

**Lab Requisition**  
**CAMD**

Name  
MRN  
DOB  
M/F

**Constitutional Cytogenetics**

Location/Institution	Reserved For CAMD Sticker
ICD Code(s) <b>REQUIRED</b> <i>(ICD-10-CM codes required as of 10/1/15.)</i>	

Collection Information			
Date	Time	Drawn by:	
		Phleb. ID	MD/RN ID

Ordering Clinician: Please print First, Last name	Clinical ID/NPI#	Contact Name & Phone Number
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Clinician Signature: (Required)	Clinician's Fax Number for Patient Reports:	Clinician's Phone Number:
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Send Duplicate Reports To: (Name/Address/Fax#/Phone)

**SPECIMEN SUBMITTED:**

<input type="checkbox"/> Amniotic Fluid	<input type="checkbox"/> Chorionic villus	<input type="checkbox"/> Peripheral Blood
<input type="checkbox"/> PUBS	<input type="checkbox"/> Cord Blood	<input type="checkbox"/> POC
<input type="checkbox"/> Tissue: Indicate type _____	<input type="checkbox"/> BWH Pathology Accession/ Block # _____	<input type="checkbox"/> Other: _____

**Clinical History:**

Tests Requested	Indication for Testing	Pregnancy History	
SNP Microarray	<i>Please provide additional details under Clinical History:</i> <input type="checkbox"/> Short stature <input type="checkbox"/> Multiple congenital anomalies <input type="checkbox"/> Developmental delay <input type="checkbox"/> Infertility <input type="checkbox"/> Azoospermia/oligospermia <input type="checkbox"/> Premature ovarian failure <input type="checkbox"/> History of recurrent SAB <input type="checkbox"/> Abnormal cffDNA result <i>List abnormal chromosome(s):</i>  <input type="checkbox"/> Abnormal maternal screen <input type="checkbox"/> Increased risk of trisomy <input type="checkbox"/> Increased risk of NTD <input type="checkbox"/> Abnormal ultrasound <i>List findings under clinical history</i> <input type="checkbox"/> Advanced maternal age <input type="checkbox"/> Family history chrom. abnormality <input type="checkbox"/> Other:	G _____ P _____ SAB _____ TAB _____	
Parental follow up to a SNP microarray (additional charges apply)			Gestational age:
Standard chromosome analysis <b>R</b>			Does Patient wish to know fetal sex? <input type="checkbox"/> Yes <input type="checkbox"/> No
FISH (not performed routinely with SNP arrays) Probe(s)/Chromosome(s) of Focus:			Multiple Gestation <input type="checkbox"/> Yes <input type="checkbox"/> No
AFP (testing performed at BWH Reproductive Endocrinology)			If yes, please indicate number: _____
ACHE (testing performed at FBR)			
Save unspun amniotic fluid for CMV/Toxoplasma testing			
Save cells			
Send out DIRECT specimen (indicate sample type and quantity):			
Send out CULTURED specimen (indicate sample type and quantity):			
Cryopreservation of cells – stored for 6 months only (Requires advance approval by Laboratory)	Reference Laboratory (name and telephone #):		

**By submission of this sample and request for genetic testing, I hereby warrant that the appropriate prior written consent has been obtained from the patient or authorized representative.**

**Provider signature:** \_\_\_\_\_ **Date:** \_\_\_\_/\_\_\_\_/\_\_\_\_

**R** Reflex or confirmatory testing, if required, will be performed, reported and billed unless indicated here:  No reflex tests