



CYTOGENETICS AND MOLECULAR GENETICS ASSESSMENT REQUISITION FORM

* Fields are mandatory

PATIENT INFORMATION							
Surname*		Given Name(s)*		Date of Birth* DD MM YYYY		Ethnicity* <input type="checkbox"/> Chinese <input type="checkbox"/> Other: _____	
Sex* <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Other		HKID/Passport No.*		Mailing Address			
Clinic/Institution No.		Phone No.		Email		FOR LAB USE ONLY <input type="checkbox"/> Tick if previously tested <div style="border: 1px dashed gray; padding: 5px; width: fit-content;">Patient Label</div> Specimen received date/ time*:	
REFERRING INFORMATION (Fill out or affix official stamp)							
Referring Physician*			Clinic/Institution Name*				
Address							
Phone No.*		Fax No.		Email			

BILLING INFORMATION	SPECIMEN DETAILS
Bill Type <input type="checkbox"/> Charge to Institution <i>(Select Referral by Physician in the Referral Type)</i> <input type="checkbox"/> Direct Payment	Specimen Type (x no. of tubes) Blood: <input type="checkbox"/> EDTA (x ___) <input type="checkbox"/> Sodium Heparin (x ___) <input type="checkbox"/> Clotted (x ___) <input type="checkbox"/> ACD Solution B (x ___) Others: <input type="checkbox"/> Amniotic Fluid <input type="checkbox"/> Specify: _____ #
Referral Type <input type="checkbox"/> Referral by Physician <input type="checkbox"/> Referral by Staff: _____ <input type="checkbox"/> Walk-in	Specimen collection date* DD MM YYYY Specimen collection time* HR : MIN

INDICATION AND TEST REQUESTED
Indication (specified by referring physician)*

Reproductive, Obstetrics and Gynecology, Pediatric Karyotyping [KT] <input type="checkbox"/> Peripheral Blood [01] / <input type="checkbox"/> Amniotic Fluid [02] Array-CGH/ Microarray [AC] <input type="checkbox"/> Peripheral Blood [01] / <input type="checkbox"/> Amniotic Fluid [03] FISH [FH] # <input type="checkbox"/> Single Probe [01]/ <input type="checkbox"/> 2-3 Probes [03]/ <input type="checkbox"/> >4 Probes [02] Specify: _____ PCR-based tests [PC] <input type="checkbox"/> α/β -Thalassemia Reverse Dot-Blot [PR] <input type="checkbox"/> Triplet Repeat Disorders (Fragile X) [PT] <input type="checkbox"/> Paternity Test [PP] <input type="checkbox"/> PCR-MLPA [PM]#: _____ <input type="checkbox"/> Thrombophilia Mutation Screening [07] Group Test <input type="checkbox"/> Infertility & Recurrent Miscarriages Test [FSA01] <input type="checkbox"/> Integrated Tests for Autism [ASD01] Next-generation Sequencing [NG] <input type="checkbox"/> Autism, ADHD & Asperger's Disorder Screening [09]	Oncology Karyotyping [KT] # <input type="checkbox"/> Bone Marrow [02] FISH [FH] (Bone Marrow in Heparin/ Paraffin Section) # <input type="checkbox"/> Single Probe [01]/ <input type="checkbox"/> 2-3 Probes [03]/ <input type="checkbox"/> >4 Probes [02] Specify: _____ PCR-based tests [PC] (Bone marrow in EDTA/ Paraffin section) # Specific Sequencing of any <input type="checkbox"/> 4 sites [96]/ <input type="checkbox"/> 3 sites [97]/ <input type="checkbox"/> 2 sites [98]/ <input type="checkbox"/> 1 site [99] Specify genes and sites: <div style="border: 1px solid gray; height: 40px; width: 100%;"></div> Next-generation Sequencing [NG] # <input type="checkbox"/> Comprehensive Cancer (14 cancer types) [01] <input type="checkbox"/> Breast Cancer [02] <input type="checkbox"/> Colorectal Cancer [03] <input type="checkbox"/> Lung Cancer [04] <input type="checkbox"/> Other Specific Cancer [99]: _____ Circulating Tumor Cells [CTC] # <input type="checkbox"/> Primary & secondary tumour origins/ specific marker(s) requested: <div style="border: 1px solid gray; height: 40px; width: 100%;"></div> Cancer Biomarker Multiplex Screening [BM] <input type="checkbox"/> Liver Cancer Biomarker Multiplex Screening [01]	Genetic Assessment PCR-based tests [PC] Triplet Repeat Disorders <input type="checkbox"/> 4 sites [95]/ <input type="checkbox"/> 3 sites [94]/ <input type="checkbox"/> 2 sites [93]/ <input type="checkbox"/> 1 site [92] Specify Disease: _____ Next-generation Sequencing [NG] <input type="checkbox"/> Comprehensive Cancer (14 cancer types) [01] <input type="checkbox"/> Breast Cancer [02] <input type="checkbox"/> Colorectal Cancer [03] <input type="checkbox"/> Lung Cancer [04] <input type="checkbox"/> Clinical Full Exome Panel [05] <input type="checkbox"/> Comprehensive Cardiovascular Diseases Panel [06A] <input type="checkbox"/> Specific Familial Cardiovascular Diseases Panel [06B] <input type="checkbox"/> Categorized Inherited Disease(s) [07] Specify: _____ <input type="checkbox"/> Full Inherited Disease Panel [08] <input type="checkbox"/> Autism, ADHD & Asperger's Disorder Screening [09] <input type="checkbox"/> Metabolic Diseases Panel [10] <input type="checkbox"/> Sensory Diseases Panel [11] <input type="checkbox"/> Neurological Diseases Panel [12] <input type="checkbox"/> Endocrine Diseases Panel [13] <input type="checkbox"/> Renal Disease Panel [14] <input type="checkbox"/> Mitochondrial Diseases Panel [20] <input type="checkbox"/> Other Specific Cancer [99]: _____ DNASalute™ Health Assessment Next-generation Sequencing [NG] <input type="checkbox"/> DNASalute™ Comprehensive Health Assessment [15] <input type="checkbox"/> DNASalute™ Weight, Management & Physiogenomics [16] <input type="checkbox"/> DNASalute™ Nutrition, Diet Impact & Metabolic Profiles [17] <input type="checkbox"/> DNASalute™ Anti-aging Parameters, Stress Tolerability & Longevity [18] <input type="checkbox"/> DNASalute™ Any 1-3 categories package [19] Specify: _____ Other Tests / Special Requests # <div style="border: 1px solid gray; height: 40px; width: 100%;"></div>
Cardiology PCR-based tests [PC] Sequencing any below <input type="checkbox"/> 4 sites [96]/ <input type="checkbox"/> 3 sites [97]/ <input type="checkbox"/> 2 sites [98]/ <input type="checkbox"/> 1 site [99] Lamin A/C Cardiomyopathy related sites <input type="checkbox"/> LMNA (R225X) <input type="checkbox"/> LMNA (Q354X) <input type="checkbox"/> LMNA (T518fs) <input type="checkbox"/> Other: _____ Familial Hypercholesterolemia related sites <input type="checkbox"/> LDLR (R257W) <input type="checkbox"/> APOB (R532W) <input type="checkbox"/> PCSK9 (R93C) <input type="checkbox"/> CETP (D459G) <input type="checkbox"/> LDLRAP1 (K30fs) <input type="checkbox"/> Other: _____ Hypertrophic Cardiomyopathy related sites <input type="checkbox"/> MYH7 (Specify: _____) <input type="checkbox"/> Other: _____ Next-generation Sequencing [NG] <input type="checkbox"/> Comprehensive Cardiovascular Diseases Panel [06A] <input type="checkbox"/> Specific Familial Cardiovascular Diseases Panel [06B]		

please contact the lab for confirmation and arrangement before ordering

REFERRAL PHYSICIAN / LABORATORY SIGNATURE

Policies, including privacy policy or sample requirement of Hong Kong Cytogenetics and Medical Genetics Centre Limited, which are available on www.hkcmgc.com, apply to this requisition. I hereby confirm the correctness of the above given information and agree with the stated policies.

Physician/ Referral laboratory/ Provider Signature*

Date (DD/MM/YYYY)*