MRI of brain	F50MRI	(1)	(2)	(3)
B.	CT scan of brain	F50CTBR	(1)	(2)	(3)
C.	PET scan of brain	F50PTBR	(1)	(2)	(3)
D.	MRA cerebral vasculature	F50MRA	(1)	(2)	(3)
E.	Transcranial Doppler	F50TCD	(1)	(2)	(3)
F.	Arteriogram	F50ARTGR	(1)	(2)	(3)
G.	Chest x-ray		(1)	(2)	(3)

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**PART V: DIAGNOSIS/PROBLEM SEVERITY AND ATTRIBUTION**

Complete PART V for each item in PART III checked YES.

1. Diagnosis/ Problem	2. Date of Onset	3. Number of Days	4. <sup>1</sup> Severity	5. <sup>2</sup> Attribution to Study Treatment	6. <sup>3</sup> Diagnosis Unexpected

<u><sup>1</sup>Severity</u>	<u><sup>2</sup>Attribution to Study Test</u>	<u><sup>3</sup>Diagnosis Unexpected</u>
1. Mild	1. Definite (clearly related)	1. Yes
2. Moderate	2. Probably (likely related)	2. No
3. Severe	3. Possible (may be related)	3. N/A
4. Life threatening	4. Unlikely (doubtfully related)	
5. Disabling	5. Unrelated (definitely not related)	
6. FATAL		
7. Unknown		

**PART VI: REPORTABLE TREATMENTS**

1. Answer each item YES      NO      N/A
- A. Transfusion **TRANSFUS** (1) (2) (3)
1. If yes, complete a. – d. Otherwise, skip to B.
- a. Transfusion Type: (1) Simple **TR\_TYPE** (2) Exchange
- b. Volume, answer b 1 or 2.
1. Whole Blood **TRVOLWBL** \_\_\_\_\_ cc
- OR
2. Packed Red Cells **TRVOLPR2** \_\_\_\_\_ cc

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c. Start Date: TSTRT\_DT \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
Month Day Year

d. Stop Date: TSTOP\_DT \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
Month Day Year

		YES	NO	N/A
B. Placement on chronic transfusion therapy	<span style="color:red">CHRTRAN</span>	(1)	(2)	(3)
C. Splenectomy	<span style="color:red">SPLCTMY</span>	(1)	(2)	(3)
D. Parenteral antibiotics	<span style="color:red">PAR_ANTI</span>	(1)	(2)	(3)
E. Dialysis, limited course	<span style="color:red">DIALYS_L</span>	(1)	(2)	(3)
F. Antibiotics		(1)	(2)	(3)
G. Pain medicine		(1)	(2)	(3)

**PART VII: HOSPITALIZATION**

1. Admission Date: ADM\_DT \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
Month Day Year

2. Discharge Date: DISCH\_DT \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_  
Month Day Year

**PART VIII: OUTCOMES**

	YES	NO
1. Resolved	(1)	(2)
2. Ongoing	(1)	(2)

ID Number

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**PART IX: COORDINATION**

1. Checked for completeness and accuracy:

A. Certification number: CERT\_NO              -          

B. Signature: \_\_\_\_\_ CERT\_SIG

C. General Comments: GEN\_CMNT

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

**Please Fax the hospital narrative along with this form to the BABY HUG FUP II Data Coordinating Center (DCC) at 443-524-2320.**

ID Number

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