Genetic Resource Centre Established Testing Menu

For testing that has more than one lab listed, please choose your preferred test/lab based on gene content

To search for a test, click "Ctrl-F" on the computer keyboard and type in part of the test name. Click "enter" until you find the test of interest.

GRC Contact Information Phone: 403-955-5400 Email: grc@albertahealthservices.ca Fax: 403-592-4238	**Ordering restrictions** - Carrier testing/presymptomatic testing is currently restricted to Clinical Genetics. Testing for symptomatic patients may be restricted based on clinical specialty. Please contact the Genetic Resource Centre if you have questions regarding ordering restrictions.	**How to use an online portal?** For assistance, please contact: Blueprint Genetics: Allison Sluyters (allison.sluyters@blueprintgenetics.com) Centogene/Life Labs: Genetic.Counsellors@lifelabs.com GeneDx: Sarah Waltho (swaltho@genedx.com) Invitae: Hana Sroka (hana.sroka@invitae.com) Prevention Genetics: Christèle du Souich (christele.dusouich@preventiongenetics.com)

Indication / Suspected Diagnosis	Test Name	Laboratory	Link to Online Portal or Requisition	Notes
Add-on testing				
Any NGS panel	Flex Testing	Blueprint Genetics	Online Portal	Can add up to 200 additional genes to Blueprint Genetics panel, free of charge
Site Specific Testing - PLEASE READ NOTES SECTION BEFORE SELECTING LAB				
Site-specific testing	Familial Variant Testing/Targeted Variant Testing	Blueprint Genetics	Online Portal	Select this lab if they performed testing for the index patient, if the lab who tested the index patient is not listed below, or if you are requesting testing to confirm a research variant. Blueprint Genetics cannot perform targeted testing for copy number variants or prenatal targeted testing if they did not test the index patient. Contact the GRC if you have any questions.
		Exeter Genomics Laboratory	Diabetes Genetic Test Referral Forms Click on "MODY Request Form"	Select this lab if they performed testing for the index patient (MODY testing)

Site-specific testing (continued)	Familial Variant Testing/Targeted Variant Testing (continued)		Hyperinsulinism Web Page Click on "Request Form"	Select this lab if they performed testing for the index patient (Congenital Hyperinsulinism testing)
		Invitae	Online Portal	Select this lab if they performed testing for the index patient
		Molecular Otolaryngology & Renal Research Laboratories	Requisition form	Select this lab if they performed testing for the index patient
		Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Select this lab if they performed testing for the index patient
Cancer/Tumour/Overgrowth				
Colon cancer	ColoSeq™ Tumor Panel	University of Washington	Website	Click on “ordering & collection” to access the link for the requisition form
Colon cancer	ColoSeq™ Tumor Single Gene	University of Washington	Website	Click on “ordering & collection” to access the link for the requisition form
Megalencephaly	Megalencephaly Panel	University of Washington	Website	Click on “ordering & collection” to access the link for the requisition form
Somatic overgrowth	Somatic Overgrowth Gene Set	Washington University (St. Louis, MO)	Requisition form	

Uveal melanoma	DecisionDx®-UM	Castle Biosciences	Requisition form	
Cardiology				
Arrhythmias, cardiomyopathy, structural heart defects	Any cardiac panel	Blueprint Genetics	Online Portal	Some of these panels include analysis of the mitochondrial genome. Visit www.blueprintgenetics.com for more information
Post-mortem cardiac testing	Arrhythmia and Cardiomyopathy Comprehensive Panel	Invitae	Online Portal	
	Any cardiac panel	Blueprint Genetics	Online Portal	Some of these panels include analysis of the mitochondrial genome. Visit www.blueprintgenetics.com for more information
Dermatology				
Albinism	Albinism Panel	Blueprint Genetics	Online Portal	
Ectodermal dysplasia	Ectodermal Dysplasia Panel	Blueprint Genetics	Online Portal	
Ichthyosis	Ichthyosis Panel	Blueprint Genetics	Online Portal	
Pseudoxanthoma elasticum	*ABCC6* single gene test	Blueprint Genetics	Online Portal	
Dysmorphology				
CHARGE syndrome	*CHD7* single gene test	Blueprint Genetics	Online Portal	
Cleft lip/palate	Cleft Lip/Palate and Associated Syndromes Panel	Blueprint Genetics	Online Portal	
Facial dysostosis	Facial Dysostosis and Related Disorders Panel	Blueprint Genetics	Online Portal	Also listed under Skeletal heading
Kabuki Syndrome	Kabuki Syndrome Panel	Blueprint Genetics	Online Portal	
Rasopathy Disorders	Noonan Syndrome Panel	Blueprint Genetics	Online Portal	
	Noonan Spectrum Disorders/RASopathies Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	
van der Woude syndrome	Invitae van der Woude Syndrome Panel	Invitae	Online Portal	

Ear, Nose & Throat				
Nonsyndromic sensorineural hearing loss	Comprehensive Hearing Loss and Deafness Panel (or sub-panel)	Blueprint Genetics	Online Portal	The Blueprint Genetics hearing loss panels include analysis of the mitochondrial genome In-house testing must be performed first, if relevant: *GJB2, GJB6,* and *MT-RNR1* targeted analysis (Edmonton Molecular Genetics Lab)
	OtoSCOPE® Panel	Molecular Otolaryngology & Renal Research Laboratories	Requisition form	
Waardenburg syndrome	Waardenburg Syndrome Panel	Blueprint Genetics	Online Portal	
Endocrinology				
Congenital adrenal hyperplasia	Congenital Adrenal Hyperplasia Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: *CYP21A2* sequencing and MLPA *CYP11B1* sequencing (Calgary Molecular Genetics Lab)
Congenital hyperinsulinism	Congenital Hyperinsulinism Panel	Exeter Genomics Laboratory	Hyperinsulinism Web Page Click on "Request Form"	
Diabetes insipidus	Diabetes Insipidus Panel	Blueprint Genetics	Online Portal	
Disorders of sexual development	Abnormal Genitalia / Disorders of Sexual Development Panel	Blueprint Genetics	Online Portal	
Familial hyperaldosteronism	*CYP11B1/CYP11B2* Fusion Gene	Center for Nephrology and Metabolic Disorders	Requisition form Consent form	For non-urgent cases where both fusion gene testing and panel testing are indicated, testing should be performed stepwise (requires two GRC funding requests)
	Primary Aldosteronism Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	

Familial hyperparathyroidism	Hyperparathyroidism Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: Endocrine Disorders Panel - *AIP, CASR, CDC73, CDKN1B, MEN1, PRKAR1A, RET* (Calgary Molecular Genetics Lab)
Familial hypocalciuric hypercalcemia	Familial Hypocalciuric Hypercalcemia (FHH) Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	In-house testing must be performed first, if relevant: *CASR* sequencing and del/dup by NGS (Calgary Molecular Genetics Lab)
GNAS-related disorders	*GNAS* – Methylation and Del/Dup analysis	Genome Diagnostics, Amsterdam UMC	Requisition form	Test includes sanger sequencing
Hypogonadotropic hypogonadism/Kallmann syndrome	Kallmann Syndrome Panel	Blueprint Genetics	Online Portal	
	Hypogonadotropic Hypogonadism/Kallmann Syndrome Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	
Hypophosphatemic rickets	Hypophosphatemic Rickets Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: *ALPL* sequencing and del/dup by NGS (Calgary Molecular Genetics Lab)
	PHEX single gene test	Blueprint Genetics	Online Portal	
Liddle syndrome	Liddle Syndrome Panel	Blueprint Genetics	Online Portal	

Mature onset diabetes of the young (MODY)	GCK sequencing and/or MODY Panel	Exeter Genomics Laboratory	Diabetes Genetic Test Referral Forms Click on "MODY Request Form"	Please order reflex testing, if indicated. Check off: 1) GCK sequencing 2) Next generation sequencing 34 gene test only if GCK or HNF1A/4A testing performed by the Exeter Laboratory is negative
	MODY Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
Monogenic obesity	Monogenic Obesity Panel	Blueprint Genetics	Online Portal	
Thyroid hormone resistance	*THRB* single gene test	Blueprint Genetics	Online Portal	
Gastroenterology				
Cholestasis	Cholestasis Panel	Blueprint Genetics	Online Portal	
Congenital diarrhea	Congenital Diarrhea Panel	Blueprint Genetics	Online Portal	
Hirschsprung disease	Hirschsprung Disease Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: *RET* and *PHOX2B* sequencing and del/dup by NGS (Calgary Molecular Genetics Lab)
Pancreatitis	Pancreatitis Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: *PRSS1* and *CFTR* targeted sequence analysis (Edmonton Molecular Genetics Lab)
Polycystic liver disease	Polycystic Liver Disease Panel	Blueprint Genetics	Online Portal	
Hematology				
Anemia	Anemia Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: *HBA1/HBA2* multiplex PCR and sequencing *HBB* sequencing and MLPA (Calgary Molecular Genetics Lab)

Bone marrow failure	Bone Marrow Failure Syndrome Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: Fanconi Anemia and DNA Repair Disorders Panel - *ATM, BLM, BRCA2, BRIP1, FANCA, FANCB, FANCC, FANCD2, FANCE, FANCF, FANCG, FANCI, FANCL, FANCM, NBN, PALB2, RAD51C, REQL4, SLX4* (Edmonton Molecular Genetics Lab) *SBDS* sequencing and del/dup by NGS (Edmonton Molecular Genetics Lab)
Congenital neutropenia	Congenital Neutropenia Panel	Blueprint Genetics	Online Portal	
	Invitae Severe Congenital Neutropenia Panel	Invitae	Online Portal	
Diamond-Blackfan anemia	Diamond-Blackfan Anemia Panel	Blueprint Genetics	Online Portal	
Hereditary angioedema	*F12* Gene Testing in Hereditary Angioedema Type III	GeneDx	Online Portal	
Hereditary leukemia	Hereditary Leukemia Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: Familial Acute Myeloid Leukemia Panel – *CEBPA, GATA2, RUN1* (Edmonton Molecular Genetics Lab) For hematological malignancies**,** the ideal tissue type for germline testing is fibroblast culture. However, buccal samples will be accepted to allow for a rapid turnaround time if urgently required for clinical management.
Hereditary spherocytosis	Red Blood Cell Membrane Disorder Panel	Blueprint Genetics	Online Portal	
Thrombocytopenia	Thrombocytopenia Panel	Blueprint Genetics	Online Portal	

Immunology				
Complement system disorder	Complement System Disorder Panel	Blueprint Genetics	Online Portal	
Familial Mediterranean fever	*MEFV* sanger sequencing	BC Molecular Genetics Laboratory	Requisition form	
Hemophagocytic lymphohistiocytosis	Hemophagocytic Lymphohistiocytosis Panel	Blueprint Genetics	Online Portal	
Periodic fever syndrome	Autoinflammatory Syndrome Panel	Blueprint Genetics	Online Portal	
Primary immunodeficiency	Primary Immunodeficiency Panel	Blueprint Genetics	Online Portal	
Primary immunodeficiency and primary ciliary dyskinesia	Primary Immunodeficiency (PID) and Primary Ciliary Dyskinesia (PCD) Panel	Blueprint Genetics	Online Portal	See Pulmonology section for Primary Ciliary Dyskinesia Panel
Severe combined immunodeficiency	Severe Combined Immunodeficiency Panel	Blueprint Genetics	Online Portal	
Metabolic				
3-Hydroxy-3-methylglutaryl-CoA lyase deficiency	*HMGCL* single gene test	Blueprint Genetics	Online Portal	
	Invitae Elevated C5-OH Panel	Invitae	Online Portal	Panel includes genes for secondary newborn screening targets (*MCCC1, MCCC2, AUH, TAZ, OPA3, SERAC1, DNAJC19, ACAT1, HSD17B10*)
Biotinidase deficiency	*BTD* single gene test	Blueprint Genetics	Online Portal	
	BTD gene sequencing	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Use test code 520 for sanger sequencing
Carnitine deficiency, systemic primary	*SLC22A5* single gene test	Blueprint Genetics	Online Portal	

Citrullinemia	Invitae Elevated Citrulline Panel	Invitae	Online Portal	
Fabry disease	*GLA* single gene test	Blueprint Genetics	Online Portal	
Familial Hypercholesterolemia	Hyperlipidemia Core Panel	Blueprint Genetics	Online Portal	
	Hyperlipidemia Panel	Blueprint Genetics	Online Portal	
Galactosemia	*GALT* single gene test	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant (**This does not apply to urgent cases**): *GALT* targeted sequence analysis (Calgary Molecular Genetics Lab)
Glutaric acidemia type 1	*GCDH* single gene test	Blueprint Genetics	Online Portal	
	Invitae Glutaric Acidemia Type I Test	Invitae	Online Portal	Can add on genes for secondary targets (*ETFA, ETFB, ETFDH*) or reflex these genes free of charge within 90 days *SUGCT* is not currently available at Invitae, but can be ordered as part of the Blueprint Genetics "Organic Acidemia/Aciduria & Cobalamin Deficiency Panel" if patient has a broad differential diagnosis
Glycogen storage disorder	Glycogen storage disorder panel	Blueprint Genetics	Online Portal	
Hypoglycemia	Hypoglycemia, Hyperinsulinism and Ketone Metabolism Panel	Blueprint Genetics	Online Portal	
	Metabolic Hypoglycemia Sequencing Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Patients with congenital hyperinsulinism are not good candidates for this panel
Isovaleric acidemia	*IVD* single gene testing	Blueprint Genetics	Online Portal	

Fatty Acid Oxidation Syndrome	Fatty Acid Oxidation Syndrome Panel	Blueprint Genetics	Online Portal	
	Invitae Fatty Acid Oxidation Defects Panel	Invitae	Online Portal	
Leukodystrophy	Leukodystrophy and Leucoencephalopathy panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome Also listed under Neurology heading
Long chain 3-hydroxyacyl-CoA dehydrogenase deficiency	Invitae Elevated C16-OH, C16:1-OH, C18-OH and C18:1-OH Panel	Invitae	Online Portal	
Maple syrup urine disease	Maple Syrup Urine Disease Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	
Medium chain acyl-CoA dehydrogenase deficiency	*ACADM* single gene test	Blueprint Genetics	Online Portal	
	ACADM gene sequencing	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Use test code 180 for sanger sequencing In-house testing must be performed first, if relevant (**This does not apply to urgent cases**): *ACADM* targeted sequence analysis (Edmonton Molecular Genetics Lab)
	Invitae Medium Chain Acyl-CoA Dehydrogenase Deficiency Test	Invitae	Online Portal	Can add on genes for secondary newborn screening targets (*ETFA, ETFB, ETFDH*) or reflex these genes free of charge within 90 days
Metabolic Myopathy and Rhabdomyolysis	Metabolic Myopathy and Rhabdomyolysis Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
Methylmalonic acidemia	Invitae Methylmalonic Acidemia Panel	Invitae	Online Portal	Can add on combined methylmalonic acidemia and homocystinuria genes

Mitochondrial DNA Depletion Syndrome	Mitochondrial DNA Depletion Syndrome Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
Niemann-Pick disease type C	Niemann-Pick Disease Type C Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	
Organic Acidemia	Organic Acidemia/Aciduria & Cobalamin Deficiency Panel	Blueprint Genetics	Online Portal	
	Invitae Organic Acidemias Panel	Invitae	Online Portal	
Phenylketonuria	Hyperphenylalaninemia panel	Blueprint Genetics	Online Portal	
	PAH single gene test	Invitae	Online Portal	Can add on additional hyperphenylalaninemia genes or reflex these genes free of charge within 90 days
Pompe disease	*GAA* single gene test	Blueprint Genetics	Online Portal	
Porphyria	Porphyria Panel	Blueprint Genetics	Online Portal	
Propionic acidemia	Invitae Propionic Acidemia Panel	Invitae	Online Portal	
Tri-functional protein deficiency	Invitae Elevated C16-OH, C16:1-OH, C18-OH and C18:1-OH Panel	Invitae	Online Portal	
Tyrosinemia, type 1	*FAH* single gene test	Blueprint Genetics	Online Portal	
	Tyrosinemia Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Includes genes for secondary newborn screening targets (*GSTZ1, TAT, HPD*)

Very long chain acyl-CoA dehydrogenase deficiency	*ACADVL* single gene test	Blueprint Genetics	Online Portal	
	ACADVL gene testing	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Use test code 184 for sanger sequencing
	Invitae Very Long Chain Acyl-CoA Dehydrogenase Deficiency Test	Invitae	Online Portal	Can add on genes for secondary newborn screening targets (*ETFA, ETFB, ETFDH, CPT2, SLC25A20*) or reflex these genes free of charge within 90 days
Wilson Disease	*ATP7B* single gene test	Blueprint Genetics	Online Portal	
Nephrology				
Adult tubulointerstitial kidney disease	Renal Malformation Panel + *UMOD* (flex)	Blueprint Genetics	Online Portal	
Alport syndrome	Alport Syndrome Panel	Blueprint Genetics	Online Portal	
Atypical hemolytic uremic syndrome	Hemolytic Uremic Syndrome Panel	Blueprint Genetics	Online Portal	
Bartter syndrome	Bartter Syndrome Panel	Blueprint Genetics	Online Portal	
Ciliopathies	Ciliopathy Panel	Blueprint Genetics	Online Portal	
Congenital Abnormalities of the Kidney and Urinary Tract	Congenital Abnormalities of the Kidney and Urinary Tract (CAKUT) Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	

Cystinosis	*CTNS* Gene, 57-kb Deletion and Sequencing	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Dosage analysis (57 kb deletion) reflex sequencing for routine cases
Focal segmental glomerulosclerosis, Thin basement membrane disease	Any nephrology panel	Blueprint Genetics	Online Portal	
Gitelman syndrome	Bartter Syndrome Panel + *HNF1B* (flex)	Blueprint Genetics	Online Portal	
Nephrolithiasis	Nephronophthisis Panel	Blueprint Genetics	Online Portal	
Nephrotic Syndrome	Nephrotic Syndrome Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
Polycystic kidney disease, adult onset	Polycystic Kidney Disease Panel	Blueprint Genetics	Online Portal	
Polycystic kidneys; young adult/child	Cystic Kidney Disease Panel	Blueprint genetics	Online Portal	
Neurology				
Amyotrophic lateral sclerosis	Amyotrophic Lateral Sclerosis Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: *SOD1* and *C9orf72* targeted sequence analysis (Edmonton Molecular Genetics Lab)
Ataxia	Ataxia Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome In-house testing must be performed first, if relevant: Spinocerebellar ataxia types 1, 2, 3, 6, 7, 8, and 17, DRPLA, Friedreich ataxia, Fragile X-associated tremor/ataxia syndrome (Calgary Molecular Genetics Lab)

Autism/Intellectual Disability	Autism Spectrum Disorders and Intellectual Disability (ASD-ID) Comprehensive Panel (Patient Plus)	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	**Parental blood samples are required for this test, and should be collected at the time you submit your funding request.** The test type for this order is Patient Plus (test code 5201).
Charcot-Marie-Tooth	Charcot-Marie-Tooth Neuropathy Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: Charcot-Marie-Tooth disease types 1A, 1B, and X (Calgary Molecular Genetics Lab)
Congenital myasthenic syndrome	Congenital Myasthenic Syndromes Panel	Blueprint Genetics	Online Portal	
Dementia	Dementia Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
	Alzheimer Disease, Familial, Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	
Dystonia	Dystonia Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
Epilepsy	Comprehensive Epilepsy Panel (or sub-panel)	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
	Epilepsy and Seizure Plus Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	
	Comprehensive Epilepsy and Seizure Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	

Facioscapulohumeral muscular dystrophy, type 2	*SMCHD1* single gene test	Blueprint Genetics	Online Portal	Testing for Facioscapulohumeral muscular dystrophy type 1 must be performed first, if relevant. This testing is performed at the Children's Hospital of Eastern Ontario, and funded by the Calgary Molecular Genetics Lab. Please contact the lab genetic counsellors at 403-955-3097 with any questions.
Familial hemiplegic migraine	Migraine Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
Hereditary spastic paraplegia	Spastic Paraplegia Panel	Blueprint Genetics	Online Portal	
Holoprosencephaly	Holoprosencephaly Panel	Blueprint Genetics	Online Portal	
Leukodystrophy	Leukodystrophy and Leucoencephalopathy panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome Also listed under Metabolic heading
Limb girdle muscular dystrophy	*DYSF* single gene test	Blueprint Genetics	Online Portal	
Malignant hyperthermia	Malignant Hyperthermia Susceptibility Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	
Microcephaly and/or pontocerebellar hypoplasia	Microcephaly and Pontocerebellar Hypoplasia Panel	Blueprint Genetics	Online Portal	
Muscular dystrophy/myopathy	Comprehensive Muscular Dystrophy / Myopathy Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
Myotonia and paramyotonia congenita	Invitae Myotonia and Paramyotonia Congenita Panel	Invitae	Online Portal	
Myotonia congenita	*CLCN1* single gene test	Blueprint Genetics	Online Portal	
Neuronal migration disorders	Neuronal Migration Disorder Panel	Blueprint Genetics	Online Portal	

Parkinson disease/parkinsonism	Parkinson Disease Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
	Parkinson Disease and Parkinsonism Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	
	Parkinson Disease Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	
Periodic paralysis	Periodic Paralysis Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: *CACNA1S* and *SCN4A* targeted sequence analysis (Edmonton Molecular Genetics Lab)
Spinal muscular atrophy	Spinal Muscular Atrophy Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: *SMN1* MLPA (Calgary Molecular Genetics Lab)
Spinocerebellar ataxia	Repeat expansion (*ATXN10, BEAN1, NOP56, PPP2R2B*)	Centogene/LifeLabs	Online Portal	In-house testing must be performed first, if relevant: Spinocerebellar ataxia types 1, 2, 3, 6, 7, 8, and 17 (Calgary Molecular Genetics Lab)
Ophthalmology				
Cone rod dystrophy	Cone Rod Dystrophy Panel	Blueprint Genetics	Online Portal	
Congenital cataracts	Cataract Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
Congenital stationary night blindness	Congenital Stationary Night Blindness Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: *CACNA1F* targeted sequence analysis (Calgary Molecular Genetics Lab)
Ectopic Lentis	Ectopia Lentis Panel	Blueprint Genetics	Online Portal	
Leber congenital amaurosis	Leber Congenital Amaurosis Panel	Blueprint Genetics	Online Portal	
Macular dystrophy	Macular Dystrophy Panel	Blueprint Genetics	Online Portal	
Microphthalmia/anopthalmia	Microphthalmia, Anophthalmia and Anterior Segment Dysgenesis Panel	Blueprint Genetics	Online Portal	
Neuro-Ophthalmology Panel	Neuro-Ophthalmology Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome

Optic atrophy	Optic Atrophy Panel	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant: mtDNA testing for Leber hereditary optic neuropathy (Edmonton Molecular Genetics Lab)
Retinal dystrophy	Retinal Dystrophy Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
Retinitis pigmentosa	Retinitis Pigmentosa Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
Retinitis pigmentosa, x-linked	Sequencing of the Mutational Hotspot RPGR (isoform C) ORF15 Region	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	
Septo-Optic Dysplasia	Septo-Optic Dysplasia Panel	Blueprint Genetics	Online Portal	
Usher syndrome	Usher Syndrome Panel	Blueprint Genetics	Online Portal	
Prenatal				
Arthrogryposis	Arthrogryposes Panel	Blueprint Genetics	Online Portal	**A maternal blood sample is required for maternal cell contamination studies** to be performed at either the Calgary or Edmonton Molecular Genetics Lab. Also listed under Skeletal heading
Ashkenazi Jewish carrier testing	Ashkenazi Jewish Carrier Screening	Blueprint Genetics	Online Portal	*HEXA* (c.805G>A, c.1278insTATC, c.1421+1G>C) *ASPA* (c.693C>A, c.854A>C) *IKBKAP* (c.2204+6T>C, c.2087G>C) *FANCC* c.456+4A>T

Familial variant(s)	Prenatal targeted testing	Blueprint Genetics	Online Portal	Select this lab if they performed testing for the index patient **A maternal blood sample is required for maternal cell contamination studies** to be performed at either the Calgary or Edmonton Molecular Genetics Lab.
		Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Select this lab if they performed testing for the index patient OR if the index patient was tested at any other lab (coordinate a positive control sample to be sent to Prevention Genetics) **A maternal blood sample is required for maternal cell contamination studies** to be performed at Prevention Genetics
Fetal aneuploidy	Harmony Prenatal Test	Dynacare	Requisition form	For fetal aneuploidy testing, only funded in cases where a invasive procedure is contraindicated and pregnancy is at increased risk of aneuploidy
Fetal sex determination	Harmony Prenatal Test	Dynacare	Requisition form	For cases where there is a family history of an x-linked condition and fetal sexing results may prevent the need for an invasive procedure
Increased nuchal translucency or nuchal fold	Noonan Spectrum Disorders/RASopathies Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	**A maternal blood sample is required for maternal cell contamination studies** to be performed at Prevention Genetics
Non-immune hydrops	Non-Immune Hydrops Fetalis Panel	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	**A maternal blood sample is required for maternal cell contamination studies** to be performed at Prevention Genetics

Skeletal dysplasia	Comprehensive Growth Disorders / Skeletal Dysplasias and Disorders Panel (or sub-panel)	Blueprint Genetics	Online Portal	**A maternal blood sample is required for maternal cell contamination studies** to be performed at either the Calgary or Edmonton Molecular Genetics Lab Also listed under Skeletal heading
Pulmonology				
Inherited pulmonary disease	Comprehensive Pulmonology Panel	Blueprint Genetics	Online Portal	This test includes analysis of the mitochondrial genome
Interstitial pulmonary disease	Interstitial Lung Disease Panel	Blueprint Genetics	Online Portal	
Neonatal respiratory distress	Neonatal Respiratory Distress – Surfactant Dysfunction Panel	Blueprint Genetics	Online Portal	
Primary ciliary dyskinesia	Primary Ciliary Dyskinesia Panel	Blueprint Genetics	Online Portal	
Single Genes				
3-Hydroxy-3-methylglutaryl-CoA lyase deficiency	*HMGCL* single gene test	Blueprint Genetics	Online Portal	
Biotinidase deficiency	*BTD* single gene test	Blueprint Genetics	Online Portal	
	BTD gene sequencing	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Use test code 520 for sanger sequencing
Carnitine deficiency, systemic primary	*SLC22A5* single gene test	Blueprint Genetics	Online Portal	
CHARGE syndrome	*CHD7* single gene test	Blueprint Genetics	Online Portal	

Congenital hypothyroidism	*TRH* Targeted Del/Dup	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	This test is designed to detect a *TRH* deletion in the Hutterite population
Cystinosis	*CTNS* Gene, 57-kb Deletion and Sequencing	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Dosage analysis (57 kb deletion) reflex sequencing for routine cases
Fabry disease	*GLA* single gene test	Blueprint Genetics	Online Portal	
Facioscapulohumeral muscular dystrophy, type 2	*SMCHD1* single gene test	Blueprint Genetics	Online Portal	Testing for Facioscapulohumeral muscular dystrophy type 1 must be performed first, if relevant. This testing is performed at the Children's Hospital of Eastern Ontario, and funded by the Calgary Molecular Genetics Lab. Please contact the lab genetic counsellors at 403-955-3097 with any questions.
Familial Mediterranean fever	*MEFV* sanger sequencing	BC Molecular Genetics Laboratory	Requisition form	
Galactosemia	*GALT* single gene test	Blueprint Genetics	Online Portal	In-house testing must be performed first, if relevant (**This does not apply to urgent cases**): *GALT* targeted sequence analysis (Calgary Molecular Genetics Lab)
Glutaric acidemia type 1	*GCDH* single gene test	Blueprint Genetics	Online Portal	
Hypophosphatemic rickets	*PHEX* single gene test	Blueprint Genetics	Online Portal	
Isovaleric acidemia	*IVD* single gene testing	Blueprint Genetics	Online Portal	

Limb girdle muscular dystrophy	*DYSF* single gene test	Blueprint Genetics	Online Portal	
Medium chain acyl-CoA dehydrogenase deficiency	*ACADM* single gene test	Blueprint Genetics	Online Portal	
	ACADM gene sequencing	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Use test code 180 for sanger sequencing In-house testing must be performed first, if relevant (**This does not apply to urgent cases**): *ACADM* targeted sequence analysis (Edmonton Molecular Genetics Lab)
Myotonia congenita	*CLCN1* single gene test	Blueprint Genetics	Online Portal	
Phenylketonuria	*PAH* single gene test	Invitae	Online Portal	Can add on additional hyperphenylalaninemia genes or reflex these genes free of charge within 90 days
Pompe disease	*GAA* single gene test	Blueprint Genetics	Online Portal	
Pseudoxanthoma elasticum	*ABCC6* single gene test	Blueprint Genetics	Online Portal	
Thyroid hormone resistance	*THRB* single gene test	Blueprint Genetics	Online Portal	
Tyrosinemia, type 1	*FAH* single gene test	Blueprint Genetics	Online Portal	
Very long chain acyl-CoA dehydrogenase deficiency	*ACADVL* single gene test	Blueprint Genetics	Online Portal	
	ACADVL gene testing	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Use test code 184 for sanger sequencing
Wilson disease	*ATP7B* single gene test	Blueprint Genetics	Online Portal	
Skeletal				
Amelogenesis Imperfecta and Dentinogenesis Imperfecta	Amelogenesis Imperfecta and Dentinogenesis Imperfecta Panel	Blueprint Genetics	Online Portal	
Arthrogryposis	Arthrogryposes Panel	Blueprint Genetics	Online Portal	Also listed under Prenatal heading

Facial dysostosis	Facial Dysostosis and Related Disorders Panel	Blueprint Genetics	Online Portal	Also listed under Dysmorphology heading
Limb malformation	Limb Malformations Panel	Blueprint Genetics	Online Portal	
Short stature, growth failure	Comprehensive Short Stature Syndrome Panel	Blueprint Genetics	Online Portal	
Skeletal dysplasia	Comprehensive Growth Disorders / Skeletal Dysplasias and Disorders Panel (or sub-panel)	Blueprint Genetics	Online Portal	Also listed under Prenatal heading
Whole Exome Sequencing				
Various (Please refer to guidelines)	Whole Exome Sequencing (Trio, Duo, Patient only)	Bluepring Genetics	Online Portal	Trio testing is preferred. **Coordinate parental blood sample collection** at the time that you submit your funding request.
	PGxome® RAPID Exome Test (Trio, Duo, Patient only)	Prevention Genetics	Website Click on "log in/sign up myPrevent" to access online portal	Trio testing is preferred. **Coordinate parental blood sample collection** at the time that you submit your funding request. **Only in cases where results are needed to inform treatment/management decisions within two weeks.**