



Medical Genetics Laboratory Constitutional Cytogenetics Requisition

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COMPLETE ALL SECTIONS. Incomplete requisitions will delay testing.

Patient Information:	Referring Physician:
Name: _____ Last First	Name: _____ (Print Clearly)
HCN: _____	Address: _____
Date of Birth: _____ DD/MONTH/YYYY	City: _____ Province: _____
Sex: <input type="checkbox"/> Male <input type="checkbox"/> Female	Telephone: _____ Fax: _____
Previously tested family members? <input type="checkbox"/> Yes <input type="checkbox"/> No	Signature: _____ Date: _____ DD/MONTH/YYYY
Index Case: _____	Copy to: _____

I. Specimen Requirements:	
Peripheral Blood collected at room temperature in sodium heparin dark green top tube	Tissue in sterile RPMI/Hanks/Saline (minimum 3 X 5 mm)
<input type="checkbox"/> Adults - Children - 4-8 mL	Specimen Collection:
<input type="checkbox"/> Infants - 2-4 mL	Date: _____ Time: _____
	Location: _____ Initials: _____
	<input type="checkbox"/> Skin <input type="checkbox"/> Cartilage <input type="checkbox"/> Placenta <input type="checkbox"/> POC

II. Testing Requested - REQUIRED (check all that apply)

Karyotype (Include indication for testing below)

Indication for Testing:	FISH:
<input type="checkbox"/> Down Syndrome <input type="checkbox"/> Klinefelter Syndrome	<input type="checkbox"/> X centromere/SRY (Yp11.3)
<input type="checkbox"/> Trisomy 13 <input type="checkbox"/> Turner Syndrome	<input type="checkbox"/> Digeorge/VCFS Syndrome (22q11.2)
<input type="checkbox"/> Trisomy 18	<input type="checkbox"/> Smith-Magenis Syndrome (17p11.2)
<input type="checkbox"/> Greater than 2 miscarriages OR <input type="checkbox"/> Stillbirth <i>(collect blood at least 8 weeks post birth)</i>	<input type="checkbox"/> Wolf-Hirschhorn Syndrome (4p16)
<input type="checkbox"/> Infertility (details: _____)	<input type="checkbox"/> Williams Syndrome (7q11.23)
<input type="checkbox"/> Amenorrhea	<input type="checkbox"/> Prader-Willi/Angelman Syndrome (15q11.2)
<input type="checkbox"/> Premature Menopause	<input type="checkbox"/> post-BMT Monitoring by XX/XY FISH
<input type="checkbox"/> IVF Treatment Pending	<input type="checkbox"/> Other: _____
<input type="checkbox"/> Azoospermia/Oligospermia	
<input type="checkbox"/> Family history of chromosome rearrangement (details: _____)	Microarray Follow up FISH:
<input type="checkbox"/> Prenatal detection of chromosome rearrangement (details: _____)	<input type="checkbox"/> Proband <input type="checkbox"/> Family Member
Patient currently pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No	Relationship: _____
<input type="checkbox"/> Multiple Congenital Malformations	Copy number change: _____
<input type="checkbox"/> Dysmorphic Facies	Lab number: _____
<input type="checkbox"/> Ambiguous Genitalia	
<input type="checkbox"/> Intellectual Disability/Developmental Delay	FOR LABORATORY USE:
<input type="checkbox"/> Delayed Puberty	GE Number: _____
<input type="checkbox"/> Short Stature	Amount Received: _____
<input type="checkbox"/> Fanconi Anemia	Date Received: _____ DD/MONTH/YYY
<input type="checkbox"/> Other: _____	Date Set up: _____ DD/MONTH/YYY
	Name: _____
	Signature: _____

Comments:

Name: _____ Signature: _____ Date: _____
DD/MONTH/YYYY