Human Genetic Laboratory
Department of Pathology, Faculty of Medicine Ramathibodi Hospital
Rama 6 Rd. Bangkok 10400
Tel. 0-2201-1267, 1369, 1463-4

For Laboratory Use Only
Lab No. / TID _____
Received By _____
Date _____ Time _____

**Patient Identification**

Patient Name _____

H.N. _____ Age _____ Gender ☐ Male ☐ Female

OPD / Ward _____ Tel. _____

**For Other Hospital /Company Use Only**

Name _____
Tel. _____
Fax. _____

**Clinical Information (Analytical process CANNOT be completed without adequate patient identification, clinical information and specimen type)**

**Clinical Diagnosis / Indication**

Dr. _____
Tel. _____
E-mail _____

**Genetic disorders (Prenatal and Postnatal genetic analysis)** *For PND cases, please fill-in the details below.

Collection date_____ Amount _____ GA by Date _____ GA by U/S _____

PARA _____ Nature _____ LMP _____

| Laboratory Test | Anticoagulant | | รหัสบริการคณะฯตาม | | |
|---|---|---|---|---|---|
| | Heparin | EDTA | AF | CV | CB |
| Prenatal chromosome analysis | | | 300001 | 300002 | 300006 |
| | Heparin | EDTA | PB | CB | HB |
| Genetic disorders chromosome | ✓ | | 300005 | 300006 | 300007 |
| Chromosome breakage analysis for Fanconi anemia | ✓ | | 300328 | | |
| Slide preparation and karyotyping | 300169 | | Other | | |
| Cell maintenance and karyotyping | 300170 | | Please contact the | | |
| Cell culture | 300141 | | officer before ordering. | | |
| VCFS/DiGeorge/CATCH22 (FISH analysis) | ✓ | | 300047 | 300048 | 300050 |
| Williams syndrome (FISH analysis) | ✓ | | 300069 | 300071 | 300072 |
| Prader-Willi syndrome (FISH analysis) (2 probes) | ✓ | | 300088 | 300090 | 300091 |
| Sex chromosome by FISH | ✓ | ✓ | 300018 | 300022 | |
| RB1 gene by FISH | ✓ | ✓ | 300062 | | 300064 |
| SRY gene by FISH | ✓ | ✓ | 300267 | 300268 | 300269 |
| Aneuploidies 13, 18, 21, X and Y chromosome by FISH | ✓ | ✓ | 300224 | 300225 | 300226 |

| Laboratory Test | รหัสบริการคณะฯตามชนิดสิ่งส่งตรวจ | | | |
|---|---|---|---|---|
| | FFPE | Breast | Stomach | Other |
| HER-2/neu gene by FISH | | 300122 | 300123 | 300124 |
| + Unstain slide (ไม่อ่าน) | | 11844 | 11844 | 11844 |
| N-Myc amplification by FISH | 300057 | | | 300058 |
| + Unstain slide (ไม่อ่าน) | 11844 | | | 11844 |
| Neuroblastoma by FISH panel | 300324 | | | 300325 |
| + Unstain slide (ไม่อ่าน) | 11844 | | | 11844 |
| | EDTA | PB | | Other |
| DNA extraction | ✓ | | | 300146 |
| BRCA1 screening by MLPA | ✓ | 300242 | | 300243 |
| BRCA2 screening by MLPA | ✓ | 300244 | | 300245 |
| BRCA1 and BRCA2 screening by MLPA | ✓ | 300246 | | 300247 |
| RB1 deletion/duplication by MLPA | ✓ | 300263 | | 300264 |
| Mitochondrial disease screening by MLPA | ✓ | 300265 | | 300266 |
| Prader-Willi/Angelman syndrome by MS-MLPA | ✓ | 300275 | | 300276 |
| Duchenne muscular dystrophy by MLPA | ✓ | 300240 | | 300241 |
| Von Hippel-Lindau (VHL) disease by MPLA | ✓ | 300331 | | 300332 |
| Spinal muscular atrophy screening by MLPA | ✓ | 300294 | | 300295 |

| Laboratory Test | EDTA | AF | CV | CB | HB | PB | POC | Stool | Other |
|---|---|---|---|---|---|---|---|---|---|
| Stool DNA (SDC2) methylation for colorectal cancer screening | | | | | | | | 300329 | 300330 |
| Prenatal Chromosomal Microarray | ✓ | 300248 | 300250 | 300249 | | | | | 300251 |
| Postnatal Chromosomal Microarray | ✓ | | | | | 300252 | | | 300253 |
| Fragile X syndrome analysis | ✓ | 300345 | | | | 300077 | | | |
| Trio Chromosomal Microarray | ✓ | | | | | 300254 | | | 300255 |
| Exon CNV detection by microarray | ✓ | | | | | 300322 | | | 300323 |
| Rapid detection of aneuploidies 13, 18, 21, X, Y and 9 Microdeletion by **BoBs** | ✓ | 300153 | 300154 | 300155 | 300179 | 300180 | 300181 | | 300156 |
| Rapid aneuploidy screening in all 24 chromosomes by **BoBs** | ✓ | 300157 | 300158 | 300159 | 300182 | 300183 | 300160 | | 300161 |
| Rapid detection of 24 chromosomal abnormalities and 9 microdeletion syndromes by **BoBs** | ✓ | 300164 | 300165 | 300166 | 300184 | 300185 | 300167 | | 300168 |
| Rapid detection of aneuploidies 13, 18, 21, X and Y by **QF-PCR** | ✓ | 300188 | 300189 | 300190 | 300191 | 300192 | 300193 | | 300194 |

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**Hematologic malignancy Tests**

☐ New diagnosis ☐ Follow up

☐ Remission ☐ Relapse

WBC _____cells, Neu _____%,

Lym _____%, Blast _____%

**Clinical Diagnosis**

Collection Date _____Time _____ Last date of chemotherapy_____

**Pre BMT / SCT**

☐ Donor ☐ Recipient

**Post BMT / SCT**

☐ Autologous ☐ Male donor ☐ Female donor

**Requisition status**

☐ Preliminary ☐ Urgent

| Laboratory Test | Anticoagulant | | รหัสบริการคณะฯตามชนิดสิ่งส่งตรวจ | | |
|---|---|---|---|---|---|
| | Heparin | EDTA | BM | PB | Other |
| ☐ Leukemia chromosome analysis | ✓ | | 300009 | 300010 | |
| ALL ☐ Multiplex RT-PCR for BCR::ABL1 p190, TEL::AML1, E2A::PBX1, MLL::AF4 in ALL and CML | | ✓ | 300174 | 300173 | |
| ☐ TEL::AML1 Fusion gene by FISH | ✓ | ✓ | 300074 | 300073 | 300076 |
| ☐ Detection of MLL by FISH | ✓ | ✓ | 300256 | 300257 | 300258 |
| ☐ IKAROS (IKZF1) and Common genetic alterations | | ✓ | 300238 | 300237 | 300239 |
| AML ☐ FLT3 Gene mutation in AML | | ✓ | 300148 | 300147 | |
| ☐ Rapid FLT3 Gene mutation for newly diagnosed AML | | ✓ | 300279 | 300280 | |
| ☐ NPM1 Gene mutation in AML | | ✓ | 300150 | 300149 | |
| ☐ CEBPA Gene mutation in AML by direct sequencing | | ✓ | 300178 | 300177 | |
| ☐ PML::RARA Fusion gene by FISH | ✓ | ✓ | 300060 | 300059 | 300061 |
| ☐ Multiplex RT-PCR for AML1::ETO, CBFB::MYH11, PML::RARA in AML | | ✓ | 300152 | 300151 | |
| ☐ PML::RARA Fusion gene by RT-PCR [bcr1 and bcr3] | | ✓ | 300187 | 300186 | |
| ☐ PML::RARA by quantitative PCR (L-subtype; bcr1) | | ✓ | 300333 | 300334 | |
| ☐ PML::RARA by quantitative PCR (V-subtype; bcr2) | | ✓ | 300335 | 300336 | |
| ☐ PML::RARA by quantitative PCR (S-subtype; bcr3) | | ✓ | 300337 | 300338 | |
| ☐ RUNX1::RUNX1T1 by quantitative PCR | | ✓ | 300339 | 300340 | |
| CML ☐ BCR::ABL1 p210 by RQ-PCR | | ✓ | 300086 | 300087 | |
| ☐ BCR::ABL1 p190 by quantitative PCR | | ✓ | 300273 | 300274 | |
| ☐ Fusion gene for BCR::ABL1 by RT-PCR | | ✓ | 300003 | 300004 | |
| ☐ BCR::ABL1 p190 Mutation detection by direct sequencing | | ✓ | 300341 | 300342 | |
| ☐ BCR::ABL1 p210 Mutation detection by direct sequencing | | ✓ | 300130 | 300129 | |
| ☐ BCR::ABL1 p230 Mutation detection by direct sequencing | | ✓ | 300343 | 300344 | |
| ☐ BCR::ABL1 Fusion gene by FISH | ✓ | ✓ | 300041 | 300040 | 300042 |
| MM ☐ Multiple myeloma by FISH [del (17) and t(4;14)] | ✓ | ✓ | 300171 | | |
| ☐ Multiple myeloma by FISH [1p/1q, del (17), t(4;14) and t(14;16)] | ✓ | ✓ | 300172 | | |
| ☐ Multiple myeloma by FISH [8 probes] | ✓ | ✓ | 300227 | | 300229 |
| ☐ IgH Clone/MRD MM, B-cell malignancies by NGS ☐ Diagnostic ☐ Induction therapy ☐ Initial therapy ☐ Transplant consolidation ☐ Maintenance (please specify the treatment) ☐ Relapse ☐ Supportive care ☐ Other..... | | ✓ | 300277<br>11866 | 300278<br>11866 | |
| MPNs ☐ JAK2 V617F Mutation by AS-PCR | | ✓ | 300126 | 300125 | |
| ☐ CALR (Exon 9) Mutation analysis | | ✓ | 300223 | 300222 | |
| ☐ JAK2, MPL, CALR and KIT mutations in MPNs | | ✓ | 300318 | 300319 | |
| ☐ JAK2 V617F mutation by quantitative PCR | | ✓ | 300320 | 300321 | |
| ☐ FISH for FIPL1::PDGFRA fusion | ✓ | ✓ | 300230 | 300231 | 300232 |
| CLL ☐ CLL panel by FISH | ✓ | ✓ | 300271 | 300270 | 300272 |
| ☐ Deletion of 6q23 by FISH | ✓ | ✓ | 300285 | 300284 | |
| ☐ ATM gene by FISH | ✓ | ✓ | 300287 | 300286 | |
| ☐ Trisomy 12 by FISH | ✓ | ✓ | 300289 | 300288 | |
| ☐ Deletion of 13q14 by FISH | ✓ | ✓ | 300291 | 300290 | |
| ☐ TP53 by FISH | ✓ | ✓ | 300293 | 300292 | |
| ☐ Sex chromosome by FISH | ✓ | ✓ | 300020 | 300018 | 300023 |
| ☐ Deletion of 5q by FISH | ✓ | ✓ | 300327 | 300326 | |
| ☐ Screening of 28 chromosome translocations in leukemia by RT-qPCR | | ✓ | 300262 | 300261 | |
| ☐ DNA Fingerprint | | ✓ | 300014 | 300013 | 300017 |
| ☐ MYD88 L265P mutation by AS-PCR | | ✓ | 300281 | 300282 | 300283 |