

# BCM-MEDICAL GENETICS LABORATORIES

PHONE: 800-411-GENE | FAX: 713-798-2787 | www.bcmgeneticlabs.org

SHIP TO: Medical Genetics Laboratories  
Baylor College of Medicine  
2450 Holcombe, Grand Blvd. -Receiving Dock  
Houston, TX 77021-2024  
Phone: 713-798-6555

## CYTOGENETICS REQUISITION

PATIENT INFORMATION	REPORTING INFORMATION
NAME: _____ LAST NAME FIRST NAME MI	PHYSICIAN: _____ INSTITUTION: _____
DATE OF BIRTH: ____ / ____ / ____ MM DD YY GENDER (Please select one): <input type="checkbox"/> FEMALE <input type="checkbox"/> MALE <input type="checkbox"/> UNKNOWN	PHONE: _____ FAX: _____ EMAIL (INTERNATIONAL CLIENT REQUIREMENT): _____
DATE OF COLLECTION: ____ / ____ / ____ MM DD YY SAMPLE TYPE (Please select one): <input type="checkbox"/> BLOOD <input type="checkbox"/> TISSUE <input type="checkbox"/> OTHER (Specify): _____	<b>ADDITIONAL PROFESSIONAL REPORT RECIPIENTS</b>
HOSPITAL#: _____ ACCESSION#: _____	NAME: _____ PHONE: _____ FAX: _____ NAME: _____ PHONE: _____ FAX: _____

## INDICATION FOR STUDY

CMA OPTIONS MOST APPROPRIATE 1ST TIER TEST FOR INDICATIONS BELOW:

- AUTISM SPECTRUM
- DEVELOPMENTAL DELAY  
 MILD  MODERATE  SEVERE
- DYSMORPHIC FEATURES
- OTHER (Specify): \_\_\_\_\_
- FAILURE TO THRIVE
- MULTIPLE CONGENITAL ANOMALIES
- SEIZURE DISORDER

CHROMOSOMES/FISH MOST APPROPRIATE 1ST TIER TEST FOR INDICATIONS BELOW:

- AUTOSOMAL TRISOMIES
- AMBIGUOUS GENITALIA
- INFERTILITY
- OTHER (Specify): \_\_\_\_\_
- KLINEFELTER/TURNER
- MULTIPLE MISCARRIAGES
- FETAL DEMISE

### CHROMOSOMAL MICROARRAY ANALYSIS (CMA) OPTIONS

- 8665 CMA-HR+SNP screen (comprehensive)  
(340K High Resolution Copy Number probes Plus 60K SNP probes (UPD/AOH))  
Specify Gene of Interest: \_\_\_\_\_  
If UPD suspected, specify chromosome: \_\_\_\_\_
- 8655 CMA-HR (180K High Resolution Copy Number ≥30Kb)  
Specify Gene of Interest: \_\_\_\_\_
- 8650 CMA-CytoScan HD (2.6M probes)  
If UPD suspected, specify chromosome: \_\_\_\_\_
- 9506 CMA-POC (paraffin-embedded tissue only)
- PARENTAL STUDIES RECOMMENDED IN CHILD'S CMA REPORT (Attach copy.)
  - Mother (Full Name): \_\_\_\_\_  
 ASYMPTOMATIC  SYMPTOMATIC (attach summary of findings)
  - Father (Full Name): \_\_\_\_\_  
 ASYMPTOMATIC  SYMPTOMATIC (attach summary of findings)

### CYTOGENETIC TESTING OPTIONS

- 8600 Chromosome Analysis
- 8480 FISH for SRY - Related Phenotypes (Metaphase & Interphase Cells)\*
- 8425 Rapid FISH - AneuVysion®: +13/+18/+21/X/Y (Interphase Cells ONLY)  
ONE of the following tests MUST accompany the 8425 request.
  - 8665 CMA-HR+SNP screen (comprehensive)
  - 8655 CMA-HR
  - 8600 Chromosome Analysis
- OR
- 8426 Rapid FISH - Sex Chromosome: X/SRY (Interphase Cells ONLY)  
ONE of the following tests MUST accompany the 8426 request.
  - 8665 CMA-HR+SNP screen (comprehensive)
  - 8655 CMA-HR
  - 8600 Chromosome Analysis
- OR

\*Testing on Metaphase Cells require cell culturing

\*\*\* ADDITIONAL FISH OPTIONS FOUND ON PAGE 2 \*\*\*

## REQUIRED: NEW YORK STATE PHYSICIAN SIGNATURE OF CONSENT

I certify that the patient specified above and/or their legal guardian has been informed of the benefits, risks, and limitations of the laboratory test(s) requested. I have answered this person's questions. I have obtained informed consent from the patient or their legal guardian for this testing.

Physician's Printed Name: \_\_\_\_\_ Physician's Signature: \_\_\_\_\_ Date (MM/DD/YY): \_\_\_\_\_

## CYTOGENETICS REQUISITION

NAME: \_\_\_\_\_ DATE OF BIRTH: \_\_\_\_ / \_\_\_\_ / \_\_\_\_ GENDER (Please select one):  FEMALE  
LAST NAME FIRST NAME MI MM DD YY  MALE  
 UNKNOWN

### FISH STUDIES

<input type="checkbox"/>	8457 Adrenal Hypoplasia Congenita
<input type="checkbox"/>	8458 JAG1-Related Alagille Syndrome
<input type="checkbox"/>	8459 Angelman Syndrome
<input type="checkbox"/>	8460 Beckwith-Wiedemann Syndrome
<input type="checkbox"/>	8462 Charcot-Marie-Tooth Neuropathy
<input type="checkbox"/>	8464 Cri-Du-Chat Syndrome
<input type="checkbox"/>	8440 DiGeorge/Velocardiofacial Syndrome (22q and 10p) Panel
<input type="checkbox"/>	8486 DiGeorge/Velocardiofacial Syndrome Type I (22q)
<input type="checkbox"/>	8465 DiGeorge/Velocardiofacial Syndrome Type II (10p)
<input type="checkbox"/>	8466 Glycerol Kinase Deficiency
<input type="checkbox"/>	8467 Hereditary Neuropathy With Liability To Pressure Palsies
<input type="checkbox"/>	8468 LIS1-Associated Lissencephaly

<input type="checkbox"/>	8469 Kallmann Syndrome
<input type="checkbox"/>	8430 Langer-Giedion Syndrome (EXT1 and TRPS1)
<input type="checkbox"/>	8470 Microphthalmia With Linear Skin Lesion (MLS/MIDAS)
<input type="checkbox"/>	8471 Miller-Dieker Syndrome
<input type="checkbox"/>	8435 Multiple Exostoses (EXT1 and EXT2) Panel
<input type="checkbox"/>	8472 Multiple Exostoses Type I (EXT1)
<input type="checkbox"/>	8473 Multiple Exostoses Type II (EXT2)
<input type="checkbox"/>	8474 Neurofibromatosis Type I
<input type="checkbox"/>	8456 1p36 Deletion Syndrome
<input type="checkbox"/>	8476 Prader-Willi Syndrome
<input type="checkbox"/>	8477 Rubinstein-Taybi Syndrome

<input type="checkbox"/>	8478 Smith-Magenis Syndrome
<input type="checkbox"/>	8479 Sotos Syndrome
<input type="checkbox"/>	8480 SRY-Related Phenotypes
<input type="checkbox"/>	8482 Trichorhinophalangeal Syndrome Type I
<input type="checkbox"/>	8450 WAGR (WT1 and PAX6) FISH Panel
<input type="checkbox"/>	8483 Williams Syndrome
<input type="checkbox"/>	8455 Wilms Tumor (WT1)
<input type="checkbox"/>	8484 Wolf-Hirschhorn Syndrome
<input type="checkbox"/>	8485 X-linked Ichthyosis

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## CYTOGENETICS REQUISITION

### BILLING INFORMATION

IMPORTANT NOTICE: ONE OF THE THREE FOLLOWING BILLING OPTIONS MUST BE INDICATED BELOW.  
PLEASE FORWARD ALL BILLING QUESTIONS TO: [MEDGENBILLING@BCM.EDU](mailto:MEDGENBILLING@BCM.EDU)

### PATIENT INFORMATION

PATIENT NAME (LAST, FIRST, MI): \_\_\_\_\_ PATIENT DATE OF BIRTH (MM/DD/YY): \_\_\_\_\_  
ADDRESS: \_\_\_\_\_ CITY, STATE, ZIP: \_\_\_\_\_  
PHONE: \_\_\_\_\_ EMAIL: \_\_\_\_\_

#### PAYMENT OPTION 1 - INSTITUTION

INSTITUTION NAME: \_\_\_\_\_ INSTITUTION CODE: \_\_\_\_\_  
CONTACT NAME: \_\_\_\_\_ EMAIL (REQUIRED): \_\_\_\_\_  
BILLING ADDRESS: \_\_\_\_\_ CITY, STATE, ZIP: \_\_\_\_\_  
PHONE: \_\_\_\_\_ FAX: \_\_\_\_\_

#### PAYMENT OPTION 2 - SELF-PAY (PAYMENT MUST ACCOMPANY SAMPLE)

CREDIT CARD (PLEASE SELECT ONE):  AMEX  DISCOVER  MC  VISA  
VALID CARD #: \_\_\_\_\_ EXPIRATION DATE (MM/YY): \_\_\_\_\_ CVC CODE: \_\_\_\_\_  
BILLING ADDRESS: \_\_\_\_\_ CITY, STATE, ZIP: \_\_\_\_\_  
CARDHOLDER PRINTED NAME: \_\_\_\_\_ CARDHOLDER SIGNATURE: \_\_\_\_\_  
 CHECK/MONEY ORDER: CHECK/MONEY ORDER #: \_\_\_\_\_ AMOUNT ENCLOSED: \_\_\_\_\_

#### PAYMENT OPTION 3 - INSURANCE

PROVIDE A LEGIBLE PHOTOCOPY OF THE FRONT & BACK OF THE INSURANCE CARD OR HMO/MEDICAID HMO AUTHORIZATION/REFERRAL.

Please refer to the Financial Policy at [www.bcmgeneticlabs.org](http://www.bcmgeneticlabs.org) for complete insurance filing information and managed care contract list. Insurance is filed to our contracted carriers as a client service courtesy. Patients are responsible for non-covered services, deductibles, co-insurance, contract exclusions, non-authorized services, and remaining balances after insurance reimbursement. HMO policies must have required approved authorizations. BCM-Medical Genetic Laboratories cannot bill out-of-state welfare programs. We accept authorized Texas Medicaid HMO covered charges for genetic testing. Please contact our office prior to submitting a Texas Medicaid sample. Contact [medgenbilling@bcm.edu](mailto:medgenbilling@bcm.edu) with questions.

ORDERING PROVIDER: \_\_\_\_\_

ICD9 Diagnosis Code(s) - must be provided or insurance cannot be filed: \_\_\_\_\_

- PPO, POS, Commercial Insurance - Provide complete member information with legible front & back photocopy of insurance card.  
 HMO - Provide approved authorization #: \_\_\_\_\_ and attach legible front & back photocopy of insurance card.  
 Texas Medicaid HMO - Provide approved authorization #: \_\_\_\_\_ and contact Billing at 713-798-6555.

### INSURED MEMBER'S INFORMATION

MEMBER NAME (Last, First, MI): \_\_\_\_\_ MEMBER DATE OF BIRTH (MM/DD/YY): \_\_\_\_\_ GENDER:  FEMALE  MALE  
MEMBER POLICY #: \_\_\_\_\_ MEMBER SS #: \_\_\_\_\_ MEMBER GROUP #: \_\_\_\_\_  
INSURANCE CO. NAME: \_\_\_\_\_ PHONE: \_\_\_\_\_  
INSURANCE CO. ADDRESS: \_\_\_\_\_ CITY, STATE, ZIP: \_\_\_\_\_

I AUTHORIZE BCM-MEDICAL GENETICS LABORATORIES TO FURNISH ANY MEDICAL INFORMATION REQUESTED ON MYSELF, OR MY COVERED DEPENDENTS. IN CONSIDERATION OF SERVICES RENDERED, I TRANSFER AND ASSIGN ANY BENEFITS OF INSURANCE TO BCM-MEDICAL GENETICS LABORATORIES. I UNDERSTAND I AM RESPONSIBLE FOR ANY CO-PAY, DEDUCTIBLES, OR NON-AUTHORIZED SERVICES AND REMAINING BALANCES AFTER INSURANCE REIMBURSEMENT. I UNDERSTAND I AM FULLY RESPONSIBLE FOR PAYMENT OF MY ACCOUNT IF THE BCM-MEDICAL GENETICS LABORATORIES IS NOT A PARTICIPANT WITH MY HEALTH PLAN, AND MY HEALTH PLAN DOES NOT FULLY REIMBURSE MY MEDICAL SERVICES DUE TO LACK OF AUTHORIZATION OR MEDICAL NECESSITY.

PRINTED NAME: \_\_\_\_\_ SIGNATURE: \_\_\_\_\_ DATE (MM/YY): \_\_\_\_\_