National University Hospital
5 Lower Kent Ridge Road
Singapore 119074
www.nuh.com.sg
Tel: (65) 6772 4346
Fax: (65) 6775 1757

MEMORANDUM

To : All Laboratory Users

From : Molecular Diagnosis Centre, Department of Laboratory Medicine

Date : 3rd December 2020

Re : **Genetic Consent Form Requirement For Genetic Tests**

Dear Users,

Please note that in accordance with the Ministry of Health (MOH) Code of Practice on the Standards for the Provision of Clinical Genetic / Genomic Testing Services and Clinical Laboratory Genetic / Genomic Testing Services (https://www.moh.gov.sg/licensing-and-regulation/regulations-guidelines-and-circulars/details/code-of-practice-on-the-standards-for-the-provision-of-clinical-genetic-genomic-testing-services-and-clinical-laboratory-genetic-genomic-testing-services), a signed informed consent form is required to be submitted with the sample for the following genetic tests performed by the Molecular Diagnosis Centre (MDC) in the list given below.

A consent form can be downloaded from the Intranet – Department of Laboratory Medicine, eForms, MDC, Informed Consent Form for Genetic/Genomuc Testing, in the link below:

http://10.11.196.15/labmedjmgs/lmdocs/eforms/mdc/MDC%20General%20consent%20(R1).pdf

Please note that the test will not be performed if the completed consent form is not received with effect from 1st January 2021.

- Alpha Thalassemia genotyping (ATHAL)
- Beta Thalassemia genotyping (BTHAL)
- Alpha thalassemia genotyping prenatal (GENEA)
- Beta Thalassemia genotyping prenatal (GENEB)
- Methylenetetrahydrofolate reductase (MTHFR) variant (MTHFR)
- Factor V Leiden Thrombophilia (FA5L)
- Prothrombin G20210A Thrombophilia (PTGT)
- Fragile X PCR (FXPCR)
- Prader Willi / Angelman syndrome (PWASM)
- Y chromosome deletion (YCHD)
- Androgen receptor mutation Exons 4-8 (ARS)
- Androgen receptor mutation Exons 1-3 (ARMX)
- Male infertility panel (MIP)
- Leber Hereditary Optic Neuropathy (LHON)
- Hearing impairment, inherited causes (HIMP)
- ATP7B Mutation Screen Wilson disease (ATPMS)

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- ATP7B Known Mutation (ATPKM)
- MODY1/2/3 Maturity Onset Diabetes of the Young (MODY)
- MODY Known Mutation (MODYK)
- Multiple Endocrine Neoplasia Type 2A/FMTC (MEN2A)
- Multiple endocrine neoplasia (MEN) 2B (MEN2B)
- Haemochromatosis gene (HFE) mutation detection (HFE)

For further queries, please contact Mr Lee Chun Kiat at 67724175 or 67724384.

Thank you.

Dr Benedict Yan
Clinical Head
Molecular Diagnosis Centre
Department of Laboratory Medicine

Dr Karen Tan
Deputy Clinical Head
Molecular Diagnosis Centre
Department of Laboratory Medicine