



## Molecular and Genetic Referral Form

### PATIENT DETAILS

Name: \_\_\_\_\_ Surname: \_\_\_\_\_ D.O.B.: \_\_\_\_/\_\_\_\_/\_\_\_\_ I.D. No.: \_\_\_\_\_ Nationality: \_\_\_\_\_  
 Gender:  Male  Female Int No.: \_\_\_\_\_  GESY,  Private, Address: \_\_\_\_\_ City: \_\_\_\_\_  
 Code \_\_\_\_\_ Country: \_\_\_\_\_ Phone: Home: \_\_\_\_\_ Work: \_\_\_\_\_ Fax: \_\_\_\_\_ e-mail: \_\_\_\_\_

### REFERRING CLINICIAN / SCIENTIST / CLINIC / LABORATORY DETAILS

Name: \_\_\_\_\_ Surname: \_\_\_\_\_ Hospital / Clinic / Lab: \_\_\_\_\_  
 Address: \_\_\_\_\_ City: \_\_\_\_\_ Code: \_\_\_\_\_ Country: \_\_\_\_\_  
 Phone: \_\_\_\_\_ Fax: \_\_\_\_\_ e-mail: \_\_\_\_\_  
 Reason for Referral: \_\_\_\_\_

Signature: \_\_\_\_\_ Date: \_\_\_\_/\_\_\_\_/\_\_\_\_

### SAMPLE DETAILS (Please tick accordingly)

Date and Time of Sample Collection: \_\_\_\_\_

Sample:  Blood (3-4ml)  CVS  SWAP  URINE  
 Other (please specify): \_\_\_\_\_

First Investigation  Repetition

For Genetic Testing 3-4ml whole blood is required in EDTA.

### TEST REQUEST (Please tick accordingly)

#### Cystic Fibrosis (CF) (FRAGMENT ANALYSIS):

- CF full mutation analysis (29 mutations)
- CF analysis for known mutation
- CF prenatal diagnosis

#### Haemochromatosis (RESTRICTION DIGEST):

- HFE C282Y & H63D mutations

#### Hypercholesterolemia (SANGER SEQUENCING):

- LDLR - Hypercholesterolemia analysis

#### Lactose Intolerance (PCR):

- MCM6 - poly T-13910C

#### FAST PCR (RT-PCR):

- HPV - human papillomavirus
- CT & NG - Chlamydia and Gonorrhoea
- TV - Trichomonas vaginalis
- CT & NG - Chlamydia and Gonorrhoea
- GBS - Group B Streptococcus
- HCV Viral Load
- HBV Viral Load
- HIV Viral Load
- HIV Qualitative
- Other DNA analysis upon request: \_\_\_\_\_
- DNA extraction/storage

#### Hereditary Recurrent Fevers (HRFs) (SANGER SEQUENCING):

##### FMF

- MEFV full mutation analysis (Exons 2, 3, 5 & 10)
- MEFV analysis for known mutation

##### MVK

- MKD full mutation analysis (Exons 8, 9 & 10)
- MKD analysis for known mutation

##### TRAPS

- TNFRSF1A full mutation analysis (Exons 2, 3 & 4)
- TNFRSF1A analysis for known mutation

##### CAPS

- NLRP3 full mutation analysis (Exon 3)
- NLRP3 analysis for known mutation

#### Multiplexing for Pathogens (RT-PCR):

- Respiratory** Adenovirus, Bocavirus, Coronavirus 229E, Coronavirus HKU1, Coronavirus NL63, Coronavirus OC43, human Metapneumovirus A/B, Influenza A, Influenza A subtype H1N1/2009, Influenza A subtype H1, Influenza A subtype H3, Influenza B, Parainfluenza virus 1, Parainfluenza virus 2/3/4, Respiratory Syncytial virus A/B, Rhinovirus/Enterovirus, Bordetella pertussis, Legionella pneumophila, Mycoplasma pneumoniae
- Gastrointestinal** Clostridium difficile toxin A/B, Enterococci (EPEC), Enteroinvasive E.coli (EIEC)/Shigella, Enteropathogenic E.coli (EPEC), Enterotoxigenic E.coli (ETEC) lt/st, Pathogenic Campylobacter spp., (C.jejuni, C.upsaliensis, C.coli), Plesiomonas shigelloides, Salmonella, Shiga-like toxin producing E.coli (STEC) stx1/stx2 & O157:H7, Vibrio cholera, parahaemolyticus, vulnificus, Yersinia enterocolitica, Cyclospora cayentanensis, Cryptosporidium spp., Entamoeba histolytica, Giardia lamblia, Adenovirus F40/41, Astrovirus, Norovirus GI/GII, Rotavirus A, Sapovirus (I, II, IV, V)

### PATIENT INFORM CONSENT (Please read and sign)

I authorize the Clinical Laboratory Bioanalysis to use my (or my child's/my foetus) sample (whole blood, serum or CVS) for genetic testing or storage. I have the right to refuse the above and request disposal of my sample. Samples are stored for future reference or use only.

I can withdraw my consent at any time by contacting the laboratory at +35725 72 62 52

Patient/Guardian Signature: \_\_\_\_\_