

# Cord For Life® Cord Blood Donation Information



Date Issued: 04FFB2020

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 <a href="https://www.cordforlife.com">www.cordforlife.com</a>. 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 34<sup>th</sup> 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</u> <u>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.

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

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 <a href="mailto:physician must sign">physician must sign</a> 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 your documentation, on receipt. You will be contacted, by phone, 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 when in active labor and after the birth of your baby, day or night. It is your responsibility to ensure that we are notified within two hours after the collection of the cord blood. We will dispatch a courier to pick up the cord blood within the next business day.

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

Form Number: B.1-6 Rev. K



## **DONOR INFORMATION AND HEALTH HISTORY**

						•		
MOTHER'S <u>LAST</u> NAME <u>FIRST</u> NAME					M.I.	LAST 4 S	S# DIGITS	
BEST CONTACT PHONE:	EMAIL				•	MOTHER	'S DOB:	
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	
BABY'S DUE DATE:								
BABT O BOL BATE.								
DELIVERY PHYSICIAN'S NAME				PHONE				
CLINIC NAME								
DELIVERY HOSPITAL NAME				PHONE				
HOSPITAL ADDRESS			CITY			STATI	ZIP CODE	
BABY'S Since certain HLA Types may be more common			HNICITY INFORI the information below		electing a	cord bloc	d unit for transplant.	
Baby's Ethnicity: Response is required,	plea	se check o	ne. 🗆 Hispanic	or Latino		Not H	spanic or Latino	
Baby's Race: Response is required. O	f whice	ch group(s)	is your baby a me	ember? (Se	lect all	that app	oly.)	
American Indian or Alaska Native	Black	or African A	American	A	sian			
Alaska Native or Aleut (ALANAM)	А	frican (AFB)			С	hinese (N	CHI)	1
North American Indian (AMIND)	Α	frican American	(AAFA)		Fi	lipino (Phi	lipino) (FILI)	İ
American Indian South or Central		lack Caribbean	, ,			ipanese (J	,	
American (AMIND) Caribbean Indian (AMIND)			entral American			orean (KO	,	
Canadan (Canada	(3	SCAMB)					(SCSEAI)	
							(SCSEAI) east Asian (SCSEAI)	
						iller South	east Asiati (SCSEAI)	ł
Native Hawaiian or Other Pacific Islander	White	)						
Guamanian (OPI)	E	astern Europear	n (CAU)				ropean (CAU)	
Hawaiian (HAWI)	N	lediterranean (C	CAU)				ropean (CAU)	
Samoan (OPI)		liddle Eastern (N	,				bean (CAU) or Central American	
Other Pacific Islander (OPI)		orth Coast of Af				AU)	TOT Certifal American	
	N	orth American (	JAU)			ther White	(CAU)	
							(0, 10)	

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.

	LAKAHAN DIBINAN DIBINA '		8/4
MANAGE AND	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		·-
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□	Ν□
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□	Ν□
	c.   Rabies vaccination in the past 12 months.		
4	In the past 8 weeks, have you had any shots or vaccinations?  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□	Ν□
6	In the past 4 months, have you experienced <b>TWO (2)</b> 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? <b>If yes</b> , which symptoms and when?  Details:	Υ□	Ν□
7	Have you ever had any type of cancer, including leukemia?  If yes, details:	Υ□	Ν□
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□	Ν□
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□	Ν□
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□	Ν□
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 <b>animal</b> ? 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 <b>animal</b> ? <b>If yes,</b> 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□	Ν□
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□	Ν□
	21. If yes, were shared or non-sterile instruments, needles, or inks used for the tattoo or piercing?	Υ□	N□



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



MOTHER'S LAST 4 SS# DIGITS: \_\_\_\_\_ MOTHER'S DOB: \_\_\_\_\_

	FOR USE \	VITH QUESTIONS #39 – 42 – COUNTRIES DEFINED AS EUROPE		
	ILBANIATravelResident Date(s):	GREECE        Travel        Resident         ROMANIA        Travel           Date(s):        Travel        Travel	el	Resident
	oate(s). Fotal Time:	Total Time:  Total Time:		
	AUSTRIATravelResident	HUNGARYTravelResidentTravelResidentTravelTravelTravel	el	_Resident
	Date(s): Fotal Time:	Date(s): Total Time:  Date(s): Total Time:		
	BELGIUMTravelResident Date(s):	IRELAND (REPUBLIC OF)	'el	_Resident
	oate(s): Fotal Time:	Total Time:  Total Time:  Total Time:		
	BOSNIA-HERZEGOVINATravelResident	Travel Resident SPAIN Travel Resident Resident Travel Resident Res	el	_Resident
	Date(s): Fotal Time:	Date(s): Total Time:  Date(s): Total Time:		
	BULGARIATravelResident	LIECHTENSTEIN Travel Resident SWEDEN Travel	el	_Resident
	Date(s): Fotal Time:	Date(s): Total Time:  Date(s): Total Time:		
	CROATIA TravelResident	LUXEMBOURG Travel Resident SWITZERLAND Travel	el	_Resident
	Date(s): Fotal Time:	Date(s): Total Time:  Date(s): Total Time:		
	CZECH REPUBLIC Travel Resident	MACEDONIA Travel Resident UNITED KINGDOM (UK) includes England, N		
	Date(s): Fotal Time:	Date(s): Scotland, Wales, Isle of Man, Channel Islan Total Time: Falkland Islands		
	DENMARKTravelResident	NETHERLANDS (HOLLAND)TravelResidentTotal Time		
	Date(s): Fotal Time:	Date(s): Total Time:		
	FINLAND Travel Resident	NORWAY Travel Resident YUGOSLAVIA (FEDERAL REPBULIC OF) Tra	ual .	Resident
	Date(s):	Date(s):	VEI	_vezinelir
	Total Time:	Total Time: Total Time:		
	FRANCETravelResident Date(s):	POLANDTravelResident KOSOVO, MONTENEGRO, SERBIATrav Date(s): Date(s):	/el	_Resident
	Total Time:	Total Time: Total Time:		
	GERMANYTravelResident Date(s):	PORTUGALTravelResident Date(s):		
	Total Time:	Total Time:		
38		raveled to Europe? ( <i>refer to chart above</i> ) <i>If no</i> , skip to question <b>42</b> . check in all the appropriate box(es) above to identify the country(ies), reason,	Υ□	Ν□
	39. From 1980 through 1996, did you chart above)?	u spend time that adds up to 3 months or more in the United Kingdom (refer to	Υ□	N□
	40. Since 1980, have you received a	transfusion of blood or blood components while in the UK or France?	Υ□	N□
	<b>41</b> . <b>Since 1980</b> , have you spent time spent in the UK between 1980 and 1980.	that <u>adds up to 5 years or more</u> in Europe ( <i>refer to chart above</i> ), including time 96?	Υ□	Ν□
42	From 1980 through 1996, were you a member of the U.S. military?	member of the U.S. military, a civilian military employee, or a dependent of a	Υ□	N□
43	From 1980 through 1990, did you spe following countries: United Kingdom, E	end a <u>total of 6 months or more</u> associated with a military base in any of the elgium, Netherlands or Germany?	Υ□	N□
44	From 1980 through 1996, did you spe following countries: Spain, Portugal, Tu	end a <u>total of 6 months or more</u> associated with a military base in any of the urkey, Italy or Greece?	Υ□	N□
		<u> </u>		



MOTHER'S LAST 4 SS# DIGITS: \_\_\_\_\_ MOTHER'S DOB: \_\_\_\_\_

	FOR USE WITH QUESTIONS 46-48: AFRICAN COUN	ITRIES		
	BENIN    Travel    Resident       Date(s):    Travel    Resident       Total Time:    Travel    Resident       Date(s):    Travel    Resident       Total Time:    Travel    Resident	SENEGALTr Date(s): Total Time:	avel	_Resident
Į.	CAMERODN    Travel    Resident       Date(s):    Travel    Resident       Total Time:    Travel    Resident       Date(s):    Travel    Resident       Total Time:    Travel    Resident	TOGOTotal Time:	ravel	Resident
l	CENTRAL AFRICAN REPUBLIC     Travel     Resident       Date(s):     Total Time:         Total Time:         KENYA    Travel    Resident       Date(s):    Travel    Resident       Total Time:	ZAMBIA Date(s): Total Time:	ravel	Resident
	CHAD     Travel     Resident       Date(s):     Date(s):       Total Time:     Total Time:			
	CONGO     Travel     Resident       Date(s):     Date(s):       Total Time:     Total Time:			
45	Since 1977, were you born in, have you lived in, or have you traveled to any African canswer question 46. <i>If no</i> , skip to question 47.  a) Use the chart above and place a check in all the appropriate box(es) above to ide date(s) and total time that apply.		Υ□	Ν□
	<b>46.</b> While in one of the African countries listed above, did you receive a blood transf treatment with a product made from blood?	usion or any other medical	Υ□	N□
47	Have you had sexual contact with anyone who was born in or lived in any African coun	try listed above since 1977?	Υ□	N□
40	a.) Were you and/or the baby's father adopted at early childhood?		Υ□	N□
48	b.) If yes, is a family medical history available for you and/or the baby's father?		Υ□	N□
49	Are you and the baby's father related, except by marriage? (e.g., first cousins)		Υ□	N□
	Did this pregnancy use either a donor egg or donor sperm?		Υ□	N□
50	If yes, is a family medical history questionnaire available for the egg or sperm donor? (ple Name of the Clinic:	ase attach copy)	Υ□	N□
	Have you ever had an abnormal result from a prenatal test (e.g., amniocentesis, blood teanswer the following questions. If no, skip to question 52.	st, and ultrasound)? If yes,	Υ□	N 🗆
51	A) Which test was abnormal?			
	B) What was the abnormal test result?			
	C) Was a diagnosis made? Specify diagnosis:			
52			Υ□	N□
	If yes, what was the cause?			
53			Υ□	N□
	If yes, what was the cause?			l –
54	(,,,,	•	Υ□	N 🗆
55		t during your pregnancy?	Υ□	Ν□
56	<ul> <li>Have you had sex during your pregnancy with a male or female who is known to have:</li> <li>a) A medical diagnosis of ZIKV (Zika) within the six months prior to that contact.</li> <li>b) Resided in, or traveled to, an area with active ZIKV (Zika) transmission within the six</li> </ul>	months prior to that contact	Υ□	N 🗆
	Sylvestace in, or navoice to, an area with active Zirve (Zirva) transmission within the six	monard prior to triat contact.	<u> </u>	<u> </u>



П	FAMILY MEDICAL HISTORY													
П	For the following questions please use the following codes to describe the relationship between the baby and a family member with a disease:  Family Relationship Codes: BM Baby's Mother BGP Baby's Grandparent BMS Baby's Mother Sibling													
П	BF Baby's Father BS Baby's sibling BFS Baby's Father's Sibling (Parents' sibling (BMS and BFS) refer to the baby's aunts and uncles by blood and does <u>not</u> include aunts and uncles who are in-laws of the parents.)													
₹		Cancer or Leukemia?	Y 🗆	N □		ia acca <u>iibt</u> iiiciat			S WIID BI	<b>5</b> III 14475		, par circ	.,	
<u>'</u>	57	If yes, please specify all th			aceed to next guestion		ВМ	ВІ	- P	s				
ı'	31				oceeu to next question.	•	BIVI			_				
4		<ul><li>A) Brain or other nervous :</li><li>B) Bone or joint cancer</li></ul>	System Ca	ancei			$+ \frac{\sqcup}{\Box}$							
		C) Kidney (including renal	nelvic) c	ancer										
		D) Thyroid Cancer	pervio) o	11001										
	E) Hodgkin's Lymphoma									_	IMI	MEDI <i>A</i>	ATE	
		F) Non-Hodgkin's Lympho	ma									AMIL		
		G) Acute or chronic myelo		nyeloid le	ukemia							ONLY		
	•	H) Acute or chronic lymph		•						J				
		I) Skin Cancer		•										
		J) Other cancer/leukemia												
		Specify Type:												
1		Specify Type:								·				
ŀ		Red Blood Cell	ΥΠ	N 🗆							_			
1		If yes, please specify all th			oceed to next guestion.		ВМ	BF	В	S B	GP	BMS	BFS	5
		A) Diamond-Blackfan Syn		, p.										
	58	B) Elliptocytosis								] [				
		C) G6PD or other red cell enzyme deficiency								] [				_
-1	•	D) Spherocytosis	-						Г	] [	]			
Ī		White Blood Cell Disea	ase?	<b>′</b> 🗆	N 🗆									
1		If yes, please specify all th	at apply.	<i>If no</i> , pro	ceed to next question.		BM	BF	В	S B	GP	BMS	BFS	\$
	59	A) Chronic Granulomatous	s Disease	)						] [				
ľ	פפ	<b>B)</b> . Kostmann Syndrome.								] [				
		C) Schwachman-Diamond	l Syndron	ne						] [				
L		D) Leukocyte Adhesion De	eficiency	(LAD)						] [				
Γ		Immune Deficiencies?	١	<b>′</b> 🗆	N 🗆									
1		If yes, please specify all th	at apply.	<i>If no</i> , pro	oceed to next question.		ВМ	BF	В	S B	GP	BMS	BFS	5
	A)	ADA or PNP Deficiency												
	B) In	Combined Immunodeficier Combined Immunodeficiency Disease (C		rome (CIE	), Common Variable									
60	C	DiGeorge Syndrome												
	D)	Hereditary Hemophagocyt	ic Lymph	ohistiocyt	osis (HLH) including FEL									
	E)	Hypoglobulinemia												
	F)	Nezeloff Syndrome												
	G	Severe Combined Immuno	odeficiend	су										
	H)	Wiskott-Aldrich Syndrome												



MOTHER'S LAST 4 SS# DIGITS: \_\_\_\_\_ MOTHER'S DOB: \_\_\_\_

	Platelet Disease?	Υ		N								
	If yes, please specify all that apply. If no, proceed to next quest	ion				вм	BF	BS	BGP	BMS	BFS	
	A) Amegakaryocytic Thrombocytopenia											
	B) Glanzmann Thrombasthenia											
61	C) Hereditary Thrombocytopenia											
	D) Platelet Storage Pool Disease											
	E) Thrombocytopenia with absent radii (TAR)											
	F) Ataxia-Telangiectasia											
<u> </u>	G) Fanconi Anemia	1										
62	Other blood disorder or problem	Y	′ 🗆	N								
Hem	oglobin Problems					ВМ	BF	BS	BGP	BMS	BFS	
63	Sickle cell disease, such as sickle-cell anemia or sickle thalassemi	ia?	Υ□	N								
63	Specify disease:											
64	Thalassemia, such as alpha thalassemia or beta-thalassemia?		Υ□	_								
	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)											
65	H) I-cell disease											
	I) 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											
1	P) Porphyria											
	<b>Q)</b> Other or unknown metabolic/storage disease, Details:											
	Acquired Immune System Disorders					ВМ	BF	BS				
66	HIV/AIDS?		<u> </u>	_								
	Severe autoimmune disorder?  If yes, please specify all that apply including providing the other re		<b>Y</b> □ ted	I N								
	information. <i>If no, proceed to next question.</i>	quos	icu			ВМ	BF	BS				
	A) Crohn's Disease or Ulcerative Colitis								IMMEDIATE FAMILY			
	B) Lupus								IMME	DIATE I		
67	C) Multiple Sclerosis (MS)								ONLI			
	D) Rheumatoid Arthritis											
	Diagnosis Date:											
	Currently under MD care? Describe:							_				
	Are you currently taking any medication? Name:											



MO	THER'S LAST 4 SS# DIGITS: MO	THER'S	DOB:						
68	Any diagnosis of other or unknown immune system disorder?	Υ□	N□	ВМ	BF	BS	IMME	DIATE F	AMILY
	Specify Disorder:		_					ONLY	
				ВМ	BF	BS	BGP	BMS	BFS
69	Required Chronic Blood Transfusions?	Υ□	N□						
70	Been told you or your family member(s) have hemolytic anemia?	Υ□	N□						
71	Had spleen removed to treat a blood disorder?	Υ□	N 🗆						
72	Had gallbladder removed before the age of 30?	Υ□	N 🗆						
73	Had Creutzfeldt-Jakob disease (CJD)?	Υ□	N□						
	Other serious or life-threatening diseases affecting the family?	Υ□	N 🗆	вм	BF	BS	BGP	BMS	BFS
	If yes, list affected family member(s) and type of disease								
74	Specify Type:								
	Specify Type:								
	Specify Type:								
		1	1						
75	In answering these questions, have you answered for both your family and the baby's father's family?	Υ□	N 🗆						
Δdde	ndum A: STATE OF NEW YORK-ONLY For collections within the State of I	NV the fo	llowing a	uestions i	must he s	nswered			
1.	Any history of acute respiratory disease? <i>If Yes</i> , please describe	141, 1110 10	nowing q	uc3(10113 1		ii i swere a		Υ□	N 🗆
2.	Any active tuberculosis disease or history of tuberculosis therapy? <i>If Yes</i> , please  Y   N						N 🗆		
3.									
Add	endum B: COVID-19 or Severe Acute Resp	iratory	Syndro	ome (SA	ARS)				
Add	endum B: COVID-19 or Severe Acute Resp	oiratory	Syndro	ome (S <i>A</i>	ARS)				
Adde	In the past 14 days, have you had any symptoms of Covid-19 or SA fatigue, muscle or body aches, headaches, new loss of taste or sme nausea or vomiting, or diarrhea?	ARS such	n as feve	r, cough	, shortne		eath,	Υ□	N 🗆
	In the past 14 days, have you had any symptoms of Covid-19 or SA fatigue, muscle or body aches, headaches, new loss of taste or sme	ARS such	n as feve nroat, col	er, cough	, shortne or runny	nose,	eath,		
1.	In the past 14 days, have you had any symptoms of Covid-19 or SA fatigue, muscle or body aches, headaches, new loss of taste or sme nausea or vomiting, or diarrhea?  In the past 14 days, have you been suspected of having or diagnos	ARS such	n as feve nroat, con Covid-19	er, cough ngestion or SAR	, shortne or runny S, by a n	nose, nedical		Υ□	N 🗆
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Form Number: B.1-1 Rev. W



## 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 <u>not</u> 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)	
I have read the above information and have had test. I allow the test results to be made availabl <b>Dr</b>	d my questions about the HIV test answered. I agree to take the HIV e 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 above	e):
PHYSICIAN'S ORDER FOR BLOOD TESTING	(REQUIRED)
<b>Rx</b> Patient Nam	ne:
It is an FDA requirement that Cord For	Life performs maternal blood testing. Tubes will be included with the hospital/birthing center during labor and delivery.
HIV-1 and HIV-2 (antibody to the AIDS virus) HEPATITIS B (HBsAg & HBcAb) HTLV-I and HTLV-II CMV, WNV	HCV/HIV NAT (Hepatitis C and AIDS virus by Nucleic Acid Test) HEPATITIS C VIRUS (Anti-HCV) SYPHILIS, ABO Rh CHAGAS DISEASE
Printed Name of Physician or Midwife:	Date:

**IMPORTANT:** 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 <a href="www.cordforlife.com">www.cordforlife.com</a> (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<sup>®</sup>. To withdraw your unit, forward a notarized letter to the Cord for Life<sup>®</sup> 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 <a href="mailto:dhudspeth@cordforlife.com">dhudspeth@cordforlife.com</a> or Deborah Sardone (*Director of Quality Assurance*) at <a href="mailto:dsardone@cordforlife.com">dsardone@cordforlife.com</a>.

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: <a href="http://parentsguidecordblood.org/">http://parentsguidecordblood.org/</a>
Save the Cord Foundation: <a href="http://www.savethecordfoundation.org/">http://www.savethecordfoundation.org/</a>

#### 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 me by Cord for Life®.

Mother's Signature	Date	

Mother's Printed Name		
	LIFE, INC. REPRESENTATIVE  enefits, and possible risks associated with donating umbilical cord blood dual and that any questions about this information have been answered.	-
Counseling Healthcare Professional	 Date	_
<b>Use of an Interpreter:</b> Complete if the subject is not f	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 administere (state language) by an individual proficient in English	ed to the subject in and	

Cord for Life<sup>®</sup>, Inc. (CFL)

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