Clinico-Etiological Spectrum of Children with Bilateral Basal ganglia lesions: An Observational Study from A Tertiary Care Centre J Dasaratha Ramaiah1 Dr. Lokesh Lingappa1, Dr. Ramesh Konanki1, Dr. Nihaal Reddy2 1-Department of Pediatric Neurology, Rainbow children's Hospital, Hyd 2-Department of Pediatric Neuroradiology, Rainbow Children's Hospital, Hyd Correspondence: <u>siriloki@gmail.com</u>; <u>jdashrath86@gmail.com</u>

INTRODUCTION

Basal ganglia, site of high adenosine triphosphate (ATP) production are affected with various disorders. Observational study aims to analyze etiological and radiological spectrum of this entity.

OBJECTIVE:

To enumerate the radiological and etiological spectrum associated with MRI evidence of "bilateral basal ganglia lesions" among children aged 2 months to 18 years, attending Rainbow Children's Hospital

Inclusion criteria:

Children attending Neuro Outpatient department or admitted at Rainbow Children's hospital with acute neurological symptoms/signs (upto 28 days) having MRI Evidence of Basal ganglia lesions on evaluation.

Age group -2 months to 18 years **Exclusion Criteria**:

Children with clinical history suggestive of perinatal asphyxia (Neonatal encephalopathy, neonatal seizures of any etiology)



¹Cluster 1- T2W HI Putamen Cluster 2 - T2W HI GP OR Inc Susceptibility in GP Cluster 3 -T2W HI, DR GP, BS, Cerebellum Cluster 4 pattern -T1W HI BG

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Conclusion: Most common basal ganglia lesions involve bilateral striatum. Common etiologies are mitochondrial disorders followed by infective and demyelinating diseases. Categorization into cluster patterns narrows down the differential diagnoses.

METHODOLOGY

Study Design: Cross sectional, observational study Study Site: Rainbow Children's Hospital, Hyderabad





1. SURF1 Homozygous mutation 2. PDHA1 Heterozygous mutation 3. Presumed Mitochondrial disorder 4. SLC30A10 Homozygous Mutation

1.Mohammad SS, Angiti RR, Biggin A, Morales-Briceño H, Goetti R, Perez-Dueñas B, et al. Magnetic resonance imaging pattern recognition in childhood bilateral basal ganglia disorders. Brain



T1. 2W HI Putamen



2. T2W HI GP OR Inc Susceptibility in GP

3. T2W HI, DR GP, BS, Cerebellum



4. T1W HI BG

Bibliography