



THE UNIVERSITY OF
ALABAMA AT BIRMINGHAM

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12/20/16

Medical Record # _____

PATIENT HISTORY AND REQUEST FORM FOR BLOOD CYTOGENETIC ANALYSIS

Patient Name: _____
Address: _____
Phone: _____

Date of Birth: _____
Race: _____
Sex: _____

Requesting Physician: _____

Phone: _____

Billing Information: _____
Address: _____

Phone: _____
FAX: _____

Reason for studies: _____

Family history of a chromosome anomaly? Yes No (If yes, please provide the information below.)

Who was tested? _____

What was the result? _____

Where were they studied? _____

Type of specimen: _____

Collected: Date: _____
Time: _____

Studies requested

		<u>FISH</u>	
<input type="checkbox"/> Routine chromosome analysis	<input type="checkbox"/> Rapid Aneuscreen (X,Y,13,18,21)	<input type="checkbox"/> Prader-Willi	
<input type="checkbox"/> HRB chromosome analysis	<input type="checkbox"/> X/Y (Sex chromosomes)	<input type="checkbox"/> Smith-Magenis	
	<input type="checkbox"/> Angelman	<input type="checkbox"/> SRY	
	<input type="checkbox"/> Cri-du-Chat	<input type="checkbox"/> Williams	
	<input type="checkbox"/> DiGeorge	<input type="checkbox"/> Wolf-Hirschhorn	
<input type="checkbox"/> aCGH+SNP analysis	<input type="checkbox"/> Kallman	<input type="checkbox"/> RP11-_____	
	<input type="checkbox"/> Miller-Dieker		
	<input type="checkbox"/> Steroid Sulfatase Deficiency		

(Indicate which
RP11 probe)

Blood specimens for chromosomes and/or FISH studies (5-6cc) should be sent in a Sodium heparinized vacutainer. If aCGH is requested, an additional lavender EDTA tube with 5-6 cc is needed.

Specimens collected in lithium heparin or EDTA cannot be used for chromosome analysis.

Specimens should be transported as soon as possible at room temperature.