SickKids®
THE HOSPITAL FOR SICK CHILDREN
Paediatric Laboratory Medicine

555 University Avenue
Room 3416, Roy C. Hill Wing
Toronto, ON, M5G 1X8, Canada
Tel: 416-813-7200 x1
Fax: 416-813-7732
(CLIA # 99D1014032)

Genome Diagnostics

www.sickkids.ca/en/care-services/for-health-care-providers/lab-testing-services

Patient Name:
Preferred Name (if different):
Date of Birth (DD/MM/YYYY):
Legal Sex: ◯ *Male* ◯ *Female* ◯ *Non-binary/U/X*
Sex Assigned at Birth (if different): ◯ *Male* ◯ *Female* ◯ *Unassigned*
Gender Identity): ◯ *Male* ◯ *Female* ◯ *Non-binary/U/X*
MRN:
Parent's Name:
Address:

For Canada Only
Provincial Health Card #: *Version:*
Issuing Province:

Testing is provided for medical purposes only and results are not intended for forensic use. The laboratory is not a forensically accredited laboratory.

Referring Physician *(required)*:

Name: _____

Facility/Ward/Clinic (***required***):_____

Address: _____

_____

Phone:_____ Fax:_____

Email address: _____

Signature:_____

Copy Report To Another Healthcare Provider *(all information is required)*:

Name: _____

Address: _____

Phone:_____ Fax:_____

Sample Information *(required)*:

Date obtained (DD/MM/YYYY):-_____-_____ Referring

Laboratory reference #: _____

☐ Blood in EDTA (purple top tube): min. 4 mL (0.5-3 mL for newborns)
☐ DNA: min.10 ug in low TE buffer (Source:_____)
* Unable to perform MLPA analysis on externally extracted DNA (contact lab)
☐ Direct CVS: min. 10 mg direct villi
☐ Cultured villi: **1 flask at 60-70% confluency and 1 flask at 80-90% confluency**
☐ Cultured amniocytes: **1 flask at 60-70% confluency and 1 flask at 80-90% confluency**
☐ Tissue (Source:_____)
☐ Other (Specify:_____)

Closed consent:
☐ (If checked, all remaining DNA will be discarded upon notification by the ordering physician that all DNA testing has been completed)

Laboratory Use:

Date (DD/MM/YYYY) | Time Received:

_____-_____-_____|_____h

Lab/Order #:_____

Specimen type, amt & # of tubes:_____

Comments:

Pedigree/Family No./Patient/Order No._____/_____

Reason for Testing (*required*):

☐ Diagnosis ☐ Carrier testing
☐ Familial mutation/variant analysis ☐ Prenatal testing
☐ Parental sample ☐ Variant re-assessment

☐ Other (Specify): _____

If expedited testing is requested, indicate reason:

☐ Pregnancy (Gestational age (weeks)_____
☐ Other (Specify): _____

Familial Mutation / Targeted Variant Analysis:

* ***If proband testing was performed elsewhere, a copy of the original report (all pages) is required. Send a positive control sample if available.***

Gene & NM #: _____

Mutation/variant(s): _____

SickKids Laboratory/Order number:_____

SickKids Pedigree/Family number:_____

Name of proband:_____

Relationship to proband:_____

Name(s) & DOB of other submitted family members: _____

_____

Clinical Diagnostics and Family History *(required)*:

Draw or attach a pedigree and provide any relevant information below, including clinical and family history details, as this is important for accurate interpretation of results.

Ethnicity: _____

Ordering Checklist:

☐ Specimen tube labeled with at least two identifiers
☐ Completed test requisition form
Clinical information must be provided for all tests. Pages 4-5 must be completed for all tests. Testing will not proceed until these are provided.
☐ Proband's report and positive control (*familial/targeted variant testing only*)
☐ Completed billing form *(page 6, if applicable)*

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LIST OF TESTS AVAILABLE BY DISEASE
For prenatal testing and cases where a familial mutations/variant is known, include information on page 1

22q11 Deletion Syndrome
☐ 22q11 deletion/duplication analysis (*external DNA not accepted*)

Angelman Syndrome
☐ Methylation and deletion/duplication analysis (*external DNA not accepted*)
☐ UPD15 analysis *(please submit parental samples)*

Ashkenazi Jewish Carrier Screening
☐ Recurrent mutation analysis (7 diseases):
Bloom syndrome, Canavan disease, Familial Dysautonomia, Fanconi Anemia Group C, Mucolipidosis Type IV, Niemann-Pick disease, Tay-Sachs disease

ETHNICITY (required):
☐ Ashkenazic ☐ Sephardic ☐ French Canadian ☐ Cajun
☐ Non-Jewish ☐ Other_____

Autoinflammatory Disease *
Clinical information must be provided on pages 4 and 5
☐ Autoinflammatory Diseases NGS panel
(excludes Recurrent Fever panel genes)
☐ Recurrent Fever Syndrome NGS panel
MEFV (FMF), MVK, NLRP12, NLRP3, TNFRSF1A
☐ Hemophagocytic Lymphohistiocytosis NGS panel
☐ Aicardi-Goutieres Syndrome NGS panel
☐ Deletion/duplication analysis

Becker Muscular Dystrophy
☐ *DMD* deletion/duplication analysis (*external DNA not accepted*)
☐ *DMD* sequence analysis

Beckwith-Wiedemann Syndrome
☐ IC1 and IC2 methylation† and 11p15 deletion/duplication analysis (*external DNA not accepted*)
☐ UPD11 analysis (parental sample required)
☐ *CDKN1C* sequence analysis
† **No methylation analysis on CVS samples**

Bone Marrow Transplantation
☐ Post-transplant monitoring

Cancer Related Tests

Li-Fraumeni Syndrome
☐ *TP53* sequence analysis
☐ *TP53* deletion/duplication analysis (*external DNA not accepted*)

Rhabdoid Tumour Predisposition Syndrome
☐ *SMARCB1* sequence analysis
☐ *SMARCB1* deletion/duplication analysis (*external DNA not accepted*)

Congenital Muscular Dystrophies
☐ Sequence analysis panel:
FKTN (FCMD), FKRP, POMGnT1, POMT1, POMT2

Connective Tissue Disease *
Clinical information must be provided on pages 4 and 5
If more than one panel is requested, rationale must be provided on page 5.
☐ Ehlers Danlos Syndrome panel
☐ Osteogenesis Imperfecta panel
☐ Osteopetrosis and Disorders of Increased Bone Density panel
☐ Bone Involvement panel
☐ Deletion/duplication analysis

Craniosynostosis
☐ Apert Syndrome (*FGFR2* recurrent mutations analysis)
☐ Crouzon Syndrome (*FGFR2, FGFR3* recurrent mutation analysis)
☐ Pfeiffer Syndrome (*FGFR1, FGFR2, FGFR3* recurrent mutation analysis)
☐ Saethre-Chotzen Syndrome (*TWIST1* sequence analysis and *FGFR3* recurrent mutation analysis)
☐ Non-Syndromic Craniosynostosis (*FGFR3* recurrent mutation analysis)
☐ *TWIST1* deletion/duplication analysis (*external DNA not accepted*)

Cystic Fibrosis and/or *CFTR*-Related Disorders **
Indication (provide additional clinical details on page 1 and/or pages 4-5):
☐ Fetal echogenic bowel (*ensure parental samples are linked to each other on both requisitions with at least two identifiers*)
☐ Clinical diagnosis of cystic fibrosis
☐ CFTR-related disorders
☐ Male factor infertility: ☐ oligo/azoospermia ☐ C(B)AVD
☐ Family history of cystic fibrosis
☐ Positive newborn screen (*ensure familial samples are linked to each other on all requisitions with at least two identifiers; send NSO report*)

Tests (indication specific):
☐ *CFTR* recurrent mutation analysis
☐ *CFTR* sequence analysis
☐ *CFTR* deletion/duplication analysis (*external DNA not accepted*)

Duchenne Muscular Dystrophy
☐ *DMD* deletion/duplication analysis (*external DNA not accepted*)
☐ *DMD* sequence analysis
☐ *DMD* mRNA analysis *(contact the laboratory before ordering)*

Fabry Disease
☐ *GLA* sequence analysis
☐ *GLA* deletion/duplication analysis (*external DNA not accepted*)
☐ *GLA* mRNA analysis *(contact the laboratory before ordering)*

Fragile X Syndrome & FMR1-related disorders
☐ Fragile X syndrome
☐ Fragile X-associated primary ovarian insufficiency
☐ Fragile X-associated tremor ataxia syndrome (FXTAS)

Fragile X E Syndrome ***
☐ *AFF2* trinucleotide repeat analysis *(See testing requirements)*

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Hereditary Hearing Loss *
Clinical information must be provided on pages 4 and 5
When the Common and Non-Syndromic Hearing Loss Panel is requested, STRC dosage is tested.
- ☐ Common and Non-Syndromic Hearing Loss panel
- ☐ Usher Syndrome panel
- ☐ Stickler Syndrome panel
- ☐ Alport Syndrome
- ☐ Norrie Syndrome, Treacher Collins Syndrome, Waardenburg Syndrome
- ☐ Deletion/duplication analysis

Hereditary Hemorrhagic Telangiectasia
- ☐ *ACVRL1* sequence analysis
- ☐ *ENG* sequence analysis
- ☐ *ACVRL1* and *ENG* deletion/duplication analysis (*external DNA not accepted*)
- ☐ *SMAD4* sequence analysis

Hereditary Spastic Paraplegia *
Clinical information must be provided on pages 4 and 5
- ☐ Comprehensive HSP (AR/AD/XL) panel including deletion/duplication analysis

Identity Testing
- ☐ Zygosity studies
- ☐ Maternal cell contamination studies *(maternal sample required)*

Neurofibromatosis type 1/Legius syndrome *
Clinical information must be provided on pages 4 and 5
- ☐ *NF1* sequence analysis
- ☐ *NF1* deletion/duplication analysis (*external DNA not accepted*)
- ☐ *SPRED1* sequence analysis
- ☐ *SPRED1* deletion/duplication analysis (*external DNA not accepted*)

Noonan Syndrome and RASopathies *
Clinical information must be provided on pages 4 and 5
- ☐ Noonan Syndrome and RASopathies panel
- ☐ Deletion/duplication analysis for *SPRED1* only (*external DNA not accepted*)

Prader-Willi Syndrome
- ☐ Methylation and deletion/duplication analysis (*external DNA not accepted*)
- ☐ UPD15 analysis *(parental samples required)*

Renal Diseases
- ☐ Atypical Hemolytic Uremic Syndrome / Membranoproliferative Glomerulonephritis sequence analysis
- ☐ Focal Segmental Glomerulosclerosis sequence analysis

Russell-Silver Syndrome
- ☐ IC1 methylation and 11p15 deletion/duplication analysis (*external DNA not accepted*)
- ☐ *UPD7* analysis *(parental samples required)*

Shwachman-Diamond Syndrome
- ☐ *SBDS* sequence analysis

Simpson-Golabi-Behmel Syndrome
- ☐ *GPC3* sequence analysis and *GPC3* and *GPC4* deletion/duplication analysis (*external DNA not accepted*)

Skeletal Dysplasia
- ☐ Achondroplasia (*FGFR3* recurrent mutation analysis)
- ☐ Hypochondroplasia (*FGFR3* recurrent mutation analysis)
- ☐ Thanatophoric Dysplasia (*FGFR3* recurrent mutation analysis)

Spinal and Bulbar Muscular Atrophy
- ☐ *AR* trinucleotide repeat analysis

Spinal Muscular Atrophy
- ☐ *SMN1* and *SMN2* deletion/duplication analysis (*external DNA not accepted*)

☐ X-Inactivation Analysis

☐ Other (*PRIOR APPROVAL REQUIRED; CONTACT LABORATORY*):

_____

_____

***Next-Generation Sequencing (NGS) testing will only be initiated if the clinical information sections (pages 4-5) are completed. For more information on our Next-Generation Sequencing (NGS) panels, including the list of genes tested, visit our website: www.sickkids.ca/en/care-services/for-health-care- providers/lab-testing-services**

**** For information on the testing algorithm for Cystic Fibrosis, visit https://www.sickkids.ca/en/care-services/for-health-care-providers/lab-tests/244-Cystic-Fibrosis/ on our website**

***** For information on the testing requirement for Fragile X E, visit the Specimen Requirements section for Fragile X E Syndrome on our website: www.sickkids.ca/en/care-services/for-health-care-providers/lab-tests/250-FRAXE**

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DISEASE SPECIFIC FEATURES

Autoinflammatory Disorders (RFS/AID/HLH/AGS)

- ☐ Abnormal inflammatory response
- ☐ Fevers
- ☐ Arthritis
- ☐ Pulmonary complications
- ☐ Gastrointestinal irritation
- ☐ Hepatosplenomegaly
- ☐ Lymphadenopathy
- ☐ Hemophagocytosis
- ☐ Oral ulcers
- ☐ Rash, specify:_____
- ☐ Ocular inflammation specify: _____
- ☐ Edema (periorbital, optic disk)
- ☐ Vision loss
- ☐ Other: _____

Hearing Loss

- ☐ Age of onset:_____
- ☐ Sensorineural hearing loss
- ☐ Conductive hearing loss
- ☐ Mixed hearing loss
- ☐ Bilateral ☐ Unilateral
- ☐ Syndromic ☐ Non-syndromic
- ☐ Ear anomalies ☐ Ear tags
- ☐ Eye anomalies ☐ Renal anomalies
- ☐ White forelock ☐ Cardiac anomalies
- ☐ Hirschsprung disease
- ☐ Other: _____

Hereditary Spastic Paraplegia (HSP)

- ☐ Abnormal corpus callosum
- ☐ Cognitive impairment
- ☐ Ataxia ☐ Spasticity
- ☐ Hyperreflexia ☐ Seizures
- ☐ Hypertonia ☐ Hypotonia
- ☐ Dystonia ☐ Dysarthria
- ☐ Extensor plantar reflex
- ☐ Other: _____

The following investigations are required before molecular testing of HSP is undertaken:

- ☐ MRI – Brain and spinal cord
- ☐ Biochemical testing - Vitamin B12, vitamin E, very long chain fatty acids, lysosomal work-up, plasma amino acids and serum lipoprotein analysis (as appropriate)

Neurofibromatosis type 1 (NF1) / Legius Syndrome

- ☐ **The patient meets the NIH criteria for a clinical diagnosis of NF1 (>2 of the clinical features below).**
- ☐ Café-au-lait macules
  - ☐ ≥6 CALS ☐ (#:_____)
- ☐ Neurofibromas, ≥ 2 or ≥ 1 Plexiform
- ☐ Freckling, axillary or inguinal
- ☐ Optic glioma
- ☐ ≥2 Lisch nodules (iris hamartomas)
- ☐ Osseous lesion (type:_____)
- ☐ First degree relative diagnosed with NF1 by above criteria
- ☐ Other: _____
- ☐ **The patient does not meet the NIH diagnostic criteria for NF1. Rationale for testing must be provided on page 5.**

Connective Tissue Disorders (CTD)

Ehlers Danlos Syndrome (EDS)

Indicate the suspected clinical diagnosis in the patient:

- ☐ Classic ☐ Vascular
- ☐ Kyphoscoliotic ☐ Other: _____

Note: Genetic testing is not offered for joint hypermobility alone. If testing is requested for joint hypermobility, provide rationale on page 5.
Check applicable CTD features below.

Osteopetrosis and Disorders of Increased Bone Density

Check applicable CTD features below.

CTD Related Clinical Features:

- ☐ Joint hypermobility: Beighton score: _____
- ☐ Arterial aneurysms, dissection or rupture
- ☐ Intestinal rupture
- ☐ Molluscoid pseudotumors
- ☐ Subcutaneous spheroids
- ☐ Loose/stretchable skin
- ☐ Smooth/velvety skin
- ☐ Widened atrophic scars

Osteogenesis Imperfecta (OI)

If the patient does not present with one of the test indications below, rationale for testing must be provided on page 5.

- ☐ Fetal findings on anatomy ultrasound consistent with OI.
- ☐ Fractures with minimal or no trauma in the absence of other known disorders of bone metabolism.
- ☐ Vertebral fractures
- ☐ Dentinogenesis imperfecta
- ☐ Low ALP for age/gender (ALPL gene analysis only will be performed – not eligible for full panel)

Check applicable CTD features below.

Bone Involvement

Check applicable CTD features below.

- ☐ Recurrent spontaneous tendon rupture
- ☐ Easy bruising
- ☐ Myopia
- ☐ Lens dislocation
- ☐ Blue/gray sclerae
- ☐ Thumb or wrist sign
- ☐ Club foot
- ☐ Scoliosis
- ☐ Marfanoid habitus
- ☐ Short stature
- ☐ Shortened long bones
- ☐ Recurrent pneumothoraces
- ☐ Joint subluxations/dislocations
- ☐ Fractures
- ☐ Bone deformity
- ☐ Wormian bones
- ☐ Increased bone mineral density
- ☐ Diaphyseal sclerosis
- ☐ Hearing loss
- ☐ Osteosclerosis
- ☐ Other: _____

Noonan Syndrome and RASopathies

- ☐ Increased nuchal translucency
- ☐ Developmental delay
- ☐ Characteristic facies
- ☐ Broad or webbed neck
- ☐ Heart defect (specify:_____)
- ☐ Hypertrophic cardiomyopathy
- ☐ Short stature (%ile:_____)
- ☐ Pectus deformity
- ☐ Lymphatic dysplasias
- ☐ Characteristic hematological abnormality (specify:_____)
- ☐ Other RASopathy features (specify:_____)
- ☐ **For postnatal patients: The patient must present with ≥ 2 of the above features for molecular testing to be undertaken.**

FAMILY HISTORY (Required)

Draw or attach a pedigree and provide any relevant information below, including clinical and family history details, as this is important for accurate interpretation of results.

Ethnicity: _____

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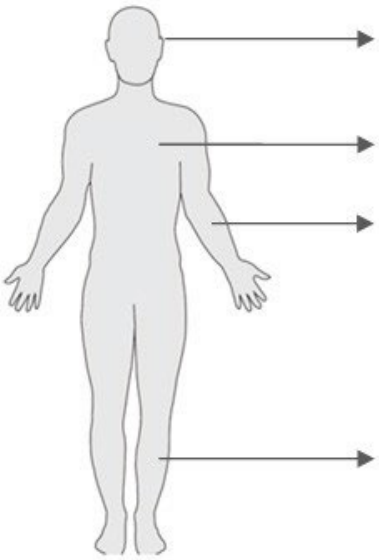
ADDITIONAL RELEVANT CLINICAL INFORMATION

Previous Genetic Testing

☐ No
☐ Yes – Test Results: _____



GENERAL CLINICAL FEATURES

Perinatal history
☐ Premature birth
☐ IUGR
☐ Oligohydramnios ☐ Polyhydramnios
☐ Other: _____

Growth
☐ Failure to thrive
☐ Growth retardation/short stature
☐ Overgrowth
☐ Macrocephaly ☐ Microcephaly
☐ Other: _____

Physical/cognitive development
☐ Delayed fine motor development
☐ Delayed gross motor development
☐ Delayed speech and language
☐ Autistic behavior
☐ Intellectual disability
☐ Developmental regression
☐ Other: _____

Behavioral
☐ Autistic features
☐ Obsessive-compulsive disorder
☐ Other psychiatric symptoms
☐ Other: _____

Cancer/Malignancy
☐ Age of onset: _____
☐ Tumor type: _____
☐ Location(s): _____

Craniofacial/Ophthalmalogic
☐ Abnormal face shape
☐ Blindness ☐ Cataracts
☐ Coloboma ☐ Optic atrophy
☐ Opthalmoplegia ☐ Ptosis
☐ Retinitis pigmentosa
☐ Oral cleft
☐ Other: _____

Brain malformations/abnormal imaging
☐ Abnormality of the basal ganglia
☐ Agenesis of the corpus callosum
☐ Brain atrophy
☐ Cortical dysplasia
☐ Hemimegalencephaly
☐ Heterotopia
☐ Holoprosencephaly
☐ Hydrocephalus
☐ Lissencephaly
☐ Periventricular leukomalacia
☐ Other: _____

Cardiac/congenital heart malformations
☐ ASD ☐ VSD
☐ Coarctation of aorta
☐ Hypoplastic left heart
☐ Tetralogy of Fallot
☐ Cardiomyopathy
☐ Arrhythmia/conduction defect
☐ Other: _____

Gastrointestinal
☐ Gastroschisis/omphalocele
☐ Gastrointestinal reflux
☐ Pyloric stenosis
☐ Tracheoesophageal fistula
☐ Hepatic failure
☐ Chronic intestinal pseudo-obstr.
☐ Hirschsprung disease
☐ Recurrent vomiting
☐ Chronic diarrhea
☐ Constipation
☐ Other: _____

Genitourinary abnormalities
☐ Ambiguous genitalia
☐ Cryptorchidism
☐ Hypospadias
☐ Hydronephrosis
☐ Kidney malformation
☐ Renal agenesis
☐ Proximal renal tubulopathy
☐ Other: _____

Endocrine
☐ Diabetes mellitus Type 1
☐ Diabetes mellitus Type 2
☐ Hypothyroidism
☐ Hypoparathyroidism
☐ Pheochromocytoma/paraganglioma
☐ Other: _____

Neurological/Muscular
☐ Ataxia ☐ Hypotonia
☐ Chorea ☐ Hypertonia
☐ Dystonia ☐ Spasticity
☐ Exercise intolerance/ easy fatigue
☐ Headache/migraine
☐ Muscle weakness
☐ Seizures (type: _____)
☐ Stroke/stroke-like episodes
☐ Other: _____

Skeletal/Limb abnormalities
☐ Contractures ☐ Club foot
☐ Polydactyly ☐ Syndactyly
☐ Vertebral anomaly ☐ Scoliosis
☐ Other: _____

Skin/Hair
☐ Abnormality of the hair pattern, quantity
☐ Abnormal nail growth
☐ Abnormal pigmentation
☐ Café-au-lait macules
☐ Neoplasms of the skin
☐ Neurofibromas
☐ Blistering
☐ Ichthyosis
☐ Other: _____

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BILLING FORM

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Completion of Billing Form NOT required for patients with an Ontario Health Card Number.

The hospital, referring laboratory, referring physician, or a patient/guardian will be billed for the services rendered, upon direction from the referring physician.

- Invoices are sent upon completion of each test/service.
- Invoices are itemized and include the date of service, patient name, CPT code, test name and charge.
- Contact SickKids' Genome Diagnostics Laboratory at 416-813-7200 x1 with billing inquiries.

How to complete the Billing Form:

- **Referring Physician completes the appropriate section below to specify billing method.**
- **Send requisition and completed "Billing Form" with specimen.**

Section 1: Complete to have the Healthcare Provider billed:

Referring Laboratory's Reference #:_____

Billing address of hospital, referring laboratory, clinic, referring physician, or medical group (if different from requisition):

Name: _____
Address: _____
City: _____ Prov/State: _____
Postal/Zip Code: _____ Country: _____
Contact Name: _____ Contact Telephone #: _____

Section 2: Complete to have Patient/Guardian billed directly:

If electing to have patient/guardian billed:

- *Patient/Guardian billing information below must be complete; otherwise, the healthcare provider will be billed.*
- *Advise the patient/guardian to expect a bill from the Genome Diagnostics laboratory.*
- *The patient's valid credit card information must be provided.*
- *Unfortunately, personal checks are not accepted as a method of payment.*
- ***In this case, the patient/guardian is solely responsible for the charges.***

Send bill to (check one): ☐ Patient ☐ Guardian

Method of Payment (check one): ☐ American Express ☐ MasterCard ☐ Visa

Name as it appears on credit card: _____

Credit card #: _____ CVS #: _____

Expiry date on credit card: _____

Signature of credit card holder (Required): _____

Mailing Address of Patient/Guardian (if different from requisition):

Name: _____
Address: _____
_____ Apt. #: _____
City: _____ Prov/State: _____
Postal/Zip Code: _____ Country: _____

Additional Contact Information

Patient's phone # with area code:

_____

-or-

Guardian's phone # with area code:

_____

SickKids®
THE HOSPITAL FOR SICK CHILDREN
Paediatric Laboratory Medicine

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(CLIA # 99D1014032)

Patient Name:
Preferred Name (if different):
Date of Birth (DD/MM/YYYY):
Legal Sex: ◯ *Male* ◯ *Female* ◯ *Non-binary/U/X*
Sex Assigned at Birth (if different): ◯ *Male* ◯ *Female* ◯ *Unassigned*
Gender Identity): ◯ *Male* ◯ *Female* ◯ *Non-binary/U/X*
MRN:
Parent's Name:
Address:

For Canada Only
Provincial Health Card #: *Version:*
Issuing Province:

Genome Diagnostics

LIST OF GENES BY TEST

Autoinflammatory Diseases (sequencing and dosage available)

Recurrent Fever Syndromes panel (5 genes)
MEFV, MVK, NLRP12, NLRP3, TNFRSF1A

Autoinflammatory Diseases panel (25 genes)
ARPC1B, CARD14, CDC42, CECR1 (ADA2), COPA, ELANE, IL1RN, IL36RN, LACC1, LPIN2, NLRC4, NOD2, OTULIN, PLCG2, POMP, PSMB8, PSTPIP1, RAB27A, RBCK1, RIPK1, SH3BP2, SLC29A3, TMEM173 (STING1), TNFAIP3, TRNT1

Hemophagocytic Lymphohistiocytosis panel (16 genes)
AP3B1, BLOC1S6, CD27, ITK, LYST, NLRC4, PRF1, CD70, RAB27A, SH2D1A, SLC7A7, STX11, STXBP2, UNC13D, XIAP, MAGT1

Aicardi-Goutières Syndrome panel (7 genes)
ADAR, IFIH1, RNASEH2A, RNASEH2B, RNASEH2C, SAMHD1, TREX1

Connective Tissue Disease (sequencing and dosage available)

Ehlers Danlos Syndrome panel (22 genes)
ACTA2, ADAMTS2, ATP7A, B4GALT7 (**no dosage**), CHST14, COL3A1, COL5A1, COL5A2, COL1A1, COL1A2, DSE, FBN2, FKBP14, PLOD1, PRDM5, SLC39A13, SMAD3, TGFB2, TGFBR1, TNXB, TGFBR2, ZNF469

Osteogenesis Imperfecta panel (20 genes)
ALPL, BMP1, COL1A1, COL1A2, CRTAP, FKBP10, IFITM5, LRP5, MBTPS2, P3H1, PLOD2, PLS3, PPIB, SERPINF1, SERPINH1, SP7, SPARC, TMEM38B, WNT1, XYLT2

Osteopetrosis and Disorders of Increased Bone Density panel (10 genes)
CA2, CLCN7, LRP5, OSTM1, PLEKHM1, SNX10, TCIRG1, TNFRSF11A, TNFRSF11B, TNFSF11

Bone Involvement panel (40 genes)
ARSL, CBS, COL11A1, COL11A2, COL2A1, COL9A1, COL9A2, COL9A3, COMP, DDR2, DYM, EBP, EIF2AK3, FBN1, FBN2, FGFR3, FLNB, HSPG2, IFT122, IFT43, IFT80, LBR, LIFR, MATN3, NEK1, NKX3-2, NSDHL, PEX7, PTH1R, SHOX, SLC26A2, SLC35D1, SLC39A13, SOX9, TRAPPC2, TRIP11, TRPV4, TTC21B, WDR19, WDR35

Hereditary Hearing Loss (sequencing and dosage available)

Common and Non-Syndromic Hearing Loss panel (61 genes)
ACTG1, ADGRV1, CDH23, CHD7, CIB2, CLDN14, COCH, DFNA5, DFNB59, DIAPH1, ESPN, ESRRB, EYA1, EYA4, GJB2, GJB6, GIPC3, GPSM2, GRHL2, GRXCR1, HGF, ILDR1, KCNQ1, KCNQ4, KCNE1, LHFPL5, LOXHD1, LRTOMT, MARVELD2, MYH14, MYH9, MYO15A, MYO3A, MYO6, MYO7A, OTOA, OTOF, OTOG, OTOGL, PCDH15, PDZD7, POU3F4, POU4F3, PRPS1, PTPRQ, RDX, SERPINB6, SIX1, SLC17A8, SLC26A4, SMPX, STRC, TECTA, TMC1, TMIE, TMPRSS3, TPRN, TRIOBP, USH2A, WFS1, WHRN

Usher Syndrome panel (11 genes)
ADGRV1, CDH23, CIB2, CLRN1, MYO7A, PCDH15, PDZD7, USH1C, USH1G, USH2A, WHRN

Stickler Syndrome panel (5 genes)
COL11A1, COL11A2, COL2A1, COL9A1, COL9A2

Alport Syndrome panel (3 genes)
COL4A3, COL4A4, COL4A5

Syndromic Hearing Loss - Treacher Collins syndrome, Waardenburg syndrome, Norrie syndrome panel (7 genes)
EDN3, EDNRB, MITF, NDP, PAX3, SOX10, TCOF1

Hereditary Spastic Paraplegia (sequencing and dosage available)

Comprehensive HSP (AR/AD/XL) panel (67 genes)
ABCD1, ADAR, ALDH18A1, ALS2, AP4B1, AP4E1, AP4M1, AP4S1, AP5Z1, ATL1, ATP13A2, B4GALNT1, BSCL2, C19orf12, CAPN1, CPT1C, CYP2U1, CYP7B1, DDHD1, DDHD2, ERLIN1, ERLIN2, FA2H, FAR1, FARS2, GBA2, HACE1, HPDL, HSPD1, IBA57, IFIH1, KIDINS220, KIF1A, KIF1C, KIF5A, L1CAM, MAG, MTRFR, NIPA1, NT5C2, PCYT2, PLP1, PNPLA6, POLG, POLR3A, POLR3B, REEP1, REEP2, RNF170, RTN2, SACS, SELENOI, SETX, SLC16A2, SPART, SPAST, SPG11, SPG21, SPG7, TECPR2, TFG, TUBB4A, UBAP1, UCHL1, VPS13D, WASHC5, ZFYVE26

Noonan Syndrome (sequencing only) (15 genes)

BRAF, CBL, HRAS, KRAS, LZTR1, MAP2K1, MAP2K2, NRAS, PTPN11, RAF1, RIT1, SHOC2, SOS1, SOS2, SPRED1-**Dosage ONLY for SPRED1**