Institut für Humangenetik
des Klinikums rechts der Isar
der Technischen Universität München

Direktor: Univ. Prof. Dr. med. Th. Meitinger

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Molecular Diagnostic Request Form

Patient Information

□ female □ male

Last Name, First Name: _____

DOB : _____

Street: _____

City, ZIP,State: _____

Billing Information

Last Name, First Name: _____

Street: _____

City, ZIP,State: _____

Please sign the added assumption of costs (download)

Clinical History (short description; please add last medical report)

□ symptomatic □ not symptomatic

Family History

Negativ □ Positiv (please add pedigree) □

Referring Physician (Please print)

Name: _____Phone_____

Institution: _____

_____

_____

Signature:

- □ Pantothenatkinase associated - Neurodegeneration (PANK2)
- □ PLA2G6 – related Neurodegeneration (PLA2G6)
- □ Neuroferritinopathy (FTL)
- □ Thiamine-responsive megaloblastic anemia syndrome (SLC19A2)
- □ Parkinson Disease (PARK8) (LRRK2)
- □ Parkinson Disease (PARK6) (PINK1)
- □ Parkinson Disease (PARK7) (DJ1)
- □ Retinitis pigmentosa (RP2, RPGR)
- □ Basal ganglia disease, biotin-responsive (SLC19A3)
- □ MTHFR Deficiency (MTHFR)
- □ Hypophosphatemia (X-linked)(PHEX)
- □ Hypophosphatemia (AD, autosomal dominant)(FGF23)
- □ Hypophosphatemia with Hypercalciuria (SLC34A3)
- □ Hypophosphatemia (AR, autosomal recessive) (DMP1)
- □ Glucocorticoid Deficiency 1 (MC2R)
- □ Glucocorticoid Deficiency 2 (MRAP)
- □ Adrenal Hypoplasia (congenital) (NR0B1)

Shipping

Material: DNA (20 µg) Prenatal diagnosis: please contact us by email or phone