

Cord For Life® Cord Blood Donation Information



Date Issued: 29AUG2023

Donating your child's cord blood is truly a *Life Saving* endeavor. To ensure the success of the donation, please read this information carefully. If you have any questions, please call our toll-free number (800) 869-8608. For additional information about the cord blood donation program or if you wish to proceed with donating your child's cord blood, visit our website www.cordforlife.com. Registration forms are available on the web or you may complete the attached forms and forward to the address listed at the bottom of this page.

IMPORTANT INFORMATION BEFORE YOU PROCEED

- You must be less than 34th week gestation to begin the application process
- You must be 18 years of age or older in order to donate your baby's cord blood
- We cannot accept cord blood donations from:
 - Multiple births (Acceptable for Private Storage-ONLY)
 - If you and your physician have chosen to delay the clamping of the cord after your child's birth (Acceptable for Private Storage-ONLY)
- Because Cord For Life is a privately funded company, we are limited to the amount of donations we can accept monthly
- Due to circumstances outside of Cord For Life control we are unable to accept cord blood units for <u>public donation</u> that were collected between Friday 3:00PM (EST) through Sunday 3:00PM (EST)

Speak with your doctor or midwife about donating the cord blood. Find out if the doctor is willing to perform this collection and if there will be any collection fees involved. Cord For Life does not charge collection fees, but some doctors may charge for this service. We suggest that you do not donate if you are going to be charged a fee. Cord For Life <u>cannot</u> reimburse you or your doctor/midwife for cord blood collection fees.

ABOUT THE ATTACHED FORMS:

The intent of the attached documents is to ensure the following:

- 1. You and your physician have a thorough understanding of the cord blood donation program.
- 2. The cord blood is collected properly and safely.
- 3. The quality of the cord blood product collected.

Please read all the documents thoroughly and complete all the requested documentation. <u>Blanks or omissions can result in the unit not meeting regulatory standards (FDA, AABB, NMDP) preventing Cord For Life from making the unit available for a transplant recipient.</u> Use only <u>BLACK or BLUE</u> ink when completing the forms. It is important that you use only your <u>legal name</u>* consistently on each document. Below is a description of the attached forms:

• Informed Consent for Cord Blood Medical Research, Form B.1-7

This form explains the National Marrow Donor Program (NMDP) protocol. Cord Blood stem cell transplantation is being investigated/studied by the NMDP, as such, it is important that you "the donor" understand the details of this program and how information regarding your donation will be utilized in this investigation.

• Cord For Life Donor Information and Health Questionnaire, Form B.1-1

Donor Information: The donor information section provides demographic data for yourself, the baby's father and your physician. In addition, it contains a "Baby's Ethnicity" section. This information is used to assist in the NMDP investigation and will not be used, in any manner, to prevent your unit from being placed on the registry. **IMPORTANT:** Your signature at the bottom of this page is <u>mandatory</u>. Remember to sign using only your legal name* as written at the top of the form.

Health Questionnaire: A thorough health history is an important part of ensuring the safety and quality of the cord blood. These questions may seem overwhelming, but they are a necessity. Any potential risk to a transplant recipient must be investigated. If you are unsure, contact a Cord For Life representative, they will assist you with any questions or concerns. Many of the items are not a reason for deferral but do require further information or clarification before acceptability is assigned. IMPORTANT: If you consider yourself to be at risk as described in the donation information, please do not donate. The standards and regulations set forth for all blood banks prevent us from accepting or using any cord blood from an at-risk donor.

• <u>Donation Informed Consent, Release-Hospital Birthing Center and The Inform Consent for HIV Test, Form B.1-9</u>

These are agreements between yourself, your physician and Cord For Life for the cord blood donation and information about HIV and consent for HIV testing. The physician must sign where indicated to allow Cord For Life to perform the required disease testing. Even if you have had previous prenatal testing done, we are required to test your blood for infectious diseases from a sample collected within 48 hours of your baby's birth. Tubes for the collection of your blood are included in the kit, and your blood will be drawn around the time of delivery by the hospital or birth center staff.

IMPORTANT: The results of your blood tests are confidential. You and your physician /midwife will be notified of any abnormal results via certified mail. It may be necessary to report certain positive test results to your local health department.

• Physician/Midwife Training Material Request for CB Collection, Form C.1-4

To ensure the cord blood is collected in a manner that results in the best possible product, Cord For Life requires the physician/midwife read and complete this form. Cord For Life offers training material options, at no cost, to the physician/midwife on request.

We thoroughly review all of your documentation on receipt. You will be contacted, by email, to review the information and obtain any additional information, if needed. If accepted as a donor, we will issue you a cord blood collection kit, which includes collection instructions for your physician or nurse/midwife. (Collection kits will be shipped to the address you provided unless we are notified otherwise).

Regulations require confirmation of your health status within <u>48 hours of delivery</u>. Included in the kit will be a copy of your completed Health Questionnaire (B.1-1) and a Verification Form (B.1-5). Form B.1-5 must be completed the day of delivery prior to forwarding the cord blood unit to Cord For Life. Update any changes to your health status between the time you filled out your original forms and the time you deliver. Please keep it with the kit so that you can fill out the verification card at that time and send it back with the cord blood.

Bring the collection kit to your hospital or birthing center when you go into labor. Your doctor or hospital's delivery staff will collect the blood remaining in the umbilical cord and placenta after your baby is delivered. Your blood will also be drawn during labor or after delivery.

Notify Cord For Life <u>when in active labor and after the birth of your baby</u>, day or night. It is <u>your</u> responsibility to ensure that we are notified within two hours after the collection of the cord blood.

*Full First Name, Middle Initial or Full Middle Name, Last Name

Form Number: B.1-6 Rev. M.



DONOR INFORMATION AND HEALTH HISTORY

MOTHER'S <u>LAST</u> NAME <u>FIRST</u> NAME						M.I.	LAST 4 S	SS# DIGITS
BEST CONTACT PHONE:	EMAIL	-					MOTHER	r'S DOB:
			L OUT!					- TIROORE
ADDRESS			CITY				STATE	ZIPCODE
FATHER'S <u>LAST</u> NAME			FIRST NAME				M.I.	LAST 4 SS# DIGITS(OPTIONAL)
BEST CONTACT PHONE:	EMAIL						FATHERS	S DOB:
ADDRESS			CITY				STATE	ZIP CODE
Abbitess			OII I				OIAIL	Zii GODE
BABY'S DUE DATE:								
DELIVERY PHYSICIAN'S NAME				PHONE				
CLINIC NAME				1				
DELIVERY HOSPITAL NAME				PHONE				
DELIVERY HOST HAE WAINE				I HOILE				
HOSPITAL ADDRESS			CITY				STATI	E ZIP CODE
BARV'	e DAG	CE AND ET	HNICITY INFORI	MATION				
Since certain HLA Types may be more common						ting a	ord bloo	od unit for transplant.
Baby's Ethnicity: Response is required	, plea	se check o	ne. 🗆 Hispanic	or Latin	10		Not Hi	ispanic or Latino
Baby's Race: Response is required.)f whic	ch group(s)	is vour baby a me	ember? (Selec	t all t	hat apr	olv.)
American Indian or Alaska Native		or African A	-		Asia		a. app	,.,
<u></u>					<i>-</i> 10.00		. //	OLU)
Alaska Native or Aleut (ALANAM) North American Indian (AMIND)		frican (AFB)	(4454)		-		ninese (N	,
American Indian (AMIND)		frican American	` '		-			lipino) (FILI)
American (AMIND)	-	lack Caribbean	(CARB) entral American				oanese (J	
Caribbean Indian (AMIND)		SCAMB)	entrai American		-		rean (KO	,
	(4	SCAIVID)			-			(SCSEAI)
					-			(SCSEAI)
						Oil	ier South	east Asian (SCSEAI)
Native Hawaiian or Other Pacific Islander	White	<u> </u>						
		-	~ (CALI)	_	Γ	No	rthern Eu	ıropean (CAU)
Guamanian (OPI)		astern Europea		_	F	_		ropean (CAU)
Hawaiian (HAWI)		lediterranean (C			-			bean (CAU)
Samoan (OPI)	Middle Eastern (MENAFC) North Coast of Africa (MENAFC)				-			n or Central American
Other Pacific Islander (OPI)	-		` '				4U)	
	N	orth American (CAU)		H		ner White	(CAU)
					<u> </u>			` '

Please read the following Health Questionnaire <u>carefully</u>. You may contact Cord For Life, Inc. (CFL), if you need help understanding any of the questions, please call: 1-800-869-8608 outside of the Orlando area, or 407-834-8333 in the Orlando area.

Completion of all the requested information on the health questionnaire is required before a cord blood unit can be eligible for transplant. This is the only opportunity the cord blood center has to gather this important information from you. An incomplete questionnaire will result in disqualification. The questionnaire should be filled out privately by the expectant mother only, or in a private interview by an approved screener. Your answers to these questions are confidential. Cord For Life, Inc. (CFL) Notice of Privacy Practices are available upon request.

If after being accepted into this program or after your baby's cord blood is collected, you learn of a reason which would exclude you from donating or feel that it should not be transfused to a patient, please call Cord For Life, Inc. (CFL). You will not be penalized from withdrawing from the program at any time.

My signature below confirms that the information provided on Pages 1-8 of Form, B.1-1 is true and accurate to the best of my knowledge.

EXPECTANT MOTHER SIGNATURE:	DATE:



MOTHER'S LAST 4 SS# DIGITS: _____ MOTHER'S DOB:

	CORD BLOOD MATERNAL QUESTIONS		
	Please <u>read carefully</u> and <u>answer each</u> of the following questions <u>individually</u> "Y" for "YES" or "N" for		·-
	Please provide details including dates, where requested, for all "Y" responses (except for #38 and	#73)	
1	Have you ever donated or attempted to donate cord blood using your current or a different name to Cryobanks International , Lifeforce Cryobank Sciences, Inc. (LFC) or Cord For Life, Inc. (CFL)? Details:	Υ□	N□
2	Have you, for any reason, been deferred or refused as a blood or cord blood donor, or been told not to donate blood or cord blood? <i>If yes</i> , why?	Υ□	N□
3	Have you taken any of the following medications (check all that apply): a. □ Insulin from cows (bovine or beef insulin) since 1980? b. □ Growth hormone from human pituitary glands ever?	Υ□	N□
<u> </u>	In the past 8 weeks, have you had any shots or vaccinations?		
4	If yes, details:	Υ□	N□
5	In the past 12 weeks, have you had contact with someone who has received the smallpox vaccine? (Examples of contact include physical intimacy, touching the vaccination site, touching the bandages or covering the vaccination site, or handling bedding or clothing that had been in contact with an unbandaged vaccination site) Details:	Υ□	N 🗆
6	In the past 4 months, have you experienced <u>TWO (2)</u> or more of the following: a fever (>100.5°F or 38.6°C), headache, muscle weakness, skin rash on trunk of the body, swollen lymph glands? If yes , which symptoms and when? Details:	Υ□	Ν□
7	Have you ever had any type of cancer, including leukemia? If yes, details:	Υ□	N□
8	During your pregnancy, have you been diagnosed with West Nile Virus or had a positive test for West Nile Virus?	Υ□	N□
9	Have you ever had a past diagnosis of clinical, symptomatic viral hepatitis after age 11? <i>If yes,</i> details, with dates:	Υ□	N□
10	Have you ever had a parasitic blood disease such as Leishmaniasis, Chagas disease or Babesiosis or any positive test for Chagas or T. cruzi, including screening tests?	Υ□	N□
11	Have you ever been diagnosed with Creutzfeldt-Jakob Disease (CJD), Variant CJD, dementia, degenerative or demyelinating disease of the central nervous system, or other neurological disease where the cause is unknown?	Υ□	N□
12	Have any of your blood relatives ever been diagnosed with Creutzfeldt-Jakob Disease (CJD), or have you been told that your family has an increased risk for CJD?	Υ□	N□
13	Have you received a dura mater (brain covering) graft?	Υ□	N□
14	Have you ever had a transplant or other medical procedure that involved being exposed to live cells, tissues, or organs from an animal ? Details:	Υ□	N□
15	Have you ever lived with or had sexual contact with anyone who had a transplant or other medical procedure that involved being exposed to live cells, tissues, or organs from an animal ? If yes, details:	Υ□	N 🗆
16	In the past 3 years, have you had malaria? If yes, details:	Υ□	N□
17	In the past 3 years, have you been outside the United States or Canada? Where:	Υ□	N 🗆
18	In the past 12 months, have you had a blood transfusion? Details:	Υ□	N□
19	In the past 12 months, have you had a transplant or tissue graft from someone other than yourself, such as organ, bone marrow, stem cell, cornea, bone, skin or other tissue?	Υ□	N□
20	In the past 12 months, have you had a tattoo or piercing (ear, skin or body)? If yes, please indicate type and answer question 21. If no, skip to question 22. Type: Tattoo Piercing, details:	Υ□	N 🗆
	21. If yes, were shared or non-sterile instruments, needles, or inks used for the tattoo or piercing?	Υ□	N□

HEALTH QUESTIONNAIRE

Form Number: B.1-1 Rev. X

Date Issued: 08MAR2024



MOTHER'S LAST 4 SS# DIGITS: MOTHER'S DOB: In the past 12 months, have you had an accidental needle stick, or have you come into contact with someone else's Υ□ $N \square$ 22 blood through an open wound (for example, a cut or sore), non-intact skin, or mucous membrane (for example, into your eye, mouth, etc.)? Details: In the past 12 months, have you had or been treated for a sexually transmitted disease, including syphilis? Υ□ 23 $N \square$ If yes, details with dates: 24 ΥΠ $N \square$ In the past 12 months have you given money, drugs, or other payment to anyone to have sex with you? In the past 12 months have you had sex with anyone who has taken money, drugs, or other payment in exchange for Υ□ 25 $N \square$ sex in the past 5 years? In the past 12 months, have you had sexual contact or lived with a person who has active or chronic viral hepatitis B or Υ□ $N \square$ 26 Hepatitis C? In the past 12 months, have you had sex, even once, with anyone who has used a needle to take drugs, steroids, or Υ□ $N \square$ 27 anything else not prescribed by a doctor in the past 5 years? In the past 12 months, have you had sex with a male who has had sex with another male, even once, in the past 5 Y□ $N \square$ 28 years? In the past 12 months, have you had sex, even once, with anyone who has HIV/AIDS or had a positive test for the AIDS 29 YΠ $N \square$ virus? Υ□ $N \square$ 30 In the past 12 months, have you been in juvenile detention, lockup, jail or prison for more than 72 continuous hours? 31 In the past 5 years have you received money, drugs, or other payment for sex? Y□ $N \square$ In the past 5 years, have you used a needle, even once, to take drugs, steroids or anything else not prescribed for you Υ□ $N \square$ 32 by a doctor? Υ□ 33 $N \square$ Do you have AIDS, or have you ever tested positive for HIV (including screening tests)? Do you have any of the following: Υ□ $N \square$ A) Unexplained night sweats? Y□ $N \square$ B) Unexplained blue or purple spots on or under the skin or mucous membranes? Υ□ $N \square$ C) Unexplained weight loss? YΠ $N \square$ D) Unexplained persistent diarrhea? 34 ΥΠ $N \square$ E) Unexplained cough or shortness of breath? YΠ F) Unexplained temperature higher than 100.5°F (38.6°C) for more than 10 days? $N \square$ Y□ $N \square$ G) Unexplained persistent white spots or sores in the mouth? Υ□ $N \square$ H) Lumps in your neck, armpits, or groin lasting longer than one month? ΥΠ $N \square$ I) Any infection during your pregnancy? Have you ever tested positive for HTLV-Human T-cell Lymphotrophic Virus (including screening tests) or had unexplained Υ□ 35 $N \square$ paraparesis (partial paralysis affecting the lower limbs)? Do you understand that if you have the AIDS virus, you can give it to someone else even though you may feel well Υ□ $N \square$ and have a negative AIDS test?



MOTHER'S LAST 4 SS# DIGITS: _____ MOTHER'S DOB: _____

FOR USE WITH QUESTIONS #39 – 42 – COUNTRIES DEFINED AS EUROPE										
	<u>ALBANIA</u>	Travel	Resident	GREECETravelResident ROMANIATrave	l	Resident				
	Date(s):			Date(s):						
	Total Time:			Total Time: Total Time:						
	AUSTRIA	Travel	Resident	HUNGARYTravelResident SLOVAK REPUBLICTrave	el	_Resident				
	Date(s):			Date(s): Date(s):						
	Total Time:			Total Time: Total Time:						
	BELGIUM	Travel	Resident	IRELAND (REPUBLIC DF)TravelResidentTravelTravelTravelTravelTravelTravelTravelTravelTravelTravelTravelTravel	el	_Resident				
	Date(s):			Date(s): Total Time: Total Time:						
	Total Time:									
	BOSNIA-HERZEGOVINA	Travel	Resident	Travel Resident SPAIN Travel Travel	el	_Resident				
	Date(s): Total Time:			Date(s): Total Time: Total Time:						
		.								
	BULGARIA Date(s):	Travel	Resident	LIECHTENSTEINTravelResident	el	_Resident				
	Total Time:			Total Time:						
		т .	П 11 г		ı	пи				
	CROATIA Date(s):	Travel	Resident	LUXEMBOURGTravelResident	<u></u>	_Resident				
	Total Time:			Total Time: Total Time:						
	CZECH REPUBLIC	Travel	Resident	MACEDONIA Travel Resident UNITED KINGDOM (UK) includes England, No	nthonn Ino	land				
	Date(s):		KGSIGGIIL	Date(s): Scotland, Wales, Isle of Man, Channel Island						
	Total Time:			Total Time: Falkland IslandsTravi	el	_Resident.				
	DENMARK	Travel	Resident	NETHERLANDS (HOLLAND) Travel Resident						
	Date(s):			Date(s):						
	Total Time:			Total Time:						
	<u>FINLAND</u>	Travel	Resident	NORWAY Travel Resident YUGOSLAVIA (FEDERAL REPBULIC OF) Trav	el	Resident				
	Date(s):			Date(s):						
	Total Time:			Total Time: Total Time:						
	FRANCE	Travel	Resident	PDLANDTravelResident KOSOVO, MONTENEGRO, SERBIATravel	el	_Resident				
	Date(s):			Date(s):						
	Total Time:			Total Time: Total Time:						
	GERMANY _	Travel	Resident	PORTUGAL Travel Resident						
	Date(s): Total Time:			Date(s): Total Time:						
				raveled to Europe? (refer to chart above) If no, skip to question 42. check in all the appropriate box(es) above to identify the country(ies), reason,						
37	a) Use the character date(s) and			brieck in all the appropriate box(es) above to identify the country(ies), reason,	Y□	N□				
	b) Answer que									
	38. From 1980	through 19	996, did you	spend time that adds up to 3 months or more in the United Kingdom (refer to	Υ□	N□				
	chart above)?					.,				
	39. Since 1980	, have you	received a	ransfusion of blood or blood components while in the UK or France?	Υ□	N□				
	40. Since 1980	, have you	spent time t	hat adds up to 5 years or more in Europe (refer to chart above), including time	Υ□	N□				
	spent in the UK					.,				
41				member of the U.S. military, a civilian military employee, or a dependent of a	Υ□	N□				
	member of the L	J.S. military	<i>'</i> ?							
42				nd a total of 6 months or more associated with a military base in any of the	Y□	N□				
	following countri	es: United	Kingdom, B	elgium, Netherlands or Germany?	- -					
43				nd a total of 6 months or more associated with a military base in any of the	Υ□	N□				
	following countri	es: Spain,	Portugal, Τι	rkey, Italy or Greece?						



MOTHER'S DOB: MOTHER'S LAST 4 SS# DIGITS: _____

	FOR USE WITH QUESTIONS 46-48: AFRICAN COUNTRIES		
l	BENINTravelResident BEUATORIAL GUINEATravelResident BEUATORIAL GUINEATravelResident BEUATORIAL GUINEATravelResident BEUATORIAL GUINEATravelResident BEUATORIAL GUINEATravelResident BEUATORIAL GUINEATravel	vel	_Resident
li	CAMEROONTravelResident GABONTravelResident TOGOTr Date(s):	avel	Resident
li	CENTRAL AFRICAN REPUBLIC Travel Resident KENYA Travel Resident ZAMBIA Travel Date(s): Oate(s): Total Time: Total Time: Total Time: Total Time:	avel	Resident
l	CHAD Travel Resident NIGER Travel Resident Date(s): Total Time: Total Time:		
l	CONGO Travel Resident NIGERIA Travel Resident Date(s): Total Time: Total Time:		
44	Since 1977, were you born in, have you lived in, or have you traveled to any African country listed above? If yes, answer question 45. If no, skip to question 46. a) Use the chart above and place a check in all the appropriate box(es) above to identify the country(ies), reason, date(s) and total time that apply.	Υ□	N□
	45. While in one of the African countries listed above, did you receive a blood transfusion or any other medical treatment with a product made from blood?	Υ□	N□
46	Have you had sexual contact with anyone who was born in or lived in any African country listed above since 1977?	Υ□	N□
47	a.) Were you and/or the baby's father adopted at early childhood?	Υ□	N
47	b.) If yes, is a family medical history available for you and/or the baby's father?	Υ□	N
48	Are you and the baby's father related, except by marriage? (e.g., first cousins)	Υ□	N□
	A.) Did this pregnancy use either a donor egg or donor sperm?	Υ□	N□
49	B.) If yes, is a family medical history questionnaire available for the egg or sperm donor? (please attach copy) Name of the Clinic:	Υ□	N□
	Have you ever had an abnormal result from a prenatal test (e.g., amniocentesis, blood test, and ultrasound)? If yes, answer the following questions. If no, skip to question 51.	Υ□	N□
50	A) Which test was abnormal?		
	B) What was the abnormal test result?		
	C) Was a diagnosis made? Specify diagnosis:		
51	Have you had any children who died within the first 10 years of life?	Υ□	N□
	If yes, what was the cause?		
52	Have you ever had a stillborn child?	Υ□	N□
F^	If yes, what was the cause?		N -
53 54	Have you had a medical diagnosis of ZIKV (Zika) infection at any point during your pregnancy?	Υ□	N□
54	Have you resided in, or travel to, an area with active ZIKV (Zika) transmission at any point during your pregnancy?		N□
55	Have you had sex during your pregnancy with a male or female who is known to have: a) A medical diagnosis of ZIKV (Zika) within the six months prior to that contact.	Υ□	N□
	b) Resided in, or traveled to, an area with active ZIKV (Zika) transmission within the six months prior to that contact.	<u></u>	

Form Number: B.1-1 Rev. X



MOTHER'S LAST 4 SS# DIGITS: _____ MOTHER'S DOB: _____

D) Hereditary Hemophagocytic Lymphohistiocytosis (HLH) including FEL E) Hypoglobulinemia F) Nezeloff Syndrome G) Severe Combined Immunodeficiency						Y MEDICAL HISTORY							
Cancer or Leukemia? Y N N					wing codes	to describe the relationship between t	he baby and BMS Bab	l a family n	nember w er Sibli	rith a dis no	sease:		
Cancer or Leukemia? Y N			BF	Baby's Fa	ther	BS Baby's sibling	BFS Bab	y's Fath	er's Sib	ling			
1	F					d uncles by blood and does <u><i>not</i> inclu</u>	de aunts an	id uncles	who are	ın-laws	of the	parents.)	
A) Brain or other nervous system cancer B) Bone or joint cancer C) Kidney (including renal pelvic) cancer D) Thyroid Cancer E) Hodgkin's Lymphoma F) Non-Hodgkin's Lymphoma G) Acute or chronic myelogenous/myeloid leukemia H) Acute or chronic myelogenous/myeloid leukemia H) Acute or chronic lymphocytic/lymphoblastic leukemia G) Skin Cancer J) Other cancer/leukemia Specify Type: Specify Type:	Į	Ļ			_			_					
B) Bone or joint cancer	5	56	If yes, please specify all that app	ly. <i>If no</i> , p	proceed	to next question.	BN	1 В	F E	S			
C) Kidney (including renal pelvic) cancer				n cancer									
D) Thyroid Cancer			<u> </u>							_			
F) Hodgkin's Lymphoma			· · · · · · · · · · · · · · · · · · ·) cancer						_			
F) Non-Hodgkins Lymphoma										_	INA	MEDIA	TE
F) Notif-Rodgin's Symforome			· · · · · · · · · · · · · · · · · · ·							_			
H) Acute or chronic myelogenous/myeloid leukemia H) Acute or chronic lymphocytic/lymphoblastic leukemia I) Skin Cancer J) Other cancer/leukemia Specify Type: Specify Type		ŀ								_			
1) Skin Cancer			· · · · · · · · · · · · · · · · · · ·							_			
J) Other cancer/leukemia Specify Type: Specify all that apply. If no, proceed to next question. BM BF BS BGP BMS BFS A) Diamond-Blackfan Syndrome Specify all that apply. If no, proceed to next question. White Blood Cell Disease? White Blood Cell Disease? Y N N Spherocytosis Specify all that apply. If no, proceed to next question. BM BF BS BGP BMS BFS A) Chronic Granulomatous Disease Specify all that apply. If no, proceed to next question. Specify Type: Spec		ŀ	<u> </u>	lymphobla	astic leuk	emia				_			
Specify Type:			,] [
Red Blood Cell			•										
Red Blood Cell	L						Iп		, ,	, I			
If yes, please specify all that apply. If no, proceed to next question. BM BF BS BGP BMS BFS			Specify Type:										
A) Diamond-Blackfan Syndrome	Г		Red Blood Cell Y	. N □			•						
B Elliptocytosis			If yes, please specify all that app	ly. <i>If no</i> , j	proceed	to next question.	BM	l BF	В	S	BGP	BMS	BFS
B) Elliptocytosis C) G6PD or other red cell enzyme deficiency D) Spherocytosis White Blood Cell Disease? Y N N White Blood Cell Disease? Y N N If yes, please specify all that apply. If no, proceed to next question. A) Chronic Granulomatous Disease B). Kostmann Syndrome. C) Schwachman-Diamond Syndrome D) Leukocyte Adhesion Deficiency (LAD) Immune Deficiencies? Y N N If yes, please specify all that apply. If no, proceed to next question. BM BF BS BGP BMS BFS A) ADA or PNP Deficiency B) Combined Immunodeficiency Syndrome (CID), Common Variable Immunodeficiency Disease (CVID) C) DiGeorge Syndrome D) Hereditary Hemophagocytic Lymphohisticoytosis (HLH) including FEL E) Hypoglobulinemia F) Nezeloff Syndrome G) Severe Combined Immunodeficiency	,	57		!]			
D) Spherocytosis	Ι,	"	B) Elliptocytosis]			
White Blood Cell Disease? Y			C) G6PD or other red cell enzym	ne deficien	су]			
If yes, please specify all that apply. If no, proceed to next question. BM BF BS BGP BMS BFS	L		D) Spherocytosis]			
A) Chronic Granulomatous Disease B). Kostmann Syndrome. C) Schwachman-Diamond Syndrome D) Leukocyte Adhesion Deficiency (LAD) Immune Deficiencies? Y													
B). Kostmann Syndrome. C) Schwachman-Diamond Syndrome D) Leukocyte Adhesion Deficiency (LAD) Immune Deficiencies? If yes, please specify all that apply. If no, proceed to next question. A) ADA or PNP Deficiency B) Combined Immunodeficiency Syndrome (CID), Common Variable Immunodeficiency Disease (CVID) C) DiGeorge Syndrome D) Hereditary Hemophagocytic Lymphohistiocytosis (HLH) including FEL E) Hypoglobulinemia F) Nezeloff Syndrome G) Severe Combined Immunodeficiency	L				proceed t	to next question.	BM	l BF	В	S	BGP	BMS	BFS
B). Kostmann Syndrome. C) Schwachman-Diamond Syndrome D) Leukocyte Adhesion Deficiency (LAD) Immune Deficiencies? If yes, please specify all that apply. If no, proceed to next question. A) ADA or PNP Deficiency B) Combined Immunodeficiency Syndrome (CID), Common Variable Immunodeficiency Disease (CVID) C) DiGeorge Syndrome D) Hereditary Hemophagocytic Lymphohistiocytosis (HLH) including FEL E) Hypoglobulinemia F) Nezeloff Syndrome G) Severe Combined Immunodeficiency C) Schwachman-Diamond Syndrome D	58		•	ase]			
D) Leukocyte Adhesion Deficiency (LAD) Immune Deficiencies? Y		'	B). Kostmann Syndrome.]			
Immune Deficiencies? Y			C) Schwachman-Diamond Synd	rome]			
If yes, please specify all that apply. If no, proceed to next question. BM BF BS BGP BMS BFS	L		D) Leukocyte Adhesion Deficien	cy (LAD)]			
A) ADA or PNP Deficiency B) Combined Immunodeficiency Syndrome (CID), Common Variable	Г		Immune Deficiencies?	Υ□	N 🗆								
B) Combined Immunodeficiency Syndrome (CID), Common Variable			If yes, please specify all that app	ly. <i>If no</i> , p	proceed	to next question.	ВМ	l BF	В	S	BGP	BMS	BFS
Immunodeficiency Disease (CVID) C) DiGeorge Syndrome D) Hereditary Hemophagocytic Lymphohistiocytosis (HLH) including FEL E) Hypoglobulinemia F) Nezeloff Syndrome G) Severe Combined Immunodeficiency		A)	ADA or PNP Deficiency]		
D) Hereditary Hemophagocytic Lymphohistiocytosis (HLH) including FEL				ndrome (C	CID), Con	nmon Variable]		
E) Hypoglobulinemia	59	C	DiGeorge Syndrome]		
F) Nezeloff Syndrome		D)	Hereditary Hemophagocytic Lym	phohistiod	cytosis (F	ILH) including FEL]		
G) Severe Combined Immunodeficiency		E)	Hypoglobulinemia]		
		F)	Nezeloff Syndrome]		
II) Wishan Aldrick Condense		G	Severe Combined Immunodefici	ency]		
H) Wiskott-Alarich Syndrome		H	Wiskott-Aldrich Syndrome]		



MOTHER'S LAST 4 SS# DIGITS: _____ MOTHER'S DOB: _____

	Platelet Disease?	Y	′ 🗆	N 🗆							
	If yes, please specify all that apply. If no, proceed to next quest	ion			ВМ	BF	BS	BGP	вмѕ	BFS	
	A) Amegakaryocytic Thrombocytopenia										
60	B) Glanzmann Thrombasthenia										
	C) Hereditary Thrombocytopenia										
	D) Platelet Storage Pool Disease										
	E) Thrombocytopenia with absent radii (TAR)										
	F) Ataxia-Telangiectasia										
<u> </u>	G) Fanconi Anemia	1									
61	Other blood disorder or problem	Υ	′ 🗆	N 🗆							
Hem	oglobin Problems				ВМ	BF	BS	BGP	BMS	BFS	
62	Sickle cell disease, such as sickle-cell anemia or sickle thalassem	ia?	Υ□	N□							
	Specify disease:		T								
63	Thalassemia, such as alpha thalassemia or beta-thalassemia?		Υ□	N 🗆							
	Metabolic/Storage Disease?		Υ□	N 🗆							
	If yes, please specify all that apply. If no, proceed to next quest	ion.			ВМ	BF	BS	BGP	BMS	BFS	
	A) Hurler Syndrome (MPS I)										
	B) Hurler-Scheie Syndrome (MPS I H-S)										
	C) Hunter Syndrome (MPS II)										
	D) Sanfilippo Syndrome (MPS III)										
	E) Morquio Syndrome (MPS IV)										
	F) Maroteaux-Lamy Syndrome (MPS VI)										
	G) Sly Syndrome (MPS VII)										
64	H) I-cell disease										
	Globoid Leukodystrophy (Krabbe Disease)										
	J) Metachromatic Leukodystrophy (MLD)										
	K) Adrenoleukodystrophy (ALD)										
	L) Sandhoff Disease										
	M) Tay-Sachs Disease										
	N) Gaucher Disease										
	0) Niemann Pick-Disease										
	P) Porphyria										
	Q) Other or unknown metabolic/storage disease, Details:										
	Acquired Immune System Disorders				ВМ	BF	BS			-	
65	HIV/AIDS?		Y 🗆	N□							
	Severe autoimmune disorder?		Y 🗆	N 🗆							
	If yes, please specify all that apply including providing the other reinformation. If no, proceed to next question.	eques	sted		ВМ	BF	BS				
	A) Crohn's Disease or Ulcerative Colitis							IMMEDIATE FAMILY ONLY			
66	B) Lupus										
66	C) Multiple Sclerosis (MS)										
	D) Rheumatoid Arthritis										
	Diagnosis Date:					_					
	Currently under MD care? Describe:	_									
	Are you currently taking any medication? Name:										



МО	MOTHER'S LAST 4 SS# DIGITS: MOTHER'S DOB:								
67	Any diagnosis of other or unknown immune system disorder? Specify Disorder:							IATE FA	AMILY
	Openiy Disorder.			ВМ	BF	BS	BGP	BMS	BFS
68	Required Chronic Blood Transfusions?	Υ□	N□						
69	Been told you or your family member(s) have hemolytic anemia?	Υ□	Ν□						
70	Had spleen removed to treat a blood disorder?	Y	N 🗆						
71	Had gallbladder removed before the age of 30?	Y	N 🗆						
72	Had Creutzfeldt-Jakob disease (CJD)?	Y	N \square						
	Other serious or life-threatening diseases affecting the family?	Y□	N 🗆						
	If yes, list affected family member(s) and type of disease			BM	BF	BS	BGP	BMS	BFS
73	Specify Type:								
13	Specify Type:								
	Specify Type:								
74	In answering these questions, have you answered for both your family and the baby's father's family?								
Adde	ndum A:								
1.	Any history of acute respiratory disease? If Yes, please describe							Υ□	N□
2.	Any active tuberculosis disease or history of tuberculosis therapy? It describe	f Yes, ple	ease					Y□	N□
3.	Any history of drug or alcohol abuse? If Yes, please describe							Υ□	N□
Add	endum B: COVID-19								
1.	Have you received a Covid-19 vaccination?						,	Υ□	N□
	TAL REVIEW TO BE COMPLETED BY CFL AFFILIATE COLLECT								
	ve performed and reviewed the above responses and have determine				-	-			
Rev	Acceptable –All CFL HQ requirements met. □ Follow Up iewed By:	– Furtne	r tollow (•	∟ require e(s):	ea for fina	ai status	aetermii	nation.
	•			Date	J(3).				
CFL	REVIEW TO BE COMPLETED BY CORD FOR LIFE, INC (CFI CLIENT SERVICES REVIEW (☑ one) □ N/A	L) ONLY		ARORAI		EVIEW (
	HQ-OK Defer] HQ-		ADONA		□ Def			
	Unusual Findings Ineligible] Unu	sual Fir	ndings			ligible		
	Other:								
∣Rev	eviewed By: Date(s): Reviewed By: Date(s):								

Form Number: B.1-1 Rev. X



DONATION INFORMED CONSENT AND RELEASE - HOSPITAL/BIRTHING CENTER

I, the undersigned, desire the collection of my unborn baby's cord blood for donation. I have elected to utilize the services of Cord For Life to achieve the desired donation. For the donation to occur it is necessary to collect and save the blood from the placenta and umbilical cord after the birth of my baby, rather than discard the blood as medical waste. The collected cord blood will be shipped to Cord For Life for processing and placement into storage.

My physician, physician's designee, midwife or a Cord For Life trained and collection specialist will perform the collection of the cord blood after the delivery of my baby, while the delivery of the placenta occurs. He/she will use methods provided by Cord For Life in their standard operational procedures. Medical conditions may arise which preclude the collection of the cord blood and will be decided at the sole discretion of the attending physician.

I understand that the donation of cord blood includes medical procedures and that there can be no guarantee or assurance of success of the results of the service. I further, on behalf of myself and my unborn baby, our respective heirs, successors and assigns, hereby release and forever hold harmless the Hospital / Birthing Center, and its affiliates, successors, assigns, officers, directors, employees and agents from any and all actions, causes of action, claims, debts, demands, liabilities, covenants, controversies, omissions and damages and any and all other claims of every kind, nature, and description whatsoever, both in law and equity, which may arise relating to the collection of the cord blood on behalf of me and my unborn baby.

I approve the sharing of any/all testing results with other medical or research facilities that are in partnership with Cord For Life and whose standards and policies follow all confidentiality measures as required by the Health Insurance Portability and Accountability Act of 1996 (*HIPAA*).

PHYSICIAN - DONATED SAMPLE

My patient desires the collection of her unborn baby's cord blood for donation to Cord For Life. For the donation to occur, it is necessary to collect and save the blood from the placenta and umbilical cord after the birth of my patient's baby, rather than discard the blood as medical waste. The cord blood obtained will be shipped to Cord For Life for processing and placement into storage.

Myself or a Cord For Life trained and approved collection specialist will perform the collection of the cord blood after the birth of her baby, while the delivery of the placenta occurs. The collection will use the methods provided by Cord For Life in their standard operational procedures. The collection period will be brief and Cord For Life will provide the protocols and collection equipment in the kit. Every effort will be used to acquire as much cord blood as is feasible and will minimize the risk of fungal, bacterial or maternal blood contamination.

The health and welfare of my patient and her baby are the primary concern and responsibility and accordingly I reserve the right to forgo the collection of the cord blood if my best medical judgment indicates this to be necessary.

I understand that the donation of cord blood includes medical procedures and that there can be no guarantee or assurance of success of the results of the service. I, on behalf of myself, my heirs and successors and assigns hereby release and forever discharge Cord For Life and its affiliates, successors, assigns, officers, directors, employees and agents from any and all actions, causes of actions, demands, debts, claims liabilities, covenants and damages and any and all other claims of every kind, nature and description whatever, both in law and equity, which may arise relating to my performing the collection of the cord blood.

perioriting the concentration of the cord	biood.		
forever discharges me and each of n demands, debts, claims, liabilities, co	ny heirs, succe ovenants and o	es, assigns, officers, directors, employees and agessors and assigns from any and all actions, caudamages and any and all other claims of every kind may arise relating to my performing the collection	ses of actions, ind, nature and
My patient.		, releases me and each of my heirs, succes	sors, and assigns from
damages and any and all other claim may arise relating to my performing t In addition, I understand that	ns of every king the collection of the donation or my services	of cord blood is a voluntary program, and as suc in the collection of the cord blood unit. I hereby	and equity, which h, I will <i>not</i> receive
Signature of Expectant Mother (Required)	Date	Signature of Physician/Midwife ((Required)	Date
Print Full Name of Expectant Mother		Print Name (Physician/Midwife)	
IMPORTANT: THIS PAGE IS REQUIRED TO BE SIGNED BY V	יחון אאח אחווף פאיכור	IVN /WIDWIEE IN UBDEB TO BECEINE Y GUBD EUB LIEE. GUBD BLUUD DUNYTIUN	LUITELLIUN KIT TU VAUIU VNA

<u>MPPORTANT:</u> THIS PAGE IS **required** to be signed by you and your physician/midwife in order to receive a cord for life cord blood donation collection kit. To avoid any delays in your paperwork review, please ensure that all required signatures are present prior to submitting your forms.



INFORMED CONSENT FOR THE INFECTIOUS DISEASE TESTING

HUMAN IMMUNODEFICIENCY VIRUS AND TRANSMISSION:

Human Immunodeficiency Virus (HIV) is a virus which can be transmitted from individuals through body fluids, primarily blood and semen. The spread is not through air or food or by casual social contact. It is passed on when the blood or body fluids of an infected person mix with your own. Sexual transmission is mainly the result of the transfer of and exposure to infected semen. Women as well as men can transmit the virus sexually. The HIV virus has also been detected in vaginal secretions, tears, and saliva, but exposure to saliva has not been proven to transmit the infection. Intravenous drug users and persons receiving blood transfusions can be exposed to the virus through infected blood or body products. A baby may become infected during pregnancy, delivery, or when breast feeding if its mother has the disease. A person may carry the virus for months before testing positive and may carry the virus for months or years before the symptoms appear. An HIV positive person can still spread the disease even though he or she may appear healthy.

When HIV enters the blood stream it invades and destroys cells in the body's infection and cancer fighting system and reduces the body's ability to fight infections. The HIV virus leads to the depletion of the immune system to a point that infections which one wouldn't normally get (opportunistic infections) start developing, at which point the patient has AIDS. The HIV virus is not what kills a person with AIDS, it is the opportunistic infections which cause death.

BEHAVIORS THAT INCREASE YOUR RISK OF BEING EXPOSED TO HIV:

Recent blood, plasma, or blood product transfusion, intravenous drug use, especially with sharing of needles or syringes, or having sexual contact with someone who: has tested positive for HIV infection, is at risk of infection through sexual practices, IV drug use, or recent blood transfusion, uses illicit intravenous drugs, received blood transfusions, plasma, or clotting factor before 1985 or within the last twelve months, has more than one sexual partner, especially ones who could be at risk of HIV infection, or is a man who has had sexual relations with another man.

THE HIV TEST AND VOLUNTARY TESTING

The HIV tests are blood tests for the presence of the HIV virus and antibodies to the HIV virus. A positive test result means that you have been exposed to the virus, and either have made antibodies or are infected. It may not mean that you have AIDS now or that you will become sick with AIDS in the future. A negative test means that you are probably not infected with the virus. It takes about 12 days to detect the virus from time of infection to time of detection.

Taking the HIV test is voluntary, and results are confidential by law. Results can only be given to people you allow, and a release form must be signed prior to releasing this information. The law requires Cord For Life to report any positive HIV test result to the County Health Department.

CONSENT (REQUIRED)	
	ad my questions about the HIV test answered. I agree to take the HIV ole to Cord For Life and to my private physician,
Printed Full Name of Expectant Mother's:	Date:
Expectant Mother's Signature (full name as printed abo	ve):
It is an FDA requirement that Cord For	me: r Life performs maternal blood testing. Tubes will be included with the
RX Patient Na It is an FDA requirement that Cord For	me:
It is an FDA requirement that Cord Forcord blood collection kit to be drawn at the	me: r Life performs maternal blood testing. Tubes will be included with the hospital/birthing center during labor and delivery.

MPORTANT: THIS PAGE IS REQUIRED TO BE SIGNED BY YOU AND YOUR PHYSICIAN/MIDWIFE IN ORDER TO RECEIVE A CORD FOR LIFE CORD BLOOD DONATION COLLECTION KIT. TO AVOID ANY DELAYS IN YOUR PAPERWORK REVIEW, PLEASE ENSURE THAT ALL REQUIRED SIGNATURES ARE PRESENT PRIOR TO SUBMITTING YOUR FORMS.



PHYSICIAN/MIDWIFE CB COLLECTION TRAINING REQUEST

Dear Healthcare Professional,

As you now know, your patient desires to have her baby's umbilical cord blood collected for either private storage or public donation. Cord For Life is registered with the FDA, a member of the National Marrow Donor Program and is accredited by the AABB, and we must ensure in every way possible that the collection process is successful. A quality collection is the biggest predictor in converting a donated cord blood unit into a transplantable product.

To ensure a high quality, high volume sample, we would like to offer you, at no cost, self training for the collection procedure. You also have the ability to state that you are well aware of the collection procedure and do not desire any further information. A one-page collection instruction sheet is also included in the collection kit sent to the donor mother. The donor mother will bring this kit to the hospital/birthing center at the time of delivery/collection.

We appreciate your time and support of this potentially life-saving program. Without your efforts, we would not be able to meet the increasing demands for stem cell transplants around the world.

IMPORTANT: Response to the following options is <u>mandatory</u> for regulatory compliance. If not completed a collection kit <u>cannot</u> be forwarded:

Phy	sician/N	lidwife Name:
	OPTION 1	I have collected umbilical cord blood before, and I am comfortable with the procedure, I do not require additional training.
	OPTION 2	I will review the electronic collection videos on the Cord For Life website www.cordforlife.com (Healthcare Professional Section).
	OPTION 3	Please contact me regarding alternate collection training options. Phone: E-mail:
Co	omment	s / Recommendations:
_		
_		
_		
For	CORD	FOR LIFE staff, only
□lh	nave verif	Midwife has stated they were previously trained for cord blood collection. ied the Physician/Midwife, named above, has reviewed the Cord For Life Training video. acted the Physician/Midwife, named above, and provided the following
ins		e to verify completion of training for the Physician/Midwife, named above. Collection (Form C.1-2) have been forwarded to ensure adherence to the Cord For Life collection
Emp	oloyee:	Date:

Date Issued: 04OCT2011

Collection Partner of the Cord for Life Foundation Informed Consent for Cord Blood Medical Research

I. INVITATION AND PURPOSE

You are invited to donate your baby's cord blood for medical research if it cannot be used as a transplant product. You are being invited because you have already agreed to donate your baby's cord blood to the **Cord for Life®** Program for patients in need of a transplant. There are many reasons that cord blood may not meet the requirements for transplant. Cord blood not meeting these requirements can be used for medical research.

Cord for Life® Program provides investigators cord blood units to use in medical research. Although the exact studies for which cord blood units may be used is not known at this time, the following are types of studies in which these units may be included: Disease Treatment or Regenerative Medicine under Clinical Trials and Investigational New Biological Drug development.

In addition, researchers will conduct research studies with cord blood units that have had all identifiers removed. In these studies, there will be no way for the unit to be linked to you. The **Cord for Life®** Program may allow researchers to use these anonymous cord blood units for many other kinds of studies. These studies are not limited to the types of studies listed above or related to transplantation in general.

You will receive a follow-up letter requesting follow-up information on your baby's health. It is important that you inform Cord For Life if the infant donor develops a serious disease.

"With this donation of your cord blood, you are agreeing that your donated cord blood may be used for stem cell transplantation (also known as bone marrow transplant), for research and development, or in commercial products that may be used to help people with their medical problems. This donation will not result in any financial payment to you. Once your cord blood is donated you will not have a say in how it will be used for transplant, research or commercial purposes and it cannot be returned once it is used."

II. PROCEDURES

If you agree to donate your baby's cord blood unit for medical research, nothing additional is required from you. After the cord blood is collected it will be tested to see if it meets all the requirements for transplant. If, **and only if**, it does **not** meet the requirements for transplant, the cord blood may be used for medical research. All research studies using cord blood must first be approved by the **Cord for Life®** scientific board, and executive and quality management.

III. POSSIBLE RISKS AND BENEFITS

There are no physical risks to you or your baby by donating the cord blood to be used in medical research. The decision to use the cord blood for medical research is only made after the cord blood is collected and it does not meet the requirements for transplant.

There is a very small risk that an unauthorized person could find out which cord blood unit is your baby's. The Cord for Life® Program has several procedures in place to keep your data private. No identifiable information about you will be given to the researchers, nor will it be published or presented at scientific meetings.

You or your baby will not be helped by donating your baby's cord blood for medical research. However, this research may help future patients who need a transplant or other therapeutic medical treatment.

IV. CONFIDENTIALITY

The Cord for Life® Program follows all HIPAA regulations and will not knowingly disclose that you donated your baby's cord blood for medical research. The Cord for Life® Program will try hard to make sure no one outside the Cord for Life® Program will know which cord blood unit is yours.

V. <u>REIMBURSEMENT AND COSTS</u>

You will not be paid for donating your baby's cord blood for medical research. It will not cost you anything to donate your baby's cord blood for medical research.



Collection Partner of the Cord for Life Foundation

VI. VOLUNTARY PARTICIPATION IN AND WITHDRAWAL

It is up to you if you want to donate your baby's cord blood for medical research. If you choose not to, your unit will be discarded as medical waste.

If you decide to donate your baby's cord blood for medical research, you may change your mind at any time in the future. If you decide you do not want your baby's cord blood used for medical research, your baby's cord blood will be destroyed if it has not already been used. This will not affect your relationship with Cord for Life[®]. To withdraw your unit, forward a notarized letter to the Cord for Life[®] Client Services Department.

VII. ALTERNATIVE TO PARTICIPATION

You may choose not to donate your baby's cord blood for medical research. If you choose not to your unit will be discarded as medical waste.

VIII. QUESTIONS OR CONCERNS

If you have questions, concerns, or complaints about donating your baby's cord blood for medical research contact *Donald Hudspeth (Director of Laboratory Operations)* at dhudspeth@cordforlife.com or Deborah Sardone (*Director of Quality Assurance*) at dsardone@cordforlife.com.

If you have questions or concerns about your rights as a research subject or about potential risks and injuries, please contact *Donald Hudspeth* (*Director of Laboratory Operations*) at dhudspeth@cordforlife.com.

IX. DONOR ADVOCACY

If you have additional concerns and desire information from an impartial source, Cord for Life® suggests visiting the following websites:

NMDP-Be the Match® Program: https://bethematch.org/Support-the-Cause/Donate-cord-blood/

Parents' Guide to Cord Blood: http://parentsguidecordblood.org/
Save the Cord Foundation: http://www.savethecordfoundation.org/

X. SUBJECT'S STATEMENT OF CONSENT

I have read both pages of this consent form and I have been given the opportunity to ask questions. I voluntarily agree to donate my baby's cord blood for medical research studies, if it cannot be used for transplantation, as defined in this consent form.

Donor Advocacy information was provided to r	ne by Cord for Life [®] .	
Mother's Signature	Date	
		

FOR COMPLETION BY CORD FOR LIFE, INC. REPRESENTATIVE Certification of Counseling Healthcare Professional I certify that the nature and purpose, the potential benefits, and possible risks associated with donating umbilical cord blood for research have been explained to the above individual and that any questions about this information have been answered. Counseling Healthcare Professional Date Use of an Interpreter: Complete if the subject is not fluent in English and an interpreter was used to obtain consent. Print name of interpreter: ______ Date: ______ Signature of interpreter: ______ Date: ______ An oral translation of this document was administered to the subject in ______ (state language) by an individual proficient in English and

Cord for Life[®], Inc. (CFL)

(state language). See the attached short form addendum for documentation.