

Suite 1020, Calgary Place Tower 1, 330-5 Ave SW Calgary, AB, T2P 0L4 T: 403.250.2221 | F: 587.349.9751 E: info@bioaro.com

	Pc	itient Information						R	eferring Physic	iar	1
								_			
								1	•		
								1			
			Label H	ere	9			1			
								; P	hysician Signature:		
	- 1								Copies to:		
								I I			
	Gei	netic Analysis	Microbiome		alysis			h			Women's Health
-	Wł	nole Genome Sequencing	☐ Gut		yaginal		Fertility				Fertility
					Oral & Dental	\Box	Men's Health				Women's Health
						R	Men's Sexua				Prenatal Screening
						ш	Menimpote	ice ai	nd Erectile Dysfunction	H	Preconception Health Panel PolyCystic Ovary Disorder
)is	sec	ase Based DNA Tests									, , , ,
		cology			Cardiology						Musculoskeletal
		Adrenocortical Carcinoma			Amyloidosis						Muscular Dystrophy
	_	Adult solid tumours cancer susce	ptibility		Arrhythmogenic ca	rdion	nyopathy				Skeletal Ciliopathies
		Astrocytoma	,		Atherosclerosis		, , ,				Skeletal Muscle Channelopathy
		Bladder cancer pertinent cancer			Atrial Fibrillation						Mental Health
	_	Bone Cancer Brain cancer			Bradycardia Cardiac arrhythmic	ıs					ADHD
	_	Breast cancer		H	Cardiomyopathy	45					Anxiety & panic attacks
		Burkitt Lymphoma			Familial non syndro						Autism Comprehensive Mental Health
		Childhood solid tumours cancer			Familial thoracic ac	ortic o	aneurysm and	d aorti	c dissection		Depression
	_	Cholangiocarcinoma Chordoma			Hyperlipidemia Hypertension						Personality disorder / Bipolar disorder (BPD)
	_	Colorectal cancer			Hypertrophic cardio	omyc	pathy				Schizophrenia
		CraniopharyngiomaCutaneous T-	-Cell Lymphoma		Long QT syndrome						Hematology
	_	Endometrial cancer Ependymoma			Mucopolysaccharic Myocardial infarction						Anemia
	_	Ewing Sarcoma			Noncompaction ca		nvopathv				Fanconi anemia
	_	Familial breast cancer			Palpitations		, , ,				Cytopenias and congenital anemias Hereditary Erythrocytosis
		Familial rhabdomyosarcoma			Progressive cardiac		duction disec	ise			Myelodysplasia
	_	Gastric Cancer GI tract tumours			Sudden cardiac de Syncope	atn					Polycythemia
	_	Head and neck cancer			Thoracic aortic ane	urysı	m and dissec	tion			Other Health Panels
		Hematological malignancies can	cer			•					AllergiAro (Allergy Panel)
		Hepatocellular Cancer			Pulmonology						EyeAro (Eye Diseases)
		Histiocytosis, Langerhans Cell Hodgkin Lymphoma			Asthma Propobitis						Gut Health Panel
		Inherited non-medullary thyroid c	ancer		Bronchitis Chronic Cough						Hearing Loss Panel
		Inherited pancreatic cancer		H	Chronic granulomo	tous	disease				Nutrient Metabolism NutriAro (NutriGenomic Panel)
		Inherited renal cancer			Chronic Obstructive	e Pulr	monary Disea	se			Renal Health Panel
		Laryngeal Cancer Medulloblastoma			Cystic Fibrosis Idiopathic Pulmond	ur i Fik	oronio				Sports Genetic Panel
	_	Melanoma			Non-CF bronchiect		310313				Sudden Death Risk Panel (Sports)
		Multiple Myeloma/Plasma Cell Ne	oplasms		Pleuritis				lan to the contract of		
		Myelogenous Leukemia, Chronic			Primary ciliary dysk		a		Clinical Notes:		
		Myeloid Leukemia, Acute Neoplasm of the heart			Pulmonary Embolis Respiratory ciliopat		includina nor	-CF	İ		
		Nephroblastoma / Wilms' tumor		ä	Sarcoidosis		morading nor	. 0.	ļ		
	_	Neuroblastoma			Sleep Apnea				-		
	_	Neuroendocrine cancer Non-Hodgkin Lymphoma			Endocrinology	& Ме	etabolism				
		Osteosarcoma			Cushing Syndrome				l I		
	_	Ovarian cancer			Diabetes						
	_	Paraganglioma			Thyroid						
		Parathyroid Cancer Pheochromocytoma			Celiac Disease	omio	•				
		Pleuropulmonary Blastoma			Hyperhomocystein Metabolic Syndrom		4				
	_	Prostate Cancer			Obesity				i i		
		Retinoblastoma Rhabdoid tumour predisposition			Glycogen storage	disea	ise II		I		
		Rhabdoid tumour predisposition Rhabdomyosarcoma (Soft Tissue	Sarcoma)		Aging						
		Sarcoma			Aging Alzheimer				-		
		Small Cell Lung Cancer			Parkinson Disease						
		Squamous Cell Carcinoma of the T-Cell Lymphoma	Skin						i		
		Thymoma			Pharmacogene				l L		
		Thyroid cancer			Pharmacogenetic ¹	Testii	ng				
		Uveal melanoma							1		



Suite 1020, Calgary Place Tower 1, 330-5 Ave SW Calgary, AB, T2P 0L4 T: 403.250.2221 | F: 587.349.9751 E: info@bioaro.com

	Date:					
!	II.	an Name:				
!	i	Physician Address: Physician Number: Physician Signature:				
Label H	1616					
i I						
	Copies t	to:				
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 Behect Disease Blepharospasm Canavan Disease Carpel Tunnel Syndrome Cereballar Hypoplasia Cerebral Cavernous Malformation Cerebral Cavernous Malformation Cerebral Palsy	Neurology Headache Hemifacial Spasm Hemiplegia Hereditary neuropathy Hereditary spastic paraplegia Holoprosencephaly Huntington's Disease Hydranencephaly Hydrocephalus Hypersomnia Hypertonia Hypertonia Hypotonia Incontinentia Pigmenti Infantile Spasms Inherited white matter disorders Kennedy's Disease Krabbe Disease Lipoid Proteinosis Lissencephaly Megalencephaly	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				
Cerebral Palsy Cerebro Oculo Facio Skeletal Syndrome	☐ Megalencephaly ☐ Meningitis	☐ Wilson Disease☐ Zellweger Syndrome				
Chiari Malformation	☐ Menkes Disease	•				
Childhood onset dystonia or chorea Chronic Inflammatory Demyelinating	☐ Microcephaly☐ Moyamoya Disease	Rheumatology				
Chronic Inflammatory Demyelinating Chronic Pain Colpocephaly Coma Common craniosynostosis syndromes Congenital Muscular dystrophy Congenital Myosthenia 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)	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	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				

CONFIDENTIALITY STATEMENT: Information contained in this communication may be confidential and is intended only for the use of the recipient(s). If the reader of this message is not the intended recipient, you are hereby notified that any dissemination, distribution, or copying of this communication or any of its content is strictly prohibited. If you receive this communication or any of its contents in error please return it to the sender and contact BioAro Inc. (1-877-748-0032)