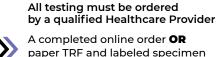


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A completed online order OR paper TRF and labeled specimen is required to initiate testing.

CLINICAL INFORMATION IS REQUIRED for PGnome®, PGxome®, and PGmaxTM panels.

Orders **MUST** include the completed clinical features checklist (preferred) or clinical notes/records. Completion of the checklist is strongly encouraged for all panel testing. The ability to interpret variants directly correlates with the quality of clinical information provided. Also include family medical history/pedigree, if available.

CLINICAL FEATURES					
PERSON COMPLETING FORM	CONTACT (DIREC	CT PHONE OR EMAIL)		DATE OF REQUEST (MM/DD/YYYY)	
	PATIENT IN	FORMATION			
LAST (FAMILY) NAME	FIRST NAME	ORMATION	MI	DATE OF BIRTH (MM/DD/YYYY)	
PATIENT ID	HAS PATIENT BE	EN TESTED PREVIOUSLY AT PREVENTION	IGENETICS?	BIOLOGICAL SEX Male Female Other SPECIFY KARYOTYPE	
	CLINICAL INFORMATION	ON (CHECK ALL THAT APPL	_Y)		
PRE/PERINATAL Abnormality of septum pellucidum Absent septum pellucidum Cavum septum pellucidum Choroid plexus cyst (CPC) Absent nasal bone Congenital heart defect Intracardiac echogenic focus (IEF) Cystic hygroma Increased nuchal translucency, Size (mm): Pleural effusion Pericardial effusion Generalized edema Fetal ascites Hydrops fetalis Diaphragmatic hernia Absent stomach bubble Omphalocele Gastroschisis Echogenic bowel Fetal pyelectasis/hydronephrosis Decreased fetal movement Encephalocele Myelomeningocele/Spina bifida Sacrococcygeal teratoma Intrauterine growth retardation (IUGR) Small for gestational age (SGA) Oligohydramnios Polyhydramnios Short long bones Small thorax Fetal demise Prematurity, Gestational Age: Other:	STRUCTURAL BRAIN ABNORMALITIES / IMAGING Abnormal/delayed myelination Abnormality of basal ganglia Abnormality of brainstem Abnormality of white matter: Periventricular Other: Abnormality of cerebral ventricles: Colpocephaly Hydrocephalus Ventriculomegaly Abnormality of corpus callosum morphology: Agenesis Complete Partial Aplasia/hypoplasia of cerebellar vermis Aplasia/hypoplasia of cerebellum Arnold-Chiari malformation: Type I Cerebral atrophy/hypoplasia Cerebral calcification Holoprosencephaly Intraventricular hemorrhage Preterm Intraventricular hemorrhage Iron deposition Leukodystrophy Neuronal migration abnormality Cortical gyration Gray matter heterotopia Other: DEVELOPMENTAL/ BEHAVIORAL Aggressive/violent behavior Anxiety Attention-deficit hyperactivity disorder Autistic behavior Autism/autism spectrum disorder	Cognitive impairment Delayed fine motor development Delayed gross motor development Developmental regression Gait disturbance Specify: Global developmental delay Hyperactivity Incoordination Intellectual disability Mild Moderate Severe/profound Learning disability Language impairment Absent speech Apraxia Articulation difficulties Delayed speech and language development Expressive Receptive Dysarthria Echolalia Loss of speech Memory impairment Self-injurious behavior: Biting Head-banging Skin picking Sensory processing disorder/neurodevelopmental abnormal Sleep disturbance Stereotypy Recurrent hand flapping Stereotypical hand wringing Other: NEUROLOGICAL Abnormality of nervous system Ataxia Athetosis	nt — ge ity — ity —	radykinesia erebral palsy horea ortical visual impairment ementia tysarthria tyskinesia tysphagia tystonia ncephalopathy ait disturbance, pecify:eadache emiplegia typotonia typertonia tifantile spasms tigraine tyoclonus teuropathy typeripheral Sensory typerityperityperityperity pasticity tyncope tremors tertigo ther: ANIOFACIAL/ SMORPHISM bnormal facial shape, pecify: bnormality of incisors, pecify: la nasi cleft Thick Underdeveloped nteverted nares rachycephaly	

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	PATIENT	
LAST NAME		
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☐ Chin abnormality,	☐ Microcephaly	EYES/VISION	☐ Macular abnormality,
Specify:	☐ Micrognathia	Age of onset of vision issues:	Specify:
Cleft lip:	☐ Nasal base abnormality:	☐ Abnormality of eye movement	Macular dystrophy
☐ Unilateral ☐ Bilateral	Narrow	☐ Esotropia	Microphthalmia
☐ Midline	☐ Wide ☐ Nasal bridge abnormality:	☐ Exotropia ☐ Nystagmus	Myopia
Cleft palate:	Depressed	Smooth pursuit	Ocular albinism
Unilateral	Narrow	Strabismus	Optic atrophy
Bilateral	Prominent	Other:	☐ Optic neuropathy ☐ Palpebral fissure abnormality:
☐ Midline ☐ Submucous cleft	Short	Abnormality of vision,	Downslanted
Cloverleaf skull	☐ Wide	Specify:	Upslanted
Columella abnormality:	☐ Nasal cartilage, absent☐ Nasal ridge abnormality:	☐ Abnormal anterior eye segment morphology	Long
☐ Broad	Depressed	☐ Ablepharon	Short
High insertion	Narrow	Achromatopsia	☐ Almond-shaped
Low hanging	☐ Wide	☐ Aniridia	☐ Proptosis ☐ Ptosis
☐ Low insertion ☐ Short	Nasal tip abnormality:	☐ Ankyloblepharon	Retinal flecks
Craniosynostosis:	Bifid	Anophthalmia	Retinal detachment
☐ Coronal	☐ Broad ☐ Depressed	☐ Blepharochalasis	Retinitis pigmentosa
Lambdoidal	Deviated	☐ Blepharophimosis	Synophrys
Metopic	☐ Narrow	☐ Cataracts	☐ Telecanthus
☐ Orbital ☐ Sagittal	Overhanging	Cataracts, congenital	Other:
☐ Dolichocephaly	Nasolabial fold abnormality:	☐ Coloboma	EARS/HEARING
Face abnormality:	☐ Prominent☐ Underdeveloped	Corneal opacity	Age of onset of hearing loss:
☐ Broad	Neck abnormality:	Corneal dystrophy	Hearing impairment
Coarse facial features	Broad	Cone/cone-rod dystrophy	Sensorineural
Flat	Long	Congenital stationary night	Congenital
☐ Long ☐ Narrow	Webbed	☐ Cryptophthalmos	Bilateral
☐ Round	☐ Short ☐ Redundant nuchal skin	Deeply set eyes	☐ Progressive ☐ Conductive
Short	Nose abnormality:	☐ Distichiasis	Congential
Square	Absent	Dyschromatopsia (color	Bilateral
Triangular	☐ Bifid	blindness)	☐ Progressive
Forehead abnormality:	Long	☐ Ectopia lentis	Mixed
☐ Broad ☐ Narrow	│	☐ Ectropion	☐ Anotia
☐ Prominent	☐ Short	☐ Entropion	Abnormal newborn screen, Specify:
Sloping	Wide	☐ Epiblepharon	Antihelix abnormality:
Creases	Occiput abnormality:	Epicanthus/epicanthal folds	Absent
Frontal bossing	☐ Flat	Epicanthus inversus	Additional crus
☐ Jaw abnormality: ☐ Broad	Prominent	☐ Eyebrow abnormality: ☐ Broad	Angulated
□ Narrow	☐ Plagiocephaly	│ │ │ Highly arched	☐ Inferior crus broad ☐ Inferior crus prominent
Lip vermilion abnormality	☐ Philtrum abnormality: ☐ Broad	Horizontal	☐ Inferior crus underdeveloped
Lip abnormality:	☐ Deep	Sparse	Superior crus prominent
Pit	Hypoplastic	Thick	Superior crus underdeveloped
☐ Thin	Long	☐ Eyelash abnormality: ☐ Absent	☐ Antitragus abnormality: ☐ Absent
☐ Thick ☐ Tented	☐ Narrow ☐ Smooth	Long	☐ Absent
☐ Exaggerated cupid's bow	Short	Prominent	☐ Everted
Absent cupid's bow	Tented	Sparse	Prominent
☐ Malar abnormality:	☐ Proboscis	☐ Eyelid cleft	Underdeveloped
☐ Flattening	Prognathism	External ophthalmoplegia	☐ Ear abnormality: ☐ Abnormality of the tragus
☐ Prominence ☐ Midface abnormality:	Retrognathia	☐ Progressive	Abnormality of the tragus Auricular pit
☐ Midiace abnormality:	☐ Scaphocephaly	☐ Glaucoma ☐ Infraorbital abnormality:	Crumpled
Prominence	☐ Supraorbital ridge abnormality: ☐ Prominent	Crease	Cupped
Retrusion	Underdeveloped	Fold	Long
Macrocephaly:	☐ Trigonocephaly	☐ Iris abnormality,	☐ Low-set ☐ Posteriorly rotated
☐ Relative ☐ True	Turricephaly	Specify:	Preauricular pit
☐ Metopic suture abnormality:	Other:	Lagophthalmos	Protruding
Depression		Leber optic atrophy	Short
Ridge		☐ Lens subluxation	☐ Satyr ☐ Tag



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	I □ Lymanhanania	Dradveardia	1
Helix abnormality:	Lymphopenia	☐ Bradycardia	GENITOURINARY
Cleft / Notching	☐ Neutropenia	Coarctation of the aorta	Abnormality of the uterus,
Crimped	☐ Pancytopenia	Congenital heart defect	Specify:
☐ Darwin notch ☐ Darwin tubercle	Recurrent infections	☐ Dilated cardiomyopathy	Ambiguous genitalia
☐ Notching	☐ Severe combined	Double outlet right ventricle	☐ Chordee
Overfolded	immunodeficiency	Ebstein anomaly	☐ Cryptorchidism
☐ Prominent	☐ Thrombocytopenia	☐ Heterotaxy	☐ Duplicated collecting system
☐ Thin	Other:	Hypertension	Horseshoe kidney
Lobe abnormality:	SKIN/HAIR	☐ Hypertrophic cardiomyopathy	Hydronephrosis
☐ Cleft	Abnormal blistering of the skin,	☐ Mitral valve prolapse	☐ Hypospadias/epispadias
Forward-facing	Specify:	☐ Noncompaction cardiomyopathy	☐ Inguinal hernia
Large	Abnormality of nail:	Patent ductus arteriosus	☐ Micropenis
Small Small		Patent ductus arteriosas	☐ Multicystic kidney dysplasia
☐ Uplifted	☐ Deep-set	Prolonged QTc interval	☐ Nephrolithiasis
☐ Macrotia	Pits Pits	☐ Pulmonary hypertension	☐ Polycystic kidney disease
Other:	☐ Albinism	☐ Pulliforlary Hypertension ☐ Arteria	l —
ENDOCRINE	☐ Alopecia	☐ Vascular	Renal agenesis/hypoplasia Unilateral agneisis
Adrenal insufficiency (Addison)	☐ Anhidrosis	Sudden death	Bilateral ageneisis
Androgen excess	Cafe-au-lait spot:	Tetralogy of Fallot	Unilateral hypoplasia
Androgen insensitivity	Single	☐ Transposition of the great vessels	☐ Blateral hypoplasia
Congenital adrenal hypoplasia	Multiple	☐ Truncus arteriosus	Sex reversal
Congenital adrenal hyperplasia	☐ Coarse hair	—	☐ Vesicoureteral reflux
_ , ,	☐ Collodion baby	☐ Ventricular septal defect	Other:
Delayed bone age	Cutaneous photosensitivity	☐ Ventricular tachycardia	
Delayed puberty	☐ Cutis laxa	Other:	MUSCULOSKELETAL
☐ Diabetes insipidus	Dry skin	GASTROINTESTINAL	Abnormal connective tissue
Diabetes Mellitus	☐ Eczema	☐ Biliary atresia	Abnormal digit morphology
Hyperandrogenism	Erythematous skin	☐ Cholestasis	☐ Broad ☐ Short
Hyperglycemia	☐ Hemangioma	Constipation:	Clinodactyly
Hyperphosphatemia	☐ Hairline:	Acute	Ectrodatyly
☐ Hyperthyroidism	☐ Anterior	☐ Chronic	Oligodactyly
☐ Hypoglycemia	Low	☐ Diarrhea	Polydactyly
☐ Hypophosphatemia	☐ High	☐ Diaphragmatic hernia	Postaxial
☐ Hypothyroidism	☐ Posterior	☐ Duodenal stenosis/atresia	☐ Preaxial
☐ Increased cortisol level (Cushing)	_ Low	Esophageal stenosis/atresia	Syndactyly
☐ Maturity-onset diabetes of the	☐ High	Exocrine pancreatic insufficiency	☐ Arachnodactyly
young	☐ Hyperextensible skin	☐ Failure to thrive	☐ Arthralgia
☐ Precocious puberty	☐ Hyperpigmentation of the skin	Feeding difficulties	☐ Arthrogryposis
Rickets	☐ Hypopigmentation of the skin	Gastroesophageal reflux	☐ Bruising susceptibility
Other:	☐ Hypohidrosis	☐ Gastroschisis	Chest abnormality:
RESPIRATORY	☐ Ichthyosis	│	Small chest
☐ Asthma	☐ Jaundice	☐ Hepatosplenomegaly	☐ Barrel-shaped
☐ Bronchiectasis	☐ Lipoma	☐ Inflammatory bowel disease	☐ Bell-shaped thorax ☐ Pectus carinatum
☐ Bronchomalacia	Lymphedema	☐ Jaundice	Pectus cannatum Pectus excavatum
☐ Hyperventilation	Palmoplantar keratoderma	Liver disease	Contractures of joint(s)
Hypoventilation	☐ Scarring of skin	☐ Liver disease	Decreased muscle mass
Laryngomalacia	☐ Skin rash	│	Delayed bone age
☐ Laryngornalacia	☐ Sparse hair	☐ Omphalecele	☐ Delayed borie age
☐ Pneumothorax	☐ Telangiectasia	☐ Pancreatitis	☐ Exercise intolerance
	☐ Vascular skin abnormality	l 😑 🗀	L
☐ Pulmonary fibrosis	☐ Velvety skin	- -	Fatigue
Respiratory insufficiency	Other:	Splenomegaly	Fracture(s)
☐ Tracheomalacia		☐ Tracheoesophageal fistula	Hemihypertrophy
☐ Tracheoesophageal fistula	CARDIAC	☐ Tube feeding ☐ Nasogastric	Hypertonia
Other:	Amyloidosis	Gastrostomy	Hypotonia
HEMATOLOGIC/IMMUNOLOGIC	Aortic root dilatation	Gastrostorny Gastrojejunal	☐ Joint hypermobility
☐ Agammaglobulinemia	Arrhythmia	☐ Umbilical hernia	☐ Kyphosis
☐ Allergic rhinitis	Atrial septal defect	│	Limb shortening:
☐ Anemia	Atrioventricular canal defect	Other:	Mesomelic
☐ Hemolytic anemia	Arrhythmogenic right ventricular		☐ Micromelic ☐ Rhizomelic
☐ Immunodeficiency,	dysplasia		Metaphyseal abnormalities:
Specify:	☐ Bicuspid aortic valve		Dumbbell



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☐ Flared ☐ Muscle weakness ☐ Myalgia ☐ Myopathic facies ☐ Myopathy ☐ Myelomeningocele/Spina Bifida/ Neural Tube Defect ☐ Osteoarthritis ☐ Osteoporosis ☐ Osteopenia ☐ Pain: ☐ Absent/decreased ☐ Abnormal sensation ☐ Episodic ☐ Limb ☐ Muscle ☐ Platyspondyly ☐ Recurrent fractures ☐ Rhabdomyolysis ☐ Rib abnormality: ☐ Cupped ☐ Fused ☐ Supernumerary ☐ Missing ☐ Short	Talipes	r:ature dysplasia novarus r:ure dysplasia abnormality: coted dalangeal al bodies, abnormal form cia/hypoplasia erfly n ivertebrae R SYSTEM m	Dissecting Thoracic Cerebral Other: Arterial calcification Arterial dissection Arterial tortuosity Arteriovenous malfore Epistaxis Lymphedema Pulmonary hypertens Arterial Vascular Stroke Other: OTHER TESTING Provide copy of report(Echocardiogram: EEG: EMG/NCV: Biopsy: Gene testing: Results:	sion:	If you would like us to comment on the presence/ absence of previously identified variants, provide a copy of the original report. Chromosomal Microarray (CMA): MRI brain: CT brain: CT (other): Muscle biopsy: Ultrasound: X-Ray:
METABOLIC FINDINGS • A Abnormal newborn screen Specify: Abnormal metabolic profile (please check each metabolite outside nor Acylcarnitine Acylglycines Amino Acids Biotindase Carnitine Cerebrospinal fluid Coenyzme/enzyme activity Creatine phosphokinase (CPK) Essential fatty acids Folate Hepatic Transaminase Homocysteine Hormones Ketones Lactic acidosis Lipase	mal limits)	Lipoproteins Lysosomal enzymes Mucopolysaccharides Oligosaccharides Porphyrin Pterins Purines Pyrimidine Pyruvate Serum alpha fetoprot Sterols/Oxysterols Transferrin Uric acid Very long chain fatty a Abnormal vitamin (please check each vitamin me	ein (AFP) acids (VLCFA) levels easuring outside normal limits)	Zinc Other Other me Abnorm studies Abnorm Abnorm activity Hyperan Hyperbil Hypergly Hyperlip Hypogly Hypolipi Plasm Urine Lactic Ac Metabol Methylm	etabolic features al cerebrospinal fluid (CSF) al glycosylation al mitochondrial respiratory chain al mitochondrial respi

ADDITIONAL NOTES

If additional space is required, please continue your response below. Include a continued notation on the other or specify line.



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		CANCE	RHISTOR	Υ				
Patient Information								
No personal history of cancer □ Ovarian/Fallopian Tube / Primary Peritoneal		☐ Pancrea Age of di			Other Age of	Other Age of diagnosis:		
Breast		Age of diagnosis:	☐ Prostate	e		Detail	S:	
Age of diagnosis: Colorectal								
☐ Triple-Negative (ER, PR, Her2 negative)☐ DCIS (Ductal Carcinoma In		Age of diagnosis: ASI/IHC results:	Metastatic Yes No Unknown Gleason Score					
Situ) DC (Invasive Ductal Carcinoma) ILC (Invasive Lobular Carcinoma) Bilateral / >1 Primary		Endometrial / Uterine Age of diagnosis: MSI/IHC results:	Polyps Age of diagnosis: Number of polyps: Pathology details:					
Family History of Cancer or In	clude Pedi	gree						
☐ No known family history	of cance	•						
relatives having lived beyo	e Limited fa ond age 45	amily history available such as	fewer than two	o female fi	rst or second	d-degree m	aternal or pate	rnal
Ashkenazi Jewish 🗌 NO	YES, Ma	ternal Yes, Paternal Ur	nknown					
RELATION TO PATIENT	SELECT	CANCER / POLYP TYPE / GLEASON S	CORE	AGE OF DIAGNOSIS	UNAVAILABLE FOR TESTING		PATIENT HAS NO CONTACT WITH WITH RELATIVE	RELATIVE DECLINES TESTING
	☐ Maternal ☐ Paternal							
	☐ Maternal ☐ Paternal							
	☐ Maternal ☐ Paternal							
	☐ Maternal ☐ Paternal							
	☐ Maternal ☐ Paternal							
PAST FAMILY GENETIC TESTING	🕽 🗌 NO pre	vious testing in family. 🔲 YES, Ir	nclude Germline, Sc	matic or Tun	nor testing result	s. Describe or	attach copies of re	eport.

PEDIGREE

Use this area to include a pedigree and/or additional relevant medical/family history.

VARIANT

KNOWN FAMILIAL VARIANT: GENE___