

CONSENT TO THAW AND TRANSFER CRYOPRESERVED EMBRYOS

I,	(Female Partner) and	(Partner, Spouse)
wish to use or understand th menstrual eyo uterus. I/We embryo transi	ryopreserved (frozen) embryos to achieve pregnancy by undergoing a frozen embryo that achieving pregnancy using cryopreserved embryos requires the preparation of the usele), followed by the thawing of embryos and embryo transfer into a recipient's (Feman acknowledge that embryo survival after thawing cannot be guaranteed, nor can pregnate if embryos survive the thawing process. I/We consent to allow the California Cent affiliate laboratory, and physicians to thaw our cryopreserved embryos for the purpose	atterus (in a medicated or natural alle Partner or gestational surrogate) ancy be guaranteed following arer for Reproductive Health
embryos and intramuscular of vaginal sup lining of the u medicated FE medications u aspirin tablet. effects are rar	and that in a FET cycle the uterus must be prepared for implantation for several weeks embryo transfer. In a medicated FET cycle this is normally first achieved using estroger injections, or transdermal patches. Several days prior to embryo thawing and transfer expositories or intramuscular injections. The purpose of estrogen treatment is to thicken atterus) and make it receptive to embryos, while progesterone serves to prepare the uter atterus, both estrogen and progesterone administration continues through the 9th to 10 used in a FET cycle may include medications to suppress ovulation, low dose oral cort. Estrogen and progesterone treatment may cause mild mood changes, bloating, breast the after treatment with steroids but low dose oral steroids can cause acne, blurred vision mach upset, bloating, and irritable mood. Monitoring for a FET cycle, will include from	gen in the form of oral tablets, or progesterone is added in the form in the endometrial lining (the inside rus for embryo implantation. In a 0 weeks of gestation. Additional ticosteroids, along with a low dose t tenderness, and fatigue. Side on, sleep disturbance, muscle
weakened usi improve the c embryos undo increased incre	ching hing (AH) is a specialized laboratory procedure whereby the shell surrounding the eming micromanipulation instruments in order to facilitate the hatching (release from its schance of implantation in the uterus. Multiple studies have suggested improved implantation assisted hatching prior to embryo transfer. The risks associated with assisted hidence of multiple gestations, and the possible damage/destruction of the embryos. A set be signed before thawing of embryos and embryo transfer.	shell) of the embryo(s), so as to ntation rates of frozen-thawed hatching include the potential of
My initials in	dicate that I consent to assisted hatching of frozen-thawed embryos:	Initials
	nsfer and that there is no guarantee than any embryo will survive the thawing process and the ansfer will be canceled.	nat if no embryos survive the thaw,
cervix (openi zygote intrafa discomfort, a	ransfer procedure involves the placement of a catheter containing thawed embryo(s) in g to the womb). Occasionally frozen-thawed embryos may be transferred into the fallopian transfer (ZIFT), or tubal embryo transfer (TET). Trans-cervical embryo transfer d on occasion scant vaginal bleeding. A separate consent for Embryo Transfer must l embryo transfer.	llopian tubes in a procedure called fer may lead to minimal to no
My initials in	dicate that I consent to embryo transfer of frozen-thawed embryos:	Initials
I/We understa	and that there is no guarantee that any of the frozen-thawed embryos transferred will re	esult in a pregnancy. I/We

understand that as in any assisted reproductive technique (ART) treatment, the transfer of a single or multiple embryo(s) into the uterus in a FET cycle may result in a risk of multiple gestation (more than one baby). The risks of multiple gestations include, but are not limited to, preterm labor and the delivery of premature infants who may require prolonged hospitalization and who may have long-term complications associated with prematurity. It is CCRH's policy to limit the number of embryos transferred according to maternal age and embryo quality in order to maximize success rates and minimize the risk of a multiple gestation.

I/We understand that pregnancies resulting from FET are subject to the same complications as pregnancies achieved with standard in vitro fertilization (IVF)/embryo transfer and those achieved without medical intervention, such as miscarriage, ectopic pregnancy, preterm labor, or other complications. There may be a risk of infants having developmental problems or congenital birth defect as a result of any ART treatment, including embryo cryopreservation and thawing/transfer; however, initial human experience and extensive experience in domestic animal species have not yet demonstrated an increase in developmental or congenital anomalies in



offspring born following cryopreservation beyond that observed in other ART treatments (such as IVF and embryo transfer). I/We understand that the health of any infant resulting from this procedure cannot be guaranteed. Separate consents for Assisted Reproductive Techniques and Embryo Cryopreservation must be signed before the IVF and embryo freezing procedures.

RELEASE			Initials	
I/We agree to absolve, agents and employees, remote, resulting from embryos, the birth of a	from any and al thawing of froze physically or m	ify, protect and hold harmless CCRH, its affiliate I liability, claims or damages including legal fees on embryos and frozen embryo transfer including entally disabled child or subsequent disputes between as a result of this procedure.	, arising from any adverse outcome, how but not limited to the loss or destruction	vever 1 of
			Initials	
	ndicates that you	or Thawing and Transfer of Cryopreserved En a have read the preceding consent, that you have your satisfaction.		d that
PATIENT NAME	(print)	PATIENT SIGNATURE	DATE	
PARTNER NAME	(print)	PARTNER SIGNATURE	DATE	
WITNESS	(print)	WITNESS SIGNATURE	DATE	