



Prenatal Diagnostic Laboratory, Tsan Yuk Hospital

Address: Rm. 210, 30 Hospital Road, Hong Kong Tel: 25892218 Fax: 25172373



REQUEST FORM FOR POSTNATAL STUDIES

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| PATIENT DETAILS Clinic / Hospital No.: _____ <i>(please affix gum label or complete in full)</i> Surname / Last Name _____ 中文姓名 _____ Given Name(s) / First Name(s) _____ HKID / Document ID _____ Date of Birth (DD-MM-YYYY) _____ Age _____ Gender: <input type="checkbox"/> Female <input type="checkbox"/> Male <input type="checkbox"/> Uncertain Relationship to proband: _____ Proband's name and ID: _____ | REFERRING DOCTOR DETAILS Name (Print / Staff Name Chop) _____ Tel: _____ Fax: _____ Address: _____ Referring institute _____ Ward / Clinic _____ |
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| SPECIMEN DETAILS Date & Time of Sampling (DD-MM-YYYY,(HH:MM)) _____ - _____ ,(:) _____ <input type="checkbox"/> Peripheral blood: _____ tubes (2 mL Heparin blood), _____ tubes (3 mL EDTA blood) <input type="checkbox"/> Skin biopsy <input type="checkbox"/> Others, please specify: _____ | Previous Report No. _____ |
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| TESTS REQUESTED <input type="checkbox"/> Karyotype <input type="checkbox"/> Chromosomal microarray <input type="checkbox"/> Genetic testing for _____ <input type="checkbox"/> QF-PCR for chromosome <input type="checkbox"/> UPD testing for chromosome _____ <input type="checkbox"/> 13 <input type="checkbox"/> 18 <input type="checkbox"/> 21 <input type="checkbox"/> Methylation PCR for _____ <input type="checkbox"/> XY <input type="checkbox"/> Del22q11.2 <input type="checkbox"/> Fragile X testing <input type="checkbox"/> FISH for _____ <input type="checkbox"/> Others: _____ | |
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| CLINICAL DETAILS Prenatal diagnosis result (if any): _____ | | | |
| Perinatal History: <input type="checkbox"/> IUGR <input type="checkbox"/> Oligohydramnios <input type="checkbox"/> Polyhydramnios <input type="checkbox"/> Premature delivery <input type="checkbox"/> Others: _____ | Behavioral: <input type="checkbox"/> Autistic behavior <input type="checkbox"/> Autism spectrum disorder <input type="checkbox"/> Obsessive-compulsive disorder <input type="checkbox"/> Oppositional defiant disorder <input type="checkbox"/> Others: _____ | Cardiac: <input type="checkbox"/> Aortic atresia <input type="checkbox"/> ASD <input type="checkbox"/> Atrioventricular canal defect <input type="checkbox"/> Coarctation of the aorta <input type="checkbox"/> Dextrocardia <input type="checkbox"/> Double outlet right ventricle <input type="checkbox"/> Ebstein anomaly <input type="checkbox"/> Echogenic intracardiac focus <input type="checkbox"/> Hypoplastic left heart <input type="checkbox"/> Hypoplastic right heart <input type="checkbox"/> Pulmonary valve atresia <input type="checkbox"/> Situs inversus <input type="checkbox"/> Tetralogy of Fallot <input type="checkbox"/> Transposition of the great vessels <input type="checkbox"/> Truncus arteriosus <input type="checkbox"/> VSD <input type="checkbox"/> Others: _____ | Musculoskeletal: <input type="checkbox"/> Abnormal vertebral morphology <input type="checkbox"/> Abnormal limb <input type="checkbox"/> Club foot <input type="checkbox"/> Diaphragmatic hernia <input type="checkbox"/> Limb contractures <input type="checkbox"/> Polydactyly <input type="checkbox"/> Syndactyly <input type="checkbox"/> Scoliosis <input type="checkbox"/> Others: _____ |
| Growth: <input type="checkbox"/> Failure to thrive <input type="checkbox"/> Growth delay <input type="checkbox"/> Increase body weight <input type="checkbox"/> Obesity <input type="checkbox"/> Short stature <input type="checkbox"/> Others: _____ | Neurological: <input type="checkbox"/> Anomaly of the brain <input type="checkbox"/> Central hypotonia <input type="checkbox"/> Cerebral palsy <input type="checkbox"/> Chorea <input type="checkbox"/> Dystonia <input type="checkbox"/> Gait ataxia <input type="checkbox"/> Muscular hypotonia <input type="checkbox"/> Neural tube defect <input type="checkbox"/> Spasticity <input type="checkbox"/> Others: _____ | Gastrointestinal: <input type="checkbox"/> Gastroschisis <input type="checkbox"/> Hirschsprung disease <input type="checkbox"/> Omphalocele <input type="checkbox"/> Pyloric stenosis <input type="checkbox"/> Tracheoesophageal fistula <input type="checkbox"/> Others: _____ | Genitourinary: <input type="checkbox"/> Ambiguous genitalia <input type="checkbox"/> Hydronephrosis <input type="checkbox"/> Hypospadias <input type="checkbox"/> Kidney malformation <input type="checkbox"/> Undescended testis <input type="checkbox"/> Urethral malformation <input type="checkbox"/> Urethral obstruction <input type="checkbox"/> Others: _____ |
| Developmental: <input type="checkbox"/> Fine motor delay <input type="checkbox"/> Gross motor delay <input type="checkbox"/> Speech delay <input type="checkbox"/> Others: _____ | Craniofacial: <input type="checkbox"/> Cleft lip <input type="checkbox"/> Cleft lip and cleft palate <input type="checkbox"/> Cleft palate <input type="checkbox"/> Coloboma of iris and retina <input type="checkbox"/> Craniosynostosis <input type="checkbox"/> Dysmorphic facial features <input type="checkbox"/> External ear malformations <input type="checkbox"/> Macrocephaly <input type="checkbox"/> Microcephaly <input type="checkbox"/> Others: _____ | Cutaneous: <input type="checkbox"/> Hyperpigmentation <input type="checkbox"/> Hypopigmentation <input type="checkbox"/> Others: _____ | Family History: <input type="checkbox"/> Family history of chromosomal / genetic disorder: _____ _____ <input type="checkbox"/> Parents with ≥ 2 miscarriages <input type="checkbox"/> Relatives with similar clinical history (please explain): _____ _____ |
| Other features: _____ | | | |

Please complete the patient consent form and return it to the laboratory with this request form.

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| Referring doctor's signature: _____ | Request date (DD-MM-YYYY) : _____ - _____ |
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| LABORATORY USE ONLY | Duty Officer: _____ |
| Date and Time of sample receipt: _____ | |