



Mitochondrial and Autoimmune Neurological Disorders Laboratory

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Patient Information			
Title Surname	First Name	M/F Date of Birth	
Street	Suburb	State Postcode	
Country	Telephone No.	UR No.	
Physician Information			
Title Surname	First Name	Provider No.	
Hospital / Medical Centre/Clinic/Institution			
Hospital / medical Centre/Onnic/institution			
Street	Suburb	State Postcode	
Country Telephone N	lo. Fax No.	E-mail address	
Specimen and Signed Consent for Genetic testing			
Note: Testing cannot proceed without a signed consent and billing details			
10-20 2 x 3 mls	20-50 mls >50mg		
Hair follicles Blood (EDTA) Urine 1 st morning void Muscle Date taken Date Sent			
Collection of two of more specimens are required for genetic testing. All genetic testing to be accompanied by genetic counselling.			
Genetic information obtained from these tests will be kept confidential and not released to anyone without prior patient permission. Does the patient consent to the testing of their DNA which may identify genetic variations (e.g. mutations) and may have implications for family members? Yes No			
Does the patient consent to the non-identified use of their specimen/s for test quality assurance and validation activities and reports? Yes No Signature. X (Patient/Next of Kin)			
Billing Information Genetic tests <u>do not</u> attract a Medicare or Private Health Care Rebate (Please note no Item numbers for rebates)			
MT-TL1 gene (m.3230-m.3304)(N	MELAS m.3243A>G, m.3271T>C)	\$150 Bill (Tick or Cross)	
MT-TK gene (m.8295-m.8364)(MERRF m.8344A>G, m.356T>C) \$150			
MT-ATP6 gene (m.8527-9207)(NARP/LS m.8993T>C/G, m.9176T>C) \$150			
LHON Multiplex RFLP m.3460G>A, 11778G>A, m.14484T>C		\$150 Hospital /	
Sanger Sequencing per mtDNA gene		\$150 Pathology / Medical Centre	
Southern Deletion screen		\$400	

Office use only SVR No Specimen No/s			
Date:/	Staff Member		
Name:	Clinical Features - Check Boxes		
Clinical Features - Check Boxes	Signs Yes No N/A		
Presenting Complaint:	Short Stature		
	Congestive Heart Failure		
Age of onset:	Respiratory Insufficiency		
2 1 1/2	Diabetes Mellitus		
Symptoms Yes No N/A	Hypothyroidism		
Development Delay	Hypoparathyroidism		
Intellectual disability	Optic Atrophy		
Exercise Intolerance	Ophthalmoplegia		
Nausea/Vomiting	Ptosis		
GI Pseudoobstruction	Lipomas		
Headache/Migraine	Retinopathy		
Stroke	Cortical Blindness		
Episodic Coma	Cerebellar Signs		
Dementia			
Seizures	Hearing Loss		
Myoclonus	Proximal Limb Weakness		
Family History	Neuropathy		
Maternal	Rhabdomyolysis		
Sporadic	Exercise related		
Autosomal dominant	Spontaneous		
Other please specify:	1 1 2 1 N N N N N N N N N N N N N N N N		
maging Studies- Norm Abnorm N/A	Laboratory Studies- Yes No N/A		
Angiogram	Elevated Blood Lactate Pyruvate		
MRI 🔲 🗀	CSF Protein		
SPECT	ECG Heart Block Pre-excitation		
CSF	EMG/NCS Myopathic		
ст 🔲 🔲	Neurogenic Other		
Basal Ganglia Calcification	(eg. Hisotopathology. Biochemistry) please specify or attach report:		
ont Name: Requisition Form NATA Accr. No. 13786 Version 19			

Information collected on this form is to determine the most appropriate mtDNA genetic tests.

This form is not a referral slip for specimen collection.

Patients must have specimens collected at their nearest pathology collection centre.

This form must accompany the specimen/s to the laboratory for testing.