

Most Common Conditions	Description, Epidemiology, and Risk Factors	Pathophysiology
<b>Cerebral Palsy</b>	<ul style="list-style-type: none"> <li>• Most common physical disability in childhood worldwide: 2/1000 live births</li> <li>• Complicated birth, genetic, multiple birth pregnancy, in vitro fertilization, hypoxic brain injury, viral infection</li> </ul>	<ul style="list-style-type: none"> <li>• Different types of cerebral palsy associated with different areas of injury</li> <li>• Spasticity: most common symptom</li> <li>• Abnormal muscle architecture</li> </ul>
<b>Spina Bifida</b>	<ul style="list-style-type: none"> <li>• Folate deficiency, lack of prenatal care</li> </ul>	<ul style="list-style-type: none"> <li>• Neural Tube Defect</li> <li>• Myelomeningocele: most common type</li> </ul>
<b>Traumatic Brain Injury</b>	<ul style="list-style-type: none"> <li>• Bimodal Distribution: 0-2 years and 15-18 years</li> <li>• Abuse, accidents, risky behavior, falls, hyperactivity, electrolyte abnormalities, seizures</li> </ul>	<ul style="list-style-type: none"> <li>• Primary Injury: direct damage, axonal injury</li> <li>• Secondary Injury: Seizures, cerebral edema, hypoxia</li> </ul>
<b>Spinal Cord Injury</b>	<ul style="list-style-type: none"> <li>• Abuse, accidents, risky behavior, falls, hyperactivity, sports injuries, infections, cancer</li> </ul>	<ul style="list-style-type: none"> <li>• More likely to have complete injury</li> <li>• Complete: extensive hemorrhage, cord disruption</li> <li>• Incomplete: contusion, partial cord disruption</li> </ul>
<b>Down Syndrome</b>	<ul style="list-style-type: none"> <li>• Most common inherited genetic syndrome</li> <li>• Trisomy 21</li> <li>• 1 in 691 births</li> </ul>	<ul style="list-style-type: none"> <li>• Complete trisomy of chromosome 21 due to nondisjunction during meiosis</li> </ul>
<b>Spinal Muscular Atrophy</b>	<ul style="list-style-type: none"> <li>• Autosomal Recessive</li> <li>• SMA-1: most common (60-70%)</li> </ul>	<ul style="list-style-type: none"> <li>• Lack a working SMN1 gene--affects motor neuron function</li> </ul>
<b>Duchenne and Becker Muscular Dystrophy</b>	<ul style="list-style-type: none"> <li>• X-linked recessive</li> <li>• Risk factor: born to female carrier</li> </ul>	<ul style="list-style-type: none"> <li>• Abnormal dystrophin gene: connection between the skeletal muscle membrane's dystrophin-associated complex and the actin filaments</li> </ul>
<b>Ehlers-Danlos Syndrome</b>	<ul style="list-style-type: none"> <li>• 1 in 5,000 births</li> </ul>	<ul style="list-style-type: none"> <li>• Genetic defects affecting collagen type I, III, IV</li> <li>• Classic, hypermobility, and vascular subtype most common</li> </ul>
<b>Juvenile Idiopathic Arthritis</b>	<ul style="list-style-type: none"> <li>• Genetic and environmental factors</li> <li>• HLA-A2 and HLA-B27 implicated in some cases</li> <li>• Most common chronic rheumatic disease in children</li> <li>• Distinctions in geography and ethnic groups</li> <li>• Younger age onset in Western Europe</li> <li>• Africa and Eastern Europe: higher rates</li> </ul>	<ul style="list-style-type: none"> <li>• Inflammatory Syndrome characterized by chronic synovial inflammation with B-Lymphocytes</li> </ul>