



University of Florida Health  
Pathology Laboratories • Cytogenetics  
4800 SW 35<sup>th</sup> Drive • Gainesville, FL 32608

## STANDARD (CONSTITUTIONAL) CYTOGENETIC TESTING REQUISITION FORM

Reference our other cytogenetics requisition forms for additional tests not listed here, or visit us online at:

[pathlabs.ufl.edu/services/cytogenetics](http://pathlabs.ufl.edu/services/cytogenetics)

Telephone: 352.265.9900  
Toll-Free: 888.375.5227  
Fax: 352.265.9920

### Patient Information

Name: \_\_\_\_\_  
Medical record #: \_\_\_\_\_  
Age or DOB: \_\_\_\_\_  
Sex/Gender:  Female  Male  Unknown

### Requesting Physician Information

Name: \_\_\_\_\_ NPI #: \_\_\_\_\_  
Location/Institution: \_\_\_\_\_  
Signature: \_\_\_\_\_  
Send additional reports to: \_\_\_\_\_

### Clinical Indication or Reason for Cytogenetic Testing

- Trisomy 21  Mental retardation  Turner syndrome  Multiple fetal loss  
 Trisomy 18  Developmental delay  Klinefelter syndrome  Infertility  
 Trisomy 13  Dysmorphic features  XYY syndrome  Ambiguous genitalia  
 Fetal demise  Failure to thrive  XXX syndrome  MCA  
 Family history of chromosome abnormality (explain): \_\_\_\_\_  
 Other: \_\_\_\_\_

### Specimen Information

- Peripheral blood  
 Skin  POC  
 Other (indicate type): \_\_\_\_\_  
Collection date: \_\_\_\_\_  
Collection time: \_\_\_\_\_

### Cytogenetic Testing Requested (must be completed to avoid delays in processing)

#### Conventional Chromosome Analyses (Karyotyping)

- Routine/Conventional chromosome study  
 Other  
Please specify: \_\_\_\_\_  
 Cell line buildup for additional and/or outside studies  
Please specify: \_\_\_\_\_

#### FISH Analyses

All requests for FISH analysis must include a routine chromosome study.

- 22q11.2 - DiGeorge/VCF syndromes  
 Yp11.3 - SRY  X/Y  
Other: \_\_\_\_\_  
(inquire on availability)

**Note:** Metaphase FISH testing previously offered for microdeletion and/or microduplication syndromes is now facilitated by comparative genomic hybridization microarray (aCGH) testing methods.

A **CGH+SNP microarray** is the current in-house aCGH platform recommended for congenital applications and can be ordered by utilizing a separate requisition form available on our website or by calling **352.265.9900**.

### For Lab Use Only

Lab #: \_\_\_\_\_  
Specimen description: \_\_\_\_\_  
# of containers: \_\_\_\_\_ Quantity (ml): \_\_\_\_\_  
 Sodium heparin tube  
Other: \_\_\_\_\_  
Additional test codes: \_\_\_\_\_  
Tech Login ID: \_\_\_\_\_

### Insurance/Billing Information (must be completed prior to sample processing)

Insurance provider: \_\_\_\_\_  
Preauthorization required?:  Yes  No  
If yes, provide the authorization number: \_\_\_\_\_