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P4	Golluri Raghuveer	Infantile Arthrogryposis-Renal Dysfunction- Cholestasis (ARC) Syndrome Associated with VIPAS39 gene mutation: A Case report		
P5	Golluri Raghuveer	ARTHROGRYPOSIS -RENAL DYSFUNCTION-CHOLESTASIS SYNDROME		
P6	Yamini Priya K	A case of hereditary spherocytosis in a newborn		
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P8	Chandana A	Clinical Profile and Management Outcomes in a Preterm Neonate with Severe Anemia and Multi-Organ Complications Following Twin Transfusion: A Case Study		
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P23	Aruru Nikhila	An unusual case of pseudoparalysis in a 1 month infant-a diagnostic dilemma
P24	DILEEP KUMAR ALLAGADDA	"Neonatal Jaundice and the Hidden Arrhythmia: The Ritual of Routine Auscultation in Early Diagnosis"
P25	DANDE NAGA MAHESH	Refractory Seizures due to Hypocalcemia (ADH1) - A Case of Autosomal Dominant Hypocalcemia
P26	Gitanjali Jain	Tracing Anhydramnios and Anuria to their Genetic Origin: ACE VariantLinked Renal Tubular Dysgenesis in a Preterm Neonate
P27	Jyothirmai Manasa	" Wiskott-Aldrich Syndrome Presenting as Bilateral CMV Retinitis in a 5-Month-Old Infant: Early Recognition of Primary Immunodeficiency "
P28	Chandana A	Profound Biotinidase Deficiency Unmasked by Whole- Exome Sequencing in a Neonate with a Significant Family History
P29	Vora Shrenik Jitendrakumar	Giant Congenital Melanocytic Nevus with Neuromelanosis, Proliferative Nodule and Malignant Melanoma in a Newborn Infant: A Case Report with Literature Review
P30	Rishee Manoharan	A Rare Syndromic Association: Trisomy 21 with Congenital Diaphragmatic Hernia and Horse Shoe Kidney
P31	Abishek Gowda	Neonatal disseminated HSV masquerading as IEM
P32	Vora Shrenik Jitendrakumar	CONGENITAL NEUROMUSCULAR VARIANT OF GLYCOGEN STORAGE DISEASE TYPE IV PRESENTING AS HYPERTROPHIC CARDIOMYOPATHY
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