

## CONSTITUTIONAL TEST REQUEST FORM

For UW Pathology use	
MRN:	Accession #

<b>1 Patient Information</b>	First Name	MI	Last Name
	Sex	DOB	SSN
	Patient Address		
	City	State	Zip
	Patient Phone #	Outside Facility Patient ID	

<b>2 Requesting Institution</b>	Institution Name		
	Institution Address		
	City	State	Zip
	Person Completing Form		
	Phone	Fax	

<b>3 Send Reports to</b>	Requesting Physician (primary):	Phone	Fax	NPI#
	Referring Physician/Surgeon:	Phone	Fax	NPI#
	Referring Pathologist:	Phone	Fax	NPI#
	Additional reports to:	Phone	Fax	NPI#

<b>4 Billing Information</b>	<b>Payment Options:</b>	<input type="checkbox"/> Patient Insurance* (If outpatient) <input type="checkbox"/> Self-Pay (No insurance) <input type="checkbox"/> Institution/Client Billing <input type="checkbox"/> Split Billing / Medicare* (Pro to Patient, Tech to Client)					
	*Medicare Billing policy does not permit tech claims on laboratory testing for hospital inpatients/outpatients. These tech charges will be billed to the requesting institution.						
	Primary Insurance			Secondary Insurance			
	ID/Policy #	Group #	ID/Policy #	Group #			
	Insurance Address		Phone	Insurance Address		Phone	
	City/State/Zip			City/State/Zip			
Insured's Name	DOB	Relation to Pt:	Insured's Name	DOB	Relation to Pt:		

Note: For sample collection requirements see <http://www.UWPathology.org/clinical/cytogenetics>

<b>5 Specimen Type</b>	<b>Date obtained:</b>
<input type="checkbox"/> Peripheral Blood <input type="checkbox"/> Amniotic Fluid (Gestational Age: _____) <input type="checkbox"/> Chorionic Villi (Gestational Age: _____) <input type="checkbox"/> Products of Conception (Gestational Age: _____) <input type="checkbox"/> Fetal Tissue (Site: _____) <input type="checkbox"/> Umbilical Cord Blood <input type="checkbox"/> Skin Biopsy (Site: _____) <input type="checkbox"/> Saliva <input type="checkbox"/> Paraffin Blocks/Slides (Site: _____) <input type="checkbox"/> DNA	

<b>6 Diagnosis or Indication for Testing</b>
Please attach copy of pedigree if indication is Family History of..
ICD-10 Code: _____
<input type="checkbox"/> This is a family follow-up study (Name of proband: _____)

\*\*\* SEE PAGE 2 FOR TESTS \*\*\*

Ordering Provider Signature Required	
Submitting a specimen with this requisition form indicates familiarity and agreement with applicable Reference Laboratory Services policies found at <a href="http://pathology.washington.edu/clinical/servicerequest">http://pathology.washington.edu/clinical/servicerequest</a>	
Signature	Date

**7 Test(s) Requested** **STAT**  **ROUTINE**

- Interphase FISH for common aneuploidies (13, 18, 21, X, Y)
- Interphase FISH after pregnancy loss (13, 15, 16, 18, 21, 22, X, Y)
- Metaphase FISH for:
 

<input type="checkbox"/> 1p36.1 deletion	<input type="checkbox"/> Prader-Willi syndrome (15q11.2 deletion)
<input type="checkbox"/> 15q11-q13 duplication (autism)	<input type="checkbox"/> SHOX-related haploinsufficiency
<input type="checkbox"/> 22q11.2 deletion (VCFS/DiGeorge)	<input type="checkbox"/> Smith-Magenis syndrome (17p11.2 deletion)
<input type="checkbox"/> 22q11.2 duplication	<input type="checkbox"/> Sotos syndrome (5q35 deletion)
<input type="checkbox"/> Angelman syndrome (15q11.2 deletion)	<input type="checkbox"/> SRY (46,XX testicular DSD/46,XY DSD/46,XY CGD)
<input type="checkbox"/> Cri du Chat syndrome (5p deletion)	<input type="checkbox"/> Subtelomeres (Specify: _____)
<input type="checkbox"/> Kallmann syndrome	<input type="checkbox"/> Williams syndrome (7q11.23 deletion)
<input type="checkbox"/> Langer-Giedion (8q24 deletion)	<input type="checkbox"/> Williams-Beuren region duplication (7q11.23 duplication)
<input type="checkbox"/> Miller-Diecker syndrome (17p13.3 deletion)	<input type="checkbox"/> Wolf-Hirschhorn (4p deletion)
<input type="checkbox"/> Pallister-Killian syndrome (iso12p mosaicism)	<input type="checkbox"/> X-linked ichthyosis (STS deletion)
<input type="checkbox"/> Potocki-Lupski syndrome (17p11.2 duplication)	<input type="checkbox"/> Other (Specify: _____)
- Cytogenomic Microarray Analysis (CMA/CGH/CGAT/SNP Array)
  - Report all findings
  - Do not report variants of uncertain clinical significance
- ddPCR (droplet digital PCR) for small deletion or duplication (Specify: \_\_\_\_\_)
- Routine G-banded chromosome analysis and karyotyping
- Mosaicism study by chromosome analysis and karyotyping
  - Mosaicism for: \_\_\_\_\_
- Limited parental follow-up study by chromosome analysis and karyotyping
- Y chromosome deletions by PCR for male infertility
- Grow cell cultures for sendout
  - Sendout instructions:

**Reflex Testing**

- If \_\_\_\_\_ is  Normal then reflex to \_\_\_\_\_  
 Abnormal
- If \_\_\_\_\_ is  Normal then reflex to \_\_\_\_\_  
 Abnormal