

## Embryology

| System                                     | Normal Development   | Abnormal Development   |
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| <b>General Embryology</b>                  | - Gametogenesis (spermatogenesis, oogenesis) - Fertilization, cleavage, implantation - Germ layer formation (ectoderm, mesoderm, endoderm) - Neurulation and folding - Embryonic and fetal circulation - Placental and umbilical development | - Teratogenesis (environmental, genetic, infectious) - Chromosomal abnormalities (aneuploidy, translocations) - Congenital malformations (multifactorial inheritance)                        |
| <b>Hematopoietic &amp; Lymphoreticular</b> | - Hematopoiesis (yolk sac → liver → bone marrow) - Development of lymphoid organs (thymus, spleen, lymph nodes)  | - Congenital immunodeficiencies (DiGeorge syndrome – thymic aplasia) - Hemoglobinopathies (sickle cell, thalassemia)   |
| <b>Central &amp; Peripheral Nervous</b>    | - Neural tube formation and closure - Brain vesicle development (forebrain, midbrain, hindbrain) - Spinal cord and neural crest derivatives - Myelination and neuronal migration   | - Neural tube defects (anencephaly, spina bifida, encephalocele) - Hydrocephalus, microcephaly, holoprosencephaly - Chiari and Dandy-Walker malformations - Congenital cranial nerve defects |
| <b>Skin &amp; Connective Tissue</b>        | - Epidermis and dermis differentiation - Hair, nails, and sweat gland formation - Development of connective tissues (cartilage, tendons, ligaments)  | - Albinism, ichthyosis, epidermolysis bullosa - Ehlers-Danlos syndrome, Marfan syndrome  |
| <b>Musculoskeletal</b>                     | - Limb bud formation, segmentation - Endochondral and intramembranous ossification - Muscle differentiation and migration  | - Limb abnormalities (polydactyly, syndactyly, amelia, phocomelia) - Achondroplasia, osteogenesis imperfecta, congenital scoliosis   |
| <b>Respiratory</b>                         | - Lung bud formation (week 4) - Bronchial branching morphogenesis - Alveolar development and surfactant production   | - Tracheoesophageal fistula - Pulmonary hypoplasia (e.g., Potter sequence) - Congenital diaphragmatic hernia   |
| <b>Cardiovascular</b>                      | - Heart tube formation and looping - Aortic arch and great vessel development - Septation of atria and ventricles - Fetal circulation changes at birth   | - Congenital heart defects (ASD, VSD, Tetralogy of Fallot, transposition of great arteries) - Patent ductus arteriosus, coarctation of the aorta   |
| <b>Gastrointestinal</b>                    | - Gut tube formation and rotation - Liver, pancreas, and biliary system development - Enteric nervous system formation   | - Malrotation, volvulus, intestinal atresia - Meckel's diverticulum, omphalocele, gastroschisis - Hirschsprung disease (aganglionosis) - Annular pancreas, biliary atresia                   |
| <b>Renal/Urinary</b>                       | - Kidney development (pronephros → mesonephros → metanephros) - Ureteric bud and nephron formation - Bladder and urethra development   | - Renal agenesis, polycystic kidney disease - Horseshoe kidney, vesicoureteral reflux - Urachal abnormalities (patent urachus)   |
| <b>Reproductive</b>                        | - Gonadal differentiation (testis, ovary) - Müllerian (paramesonephric) and Wolffian (mesonephric) duct development - External genitalia formation   | - Disorders of sexual development (androgen insensitivity, Turner syndrome) - Cryptorchidism, hypospadias, epispadias - Müllerian agenesis, uterine malformations                            |
| <b>Endocrine</b>                           | - Development of hypothalamus, pituitary, thyroid, adrenal glands - Pancreatic islet cell formation  | - Congenital adrenal hyperplasia - Thyroid dysgenesis, ectopic thyroid - Diabetes-related genetic syndromes  |