



PRENATAL MICROARRAY REQUISITION

GENETICS LABORATORY

2200 Eglinton Ave. W., Rm 2H144 Tel: (905) 813-1100 x6288
 Mississauga, ON L5M 2N1 Canada Fax: (905) 813-3854

Acct # _____
 Last Name: _____
 First Name: _____
 DOB: _____ (DD/MM/YYYY) Sex: _____
 Healthcard #: _____ VC: _____
 Address: _____

 Primary Phone #: _____
 Unit # _____

To ensure there is no delay in reporting results; please ensure the following accompany each sample:

- Completed requisition form
- Parental samples. *Mother's sample is mandatory for MCC studies. Father's sample is highly recommended (Complete a Separate Prenatal Microarray Requisition for each parent)*

PHYSICIAN INFORMATION	
Referring Dr: _____ Registration #: _____ Address: _____ Telephone: _____ Fax: _____ Signature (required): _____	Copy To: _____ Registration #: _____ Address: _____ Telephone: _____ Fax: _____
CLINICAL INFORMATION	SPECIMEN TYPE
Gestation at collection date: _____ weeks Karyotype result (if known) : _____ Sex of Fetus (if known): _____ (by <input type="checkbox"/> QF-PCR or <input type="checkbox"/> FISH) Father's name: _____ DOB: _____ <i>Please attach pedigree if relevant</i>	<input type="checkbox"/> Direct CVS (10-15 mg cleaned villi) <input type="checkbox"/> Cultured CVS (1 T75 or 2 T25 flasks 70% confluent) <input type="checkbox"/> Direct Amniotic Fluid (15cc) <input type="checkbox"/> Cultured Amniocytes (1 T75 or 2 T25 flasks 70% confluent) <input type="checkbox"/> DNA: 2ug total (at a minimum concentration of 70 ng/uL) <input type="checkbox"/> Other: _____
INDICATION FOR TESTING:	
<input type="checkbox"/> Fetal Ultrasound Abnormality <i>(please specify below and attach copy of fetal ultrasound)</i> <input type="checkbox"/> Brain malformation: _____ <input type="checkbox"/> Facial cleft: _____ <input type="checkbox"/> Heart defect: _____ <input type="checkbox"/> Lung abnormality: _____ <input type="checkbox"/> Diaphragmatic hernia: _____ <input type="checkbox"/> GI abnormality: _____ <input type="checkbox"/> GU abnormality: _____ <input type="checkbox"/> Skeletal abnormality: _____ <input type="checkbox"/> NT greater than or equal to 3.5mm: _____ <input type="checkbox"/> Other: _____ <input type="checkbox"/> Known familial microarray deletion/duplication syndrome <i>(provide copy of the result)</i>	<i>Parental Sample:</i> <input type="checkbox"/> Maternal Blood: 5-10mL EDTA <input type="checkbox"/> Paternal Blood: 5-10mL EDTA
SPECIMEN COLLECTION:	
Date: _____ (DD/MM/YYYY) Time: _____	
CVH Lab Use Only	
Date Rec'd:(DD/MM/YYYY) _____ Time: _____ # Tubes Rec'd: _____ Comments: _____	
Unit #	LAB #

3993 D HR (July/2013)

