



DEFINITION

- Group of non-progressive neurodevelopmental conditions
- Characterized by impairments in motor function that limit activity
- Caused by a disturbance to developing fetal or infant brain



CLASSIFICATION

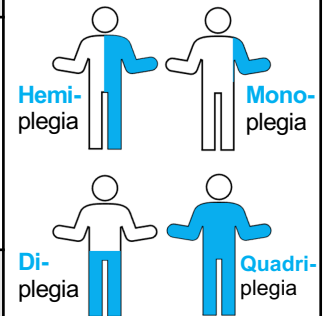
Gross Motor Function Classification System (GMFCS) Levels:

- Walk without limitations (least functional impact)
- Walk with limitations
- Walk using handheld mobility device
- Self-mobility with limitations
- Manual wheelchair (most functional impact)

Motor Types:

spastic, dyskinetic, ataxic, mixed

Affected Limb(s):



DIAGNOSIS

HISTORY

- Inquire about risk factors
- Motor development and milestones
- Comorbidities: epilepsy, learning difficulty, sensory impairments, hip subluxation, scoliosis



PHYSICAL EXAM

- Hypertonia or hypotonia
- Hyperreflexia
- Persistent primitive reflexes: Moro, asymmetric tonic neck
- Spasticity



INVESTIGATIONS

1. Neuroimaging (MRI) for correlation with clinical exam
2. If normal MRI, consider metabolic screen and/or genetic testing



CLINICAL RED FLAGS

- Hand preference <12 months
- Stiffness in legs <12 months
- Inability to sit by 9 months
- Makings fists continuously after 4 months
- Other motor or posture delays



ETIOLOGY

- Genetic
- Birth asphyxia (eg. hypoxic ischemic encephalopathy) ($\leq 10\%$)
- White matter injury
- Intraventricular hemorrhage
- Perinatal stroke
- Infection (TORCH, meningitis)



SIGNIFICANT RISK FACTORS

- Premature birth
- Low birthweight



SUPPORTIVE MANAGEMENT

