MUHC-GLEN CYTOGENETICS LABORATORY REQUISITION

POSTNATAL CHROMOSOMAL SNP MICROARRAY

COMPLETE IN FULL TO AVOID DELAY IN REPORTING RESULTS			
PRESCRIBER		Patient identification	
DD)	First name : MRN or RAMO Address : Date of birth (Y) Gender :	Q: YYY/MM/DD):	
		ninimum (1 mL minimum for newborns) (requires consult with Cytogeneticist)	
Date (YYYY/MM/DD):		Time:	
		Phenotype : Normal Abnormal	
INDICATIONS FOR TESTING: □ Developmental delay or intellectual disability □ Developmental delay or intellectual disability and additional clinical features (complete page 2). □ Congenital anomalies (complete page 2).			
PRENATAL, PERINATAL AND FAMILY HISTORY			
ns found prenatally, perinatally	, in family (describe		
	PRESCRIBER DD)	PRESCRIBER	

DECRIP + TORY
RESPIRATORY Display a greatic harrie
☐ Diaphragmatic hernia☐ Lung abnormality (Specify)
Lung abnormanty (Specify)
Othor
□ Other:
——— MUSCULOSKELETAL
☐ Upper limb abnormality
☐ Lower limb abnormality
\Box Camptodactyly (\Box finger / \Box toe)
\square Syndactyly (\square fingers / \square toes)
□ Polydactyly (□ finger / □ toe
□ Preaxial □ Postaxial
□ Oligodactyly (□ finger / □ toe)
☐ Clinodactyly (☐ finger / ☐ toe)
☐ Vertebral anomaly
□ Club foot
□ Other:
GASTROINTESTINAL
☐ Esophageal atresia
——— ☐ Tracheoesophageal fistula
☐ Gastroschisis
□ Omphalocele
☐ Pyloric stenosis
Other:
n Tag
""
GENITOURINARY
☐ Kidney malformation (Specify)
)
☐ Hydronephrosis
☐ Ambiguous genitalia
☐ Hypospadias
☐ Cryptorchidism
Other:

THIS REQUISITION MUST BE ACCOMPANIED BY A SIGNED INFORMED CONSENT

available at https://muhc.ca/muhc-clinical-laboratories/muhc-laboratory-requisitions-and-consultations