

PRENATAL GENETICS LABORATORY REQUISITION

BC Children's Hospital and BC Women's Hospital Facility Code L1050

4500 Oak Street, Vancouver B.C. V6H 3N1

Cytogenetics Tel: 604-875-2304, Fax: 604-875-3601

Molecular Genetics Tel: 604-875-2852, Fax: 604-875-2707

www.genebc.ca

Ordering Physician		
Last Name	First Name	Billing #
Address		
Phone	Fax	
Contact Person / Genetic Counsellor	Phone	
Copy Physician	Billing #	
Copy Physician	Billing #	

Patient Information		
Last Name	First and Middle Names	
Gender	Date of Birth (DD/MMM/YY)	
X Female		
Personal Health Number (PHN)	Referring Hospital ID	Referring Clinic ID
Address		
Patient Phone Number	Multiple Gestation – select only ONE:	
	<input type="checkbox"/> Fetus A <input type="checkbox"/> Fetus B <input type="checkbox"/> Fetus C	

GENETICS LABORATORY
LABELS ONLY

Reproductive History G: _____ T: _____ P: _____ A: _____ L: _____	Procedure Date (DD/MMM/YY)	Current Gestational Age _____ weeks _____ days	<input type="checkbox"/> EDD by U/S <input type="checkbox"/> LMP (DD/MMM/YY)
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Reason For Referral REQUIRED	
<i>Choose all that apply; analysis will not be performed unless appropriate history is provided.</i>	
<input type="checkbox"/> 40+ years <input type="checkbox"/> 35 – 39 years, no prior screening, singleton, ≥ 21 wks gestation (at first prenatal visit) <input type="checkbox"/> ≥35 years, no prior screening, multiple gestation <input type="checkbox"/> Positive maternal screen: <input type="checkbox"/> IPS <input type="checkbox"/> SIPS <input type="checkbox"/> Quad <input type="checkbox"/> NIPT <input type="checkbox"/> Other: _____ <input type="checkbox"/> In vitro fertilization with intracytoplasmic sperm injection (IVF with ICSI)	Age at projected birth: _____ Positive for: <input type="checkbox"/> T21 <input type="checkbox"/> T18 <input type="checkbox"/> Other: _____ Risk: _____

<i>Additional information must be provided for the following:</i> <input type="checkbox"/> Fetal ultrasound abnormality <input type="checkbox"/> Fetus at risk of inherited disease <input type="checkbox"/> Personal/family history of chromosome abnormality <input type="checkbox"/> Follow up after CVS <input type="checkbox"/> Culture for: <input type="checkbox"/> Biochemical Genetics <input type="checkbox"/> Possible future testing (please specify) <input type="checkbox"/> Supernatant fluid to: <input type="checkbox"/> Biochemical Genetics <input type="checkbox"/> Clinical Biochemistry	<i>Relevant clinical and family history:</i>
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Test(s) Requested	Sample Type <i>Use this requisition for prenatal samples ONLY</i>
<input type="checkbox"/> QF-PCR Rapid Aneuploidy Detection (RAD) <input type="checkbox"/> Chromosome microarray (CMA) <input type="checkbox"/> Molecular test: _____ <i>Specify test & FAX req to MGL (2707) at time of scheduling</i> <input type="checkbox"/> Other: _____ <i>Specify (ex. Karyotype, FISH probe)</i>	<input type="checkbox"/> 25 mL Amniotic fluid <input type="checkbox"/> 2 mL Fetal Blood <i>(Kleihauer required for cordocentesis)</i> <input type="checkbox"/> 20 mg CVS <input type="checkbox"/> Other: _____ Path # _____
Collection Details	
Date Collected (DD/MMM/YY)	Time Collected (HH:MM)

Ordering Physician Signature REQUIRED	Date
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Date Arrived: _____ Previous Studies: _____

GENETICS LABORATORY USE ONLY