Molecular Genetics Laboratory
BC Children's Hospital & BC Women's Hospital
2J40 - 4500 Oak Street, Vancouver, BC V6H 3N1
Phone (604) 875-2852 • Fax (604) 875-2707

Test Pricing*

Test	Price
Achondroplasia	$200
Androgen Insensitivity Syndrome	$600
Angelman Syndrome	$200
Ashkenazi Carrier Screening	$150
Brugada Syndrome	$975
CADASIL	$300
Charcot-Marie-Tooth Type 1A	$400
Cystic Fibrosis	$300
Dystonia, Early Onset Primary (DYT1)	$150
Dystrophinopathies (DMD, BMD)	$700
Familial Mediterranean Fever	$500
Friedreich Ataxia	$300
Glucose Transporter Deficiency Type 1	$1,350 – sequencing + MLPA $950 – sequencing only $400 – MLPA only
HFE-Related Hemochromatosis	$150
Hereditary Neuropathy with Liability to Pressure Palsies	$400
Huntington Disease	$150
Hyper IgD Syndrome	$600
Hyperkalemic Periodic Paralysis	$300
Hypokalemic Periodic Paralysis	$400
Hypochondroplasia	$200
Muenke Syndrome	$200
Myotonic Dystrophy Type 1	$300
Myotonic Dystrophy Type 2	$300
Oculopharyngeal Muscular Dystrophy	$150
Prader-Willi Syndrome	$200
Sensorineural Hearing Loss (GJB2/6)	$400
Spinal Muscular Atrophy (SMA)	$400
Spinobulbar Muscular Atrophy (SBMA)	$150
Spinocerebellar Ataxia Panel (SCA1,2,3,6,7)	$300
Steroid 5-alpha-reductase deficiency	$500
Thanatophoric Dysplasia	$200
Transthyretin Amyloidosis	$400
TRAPS	$600
Weaver syndrome	$700
X-linked Ichthyosis (STS Deficiency)	$150
Targeted Testing for known mutation(s)** (carrier or presymptomatic testing)	$300 – single mutation $400 - two mutations in same gene

For prenatal test pricing, please contact the laboratory to discuss.

** Prices are subject to change without notice*

*** Targeted testing for a previously identified family mutation(s) requires a copy of the original mutation report if testing was not completed at our laboratory; may require a sample from the proband as a positive control.*