Background The clinical spectrum and associated factors of cerebral palsy may differ between developing and developed countries.

Aim To evaluate the predisposing factors, clinical spectrum, and associated problems of cerebral palsy (CP) in children.

Setting and design In this retrospective study patient data were extracted from file records in our center for early diagnosis of childhood disabilities.

Patients and Methods Our study population included one hundred and twenty two children with age range from 7 months to 17 years. Patients were followed and reviewed in a 32 months period from September 2007 to April 2010.

Simple statistical analysis was used for percentage calculation.

Results and Conclusions Spastic type was the predominant (82.7%), with quadriplegic subtype being the most common (34.4%). The other types were choreoathetoid (8.2%) mixed type (6.6%) and ataxic (2.5%) being the least. Speech delay was the most common associated problem (71.3%) followed by mental retardation (61.5%), seizures (35.2%), hearing problems (26.2), and autism (4.9%) being the least.

The clinical spectrum of CP in our country may differ from that reported from the western counties. Prospective studies are needed to evaluate the clinical spectrum and predisposing factors in Jordan.

Obstetrical brachial plexus palsy (OBPP) is a complication of difficult delivery and resulted from excessive traction on the brachial plexus during delivery. Erb palsy, klumpke paralysis and panplexus palsy reported in 46%, 0.6% and 20% of patients, respectively. Unilateral injury is more common than bilateral injury. Risk factors include macrosomia, multiparity, prior delivery of a child with OBPP, breech delivery shoulder dystocia, vacuum and forceps assisted delivery and excessive maternal weight gain. The recovery rate is usually reported to be between 80 and 90%. Management based on medical and surgical interventions. We evaluated 42 children with OBPP. Of them, we could follow only 28 cases during two years. Most of the patients were females. Right side palsy was more prevalent than left side palsy. Vaginal delivery without forceps was the most mode of delivery. Vertex was the most common presentation. Most of the patients were term. The mean weight of the birth was 3.8 kg. Erb palsy and pan-plexus palsy consisted of 71.4% and 28.6% of lesions. Poor to moderate recovery occurred in 13 cases. Good to complete (expected) recovery occurred in 15 cases.

Introduction Narcolepsy often begins in childhood (30% of cases) but is infrequently recognized early in the course; only around 4% are diagnosed before age 15. There have been many reports linking H1N1 and narcolepsy. We present a video series of 4 patients that highlight the diagnostic dilemma in pediatric cases as well as association with H1N1 flu.

Case Reports 4 children (5–11 years) were referred to the Neurologists for excessive daytime sleepiness. Disturbed sleep and cataplexy developed soon after. Immunosuppressive therapy was attempted with no response in one child with rapid progression. Cataplexy initially was focal or partial and progressed to the classical description. The Multiple Sleep Latency Test (MSLT) was grossly abnormal in all them. Hypocretin was absent in the one child investigated. Autoimmune screen was negative in all. One child was treated for H1N1 and 2 had the H1N1 vaccine prior to onset of symptoms.

Conclusion The key to early diagnosis and treatment is recognition of the condition. The videos highlight the facial features and other symptoms that are clues to narcolepsy (Pentad of narcolepsy). Rapid progression is possible related to H1N1 exposure.