

LAB RECEPTION DATE/TIME:

LAB DESCRIPTION:

STAMP OR LABEL HERE

Centre universitaire de santé McGill
McGill University Health Centre

SEND TO: DR. ANDREA RUCHON
MOLECULAR DIAGNOSTIC LABORATORY
E05.3028 LABORATORY CENTRALIZED RECEPTION
1001 DECARIE BLVD, MONTREAL, QC, H4A 3J1
TEL: 514-934-1934 EXT 23383/23298
FAX: TBD
CLIA #: 99D1042152



Montreal Children's Hospital
Molecular Genetics

*SAMPLES CANNOT BE PROCESSED
WITHOUT REQUIRED INFORMATION

PATIENT INFORMATION

1 REQUISITION PER PATIENT

REQUIRED

GENDER: MALE FEMALE UNKNOWN

PATIENT'S NAME:

LEGAL NAME FIRST LAST

DATE OF BIRTH:

MONTH / DAY / YEAR

N/A

PROVINCIAL HEALTH #:

CANADA ONLY

QUEBEC RESIDENTS RAMQ#

HOSPITAL REF #:

ADDRESS (PATIENT):

FATHER:

MOTHER:

ETHNICITY (CHOOSE ALL THAT APPLY):

- FRENCH CANADIAN (SELECT REGION IF RELEVANT)
- SAGUENAY-LAC-ST-JEAN CHARLEVOIX
- ASHKENAZI JEWISH
- OTHER (PLEASE SPECIFY): _____

PHYSICIAN INFORMATION (REQUIRED)

REQUIRED

PHYSICIAN'S NAME:

LICENSE #:

INSTITUTION:

ADDRESS:

PHONE:

FAX:

GENETIC COUNSELLOR:

PHONE:

FAX:

PHYSICIAN/GENETIC COUNSELLOR

SIGNATURE: _____ DATE: _____
MONTH/DAY/YEAR

I ACKNOWLEDGE THAT THE PATIENT/GUARDIAN ARE AWARE OF THE BENEFITS, LIMITATIONS & RISKS ASSOCIATED WITH THE REQUESTED TEST(S) AND THAT I HAVE OBTAINED INFORMED CONSENT TO PERFORM GENETIC TESTING FOR THIS PATIENT.

SAMPLE INFORMATION (REQUIRED)

LABORATORY RECEPTION HOURS: MON-FRI 8:00-16:00
PRENATAL SAMPLES MON-THR 8:00 - 16:00

COLLECTION DATE ____ / ____ / 20 ____ TIME: ____ AM / PM
MONTH DAY YEAR

BLOOD (EDTA/LAVENDER) 2cc NEWBORN 5cc ADULT

PRENATAL (CONTACT LAB FOR INSTRUCTIONS)

AMNIOTIC FLUID CULTURED AMNIOCYTES CVS min 10 mg (DIRECT)

DNA : SOURCE _____ CONC. ____ (/)

OTHER (SPECIFY): _____

EACH SAMPLE MUST BE LABELLED WITH THE PATIENT'S NAME AND DOB. IF BLOOD SAMPLES CANNOT BE SHIPPED THE SAME DAY STORE AT 4°C. DO NOT FREEZE. BLOOD TUBES SHOULD BE PROPERLY PACKAGED TO PROTECT AGAINST SHOCK AND DAMAGE.

PEDIGREE/CLINICAL INFORMATION (PLEASE COMPLETE)

ONGOING PREGNANCY INVOLVED

GESTATIONAL AGE: ____ WEEKS ON ____ / ____ / 20 ____ LMP US
MONTH DAY YEAR

IF OTHER SAMPLES SUBMITTED INDICATE NAME/DOB & RELATIONSHIP TO PATIENT.

PREVIOUSLY TESTED FAMILY MEMBERS

LAB FAMILY #: _____

NAME(S): _____

MUTATION(S) UNKNOWN KNOWN (SPECIFY) _____

TEST REQUESTED (REQUIRED)

CFTR-RELATED DISORDERS (SELECT ALL THAT APPLY)

SWEAT CHLORIDE VALUES: UNAVAILABLE NO YES (RESULT _____)

- FETAL ECHOGENIC BOWEL CAVD/CBAVD
- PANCREATIC INSUFFICIENCY FAILURE TO THRIVE MECONIUM ILEUS
- CHRONIC PANCREATITIS BRONCHIECTASIS ASTHMA
- SINUSITIS RECTAL PROLAPSE NASAL POLYPS

RESPIRATORY INFECTION (SPECIFY) _____

OTHER: _____

ANGELMAN SYNDROME METHYLATION UPD15

PRADER-WILLI SYNDROME METHYLATION UPD15

ASHKENAZI JEWISH PANEL (FAMILIAL DYSAUTONOMIA/TAYSACHS/CANAVAN)

DEAFNESS (CONNEXIN26/30)

CLOUSTON (HED2)

MCAD DEFICIENCY

PAH DEFICIENCY PKU HYPERPHENYLALANINEMIA

TAY-SACHS ENZYMES: NO YES (PLEASE ATTACH RESULTS)

UPD15

Y-CHROMOSOME MICRODELETIONS

HBB-RELATED DISORDERS* (SELECT ALL THAT APPLY)

HbS HbC HbE OTHER: _____

β -THALASSEMIA: MAJOR INTERMEDIA TRAIT/MINOR

* INCLUDE RELEVANT LAB FINDINGS (EG. CBC; ELECTROPHORESIS; HPLC)

INDICATION FOR TESTING (REQUIRED)

CONFIRM CLINICAL DIAGNOSIS

POSITIVE NEWBORN SCREENING

CARRIER STATUS (ASYMPTOMATIC)

FAMILY HISTORY (PLEASE INDICATE RELATIONSHIP IN PEDIGREE)

OTHER (SPECIFY REASON) _____

PRENATAL DIAGNOSIS (MATERNAL SAMPLE REQUIRED)

MATERNAL CELL CONTAMINATION STUDIES

OTHER (SPECIFY) _____

DRAW ON BACK OR ATTACH PEDIGREE IF NECESSARY

REQUIRED INFORMATION