MyCare
Expert Diagnosis & Treatment Options

Genetic Analysis

- ☑ Whole Genome Sequencing
- ☑ Whole Exome Sequencing

Men's Health

Fertility
Physical Health
Sexual Health
Impotence and Erectile Dysfunction

Women's Health

Fertility
Women's Health
Prenatal Screening
Preconception Health Panel
Polycystic Ovary Disorder

DNA Tests Include:

Oncology

Adrenocortical Carcinoma
Adult solid tumours cancer susceptibility
Astrocytoma
Bladder cancer pertinent cancer
Bone Cancer
Brain cancer
Breast cancer
Burkitt Lymphoma
Childhood solid tumours cancer
Cholangiocarcinoma
Chordoma
Colorectal cancer
Craniopharyngioma Cutaneous T-Cell Lymphoma
Endometrial cancer
Ependymoma
Ewing Sarcoma
Familial breast cancer
Familial rhabdomyosarcoma
Gastric Cancer
GI tract tumours
Head and neck cancer
Hematological malignancies cancer
Hepatocellular Cancer
Histiocytosis, Langerhans Cell
Hodgkin Lymphoma
Inherited non-medullary thyroid cancer
Inherited pancreatic cancer
Inherited renal cancer
Laryngeal Cancer
Medulloblastoma
Melanoma
Multiple Myeloma/Plasma Cell Neoplasms
Myelogenous Leukemia, Chronic
Myeloid Leukemia, Acute
Neoplasm of the heart
Nephroblastoma / Wilms' tumor
Neuroblastoma
Neuroendocrine cancer
Non-Hodgkin Lymphoma
Osteosarcoma
Ovarian cancer
Paraganglioma
Parathyroid Cancer
Pheochromocytoma
Pleuropulmonary Blastoma
Prostate Cancer
Retinoblastoma
Rhabdoid tumour predisposition
Rhabdomyosarcoma (Soft Tissue Sarcoma)
Sarcoma
Small Cell Lung Cancer
Squamous Cell Carcinoma of the Skin
T-Cell Lymphoma
Thymoma
Thyroid cancer
Uveal melanoma

Cardiology

Amyloidosis
Arrhythmogenic cardiomyopathy
Atherosclerosis
Atrial Fibrillation Bradycardia
Cardiac arrhythmias
Cardiomyopathy
Familial non syndromic congenital heart disease
Familial thoracic aortic aneurysm and aortic dissection
Hyperlipidemia
Hypertension
Hypertrophic cardiomyopathy
Long QT syndrome
Mucopolysaccharidosis
Myocardial infarction
Noncompaction cardiomyopathy
Palpitations
Progressive cardiac conduction disease
Sudden cardiac death
Syncope
Thoracic aortic aneurysm and dissection

Pulmonology

Asthma
Bronchitis
Chronic
Cough
Chronic granulomatous disease
Chronic Obstructive Pulmonary Disease
Cystic Fibrosis
Idiopathic Pulmonary Fibrosis
Non-CF bronchiectasis
Pleuritis
Primary ciliary dyskinesia
Pulmonary Embolism
Respiratory ciliopathies including non-CF
Sarcoidosis
Sleep Apnea

Endocrinology & Metabolism

Cushing Syndrome
Diabetes
Thyroid
Celiac Disease
Hyperhomocysteinemia
Metabolic Syndrome
Obesity
Glycogen storage disease II

Aging

Alzheimer Parkinson Disease

Pharmacogenetics

Pharmacogenetic Testing

Musculoskeletal

Muscular Dystrophy
Skeletal Ciliopathies
Skeletal Muscle Channelopathy

Mental Health

ADHD
Anxiety & panic attacks
Autism
Comprehensive Mental Health
Depression
Personality disorder / Bipolar disorder (BPD)
Schizophrenia

Hematology

Anemia
Fanconi anemia
Cytopenias and congenital anemias
Hereditary Erythrocytosis
Myelodysplasia
Polycythemia

Other Health Panels

AllergiAro (Allergy Panel)
EyeAro (Eye Diseases)
Gut Health Panel
Hearing Loss Panel
Nutrient Metabolism
NutriAro (NutriGenomic Panel)
Renal Health Panel
Sports Genetic Panel
Sudden Death Risk Panel (Sports)

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Neurology

- Adult onset movement disorder
- Agnosia
- Aicardi Goutieres Syndrome Disorder
- Alexander Disease
- Amyotrophic Lateral Sclerosis ALS
- Anencephaly
- Aphasia
- Arteriovenous Malformation
- Ataxia Telangiectasia
- Barth Syndrome
- Beckwith-Wiedemann syndrome & other congenital overgrowth disorders
- Behcet Disease
- Blepharospasm Canavan Disease
- Carpel Tunnel Syndrome
- Cerebellar Hypoplasia
- Cerebral Atrophy
- Cerebral Cavernous Malformation
- Cerebral Palsy
- Cerebro Oculo Facio Skeletal Syndrome
- Chiari Malformation
- Childhood onset dystonia or chorea
- Chronic Inflammatory Demyelinating
- Chronic Pain
- Colpocephaly
- Coma
- Common craniosynostosis syndromes
- Congenital Muscular dystrophy
- Congenital Myasthenia
- Congenital Myopathy
- Cranioectodermal Dysplasia
- Craniosynostosis
- Dementia with Lewy Bodies
- Developmental and epileptic encephalopathy
- Dysautonomia
- Dysgraphia Dyslexia
- Friedreich's Ataxia
- Early onset dementia
- Encephalopathy
- Epilepsy
- Facial Palsy
- Familial dysautonomia
- Febrile Seizures (3 Months to 6 Years)
- Foot Drop (foot dorsiflexor weakness)

Neurology

- Headache
- Hemifacial Spasm
- Hemiplegia
- Hereditary neuropathy
- Hereditary spastic paraplegia
- Holoprosencephaly
- Huntington's Disease
- Hydranencephaly
- Hydrocephalus Hypersomnia
- Hypertonia Hypotonia
- Incontinentia Pigmenti
- Infantile Spasms
- Inherited white matter disorders
- Kennedy's Disease
- Krabbe Disease
- Lipoid Proteinosis
- Lissencephaly
- Megalencephaly
- Meningitis
- Menkes Disease
- Microcephaly
- Moyamoya Disease
- Multiple system atrophy
- Myoclonus
- Myopathy
- Narcolepsy
- Neurodegeneration with brain iron accumulation
- Neurofibromatosis Type 1
- Neuroma
- Normal Pressure Hydrocephalus
- Olivopontocerebellar Atrophy
- Opsoclonus Myoclonus
- Orthostatic Hypotension
- Paresthesia
- Parkinson Disease and Complex Parkinsonism
- Paroxysmal Choreoathetosis
- Periodic Paralysis
- Peripheral Neuropathy
- Periventricular Leukomalacia
- Pick's Disease
- Primary Lateral Sclerosis
- Rare multisystem ciliopathy disorders
- Refsum Disease

Neurology

- Retinoblastoma
- Rett Syndrome
- Sandhoff Disease
- Schizencephaly
- Severe microcephaly
- Sotos syndrome
- Spasticity
- Spinal Muscular Atrophy
- Spina Bifida
- Stroke
- Syringomyelia
- Transient Ischemic Attack
- Tremor
- Trigeminal Neuralgia
- Troyer Syndrome
- Tuberous sclerosis
- Von Hippel Lindau Disease
- White matter disorders - adult onset
- Williams Syndrome
- Wilson Disease
- Zellweger Syndrome

Rheumatology

- Antinuclear Antibodies
- Arthritis
- Avascular Necrosis (Osteonecrosis)
- Bursitis
- Familial Mediterranean Fever
- Gout
- Granulomatosis
- Osteoarthritis
- Osteoporosis
- Periodic Fever, Aphthous Stomatitis, Pharyngitis
- Purpura
- Raynaud's Phenomenon
- Scleroderma
- Spinal Canal Stenosis
- Takayasu's Arteritis
- Temporal Arteritis
- Vasculitis