Poster Presentation Abstracts

P-1 EVALUATION OF LOWER URINARY TRACT DYSFUNCTION IN CHILDREN WITH ALLERGIC RHINITIS

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P-2 A CASE OF EARLY ONSET SUBACUTE SCLEROSING PANENCEPHALITIS

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P-3 SYRINGOMYELIA AND VITAMIN B12 DEFICIENCY CONCIDENCE CASE PRESENTED WITH HAND AND FINGERS NUMBNESS COMPLAINTS; THE IMPORTANCE OF PHYSICAL EXAMINATION

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P-4 A RARE DISEASE COCKAYNE SYNDROME IN TWO SUBLINGS; CASE REPORT

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P-5 A RARE NEUROCUTANEOUS DISEASE INCONTINENTIA PIGMENTI; CASE REPORT

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P-6 A AGRANULOCYTOSIS CASE DIAGNOSED WITH STURGE WEBER, ASSOCIATED WITH PHENYTOIN TREATMENT

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P-7 ATTENTION TO THE DIFFERENTIAL DIAGNOSIS OF HYPOPIGMENTE LESIONS;TUBEROUS SCLEROSIS AND VITILIGO ASSOCIATION CASE REPORT

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P-8 CULTURE NEGATIVE ENDOCARDITIS CAUSING MITRAL VALVE PERFORATION AND CEREBRAL INFARCTION

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P-9 RICKETS IN CHILDREN WITH CEREBRAL PALSY AND ITS FOLLOW-UP

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P-10 GUILLAIN BARRE SYNDROME VARIANT FACIAL DIPLEGIA AND PARESTHAESIA WHO REFERRED WITH A COMPLAINT OF BILATERAL FACIAL PARALYSIS AND WEAKNESS: A CASE REPORT

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P-11 COMORBITIDY OF PROLONGED JITTERINES, CORTICAL DYSPLASIA AND HETEROTROPIA IN NEONATAL PERIOD: A CASE REPORT

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P-12 GUILLAIN BARRE SYNDROME WHICH DEVELOPED AFTER VARICELLA: A CASE REPORT

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Murat Cansever[1], Alper Özcan[4], Gülşah Uçan[2], Alperen Vural[5], Sevgi Keleş[3], Ekrem Ünal[4], Musa Karakükcü[4], Talal Chatila[6], Türkan Patiroğlu[1]

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P-14 SEVERE PERTUSSIS PNEUMONIA IN AN INFANT: TREATED WITH EXCHANGE TRANSFUSION: CASE REPORT

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P-15 ACUTE LIVER TOXICITY DUE TO ORAL ABSORPTION OF YELLOW PHOSPHORUS: A CASE REPORT

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P-16 TRACHEO-OESOPHAGEAL FISTULA: HOW CAN IT BECOME LATE-ONSET? A CASE REPORT

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P-17 LATE-ONSET HYPERTHROPHIC PYLORIC STENOSIS: A CASE REPORT

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P-18 PEPTIC ULCER DEVELOPING DUE TO CONSUMING EXCESSIVE COLA: A CASE REPORT

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P-19 CO-EXISTENCE OF HEREDITARY SPHEROCYTOSIS AND AUTOIMMUNE HEPATITIS: A CASE REPORT

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P-20 LIVER TRANSPLANTATION TO AN INFANT DIAGNOSED WITH CRIGLER NAJJAR TYPE I FROM THE FATHER WITH GILBERT'S SYNDROME: A CASE REPORT

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P-21 POSTERIOR REVERSIBLE ENCEPHALOPATHY SYNDROME EXPERIENCE AT A PEDIATRIC INTENSIVE CARE UNIT

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P-22 A RARE DIAGNOSIS: CORNELIA DE LANGE SYNDROME

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P-23 ABNORMAL COAGULATION TEST IN A CHILD WITH HODGKIN'S LYMPHOMA

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P-24 CASE REPORT: MICROVILLUS INCLUSION DISEASE

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P-25 PSEUDOTUMOR CEREBRI SECONDARY TO RETINOIC ACID IN CHILDHOOD: CASE PRESENTATION

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P-26 THE ROLE OF HEMODIALYSIS IN THE TREATMENT OF ACUTE CYANIDE POISONING AFTER APRICOT KERNEL EATING

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P-27 AN INCIDENTAL DIAGNOSIS OF MORGAGNI HERNIA

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P-28 ALSTROM SYNDROME: A RARE CASE OF SYNDROMIC OBESITY

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P-29 DYKE-DAVIDOFF-MASSON SYNDROME-CASE REPORT

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P-30 IMMUNE THROMBOCYTOPENIA AND HEMOLYTIC ANEMIA IN A CHILD WITH HODGKIN LYMPHOMA

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P-31 A CHARACTERISTIC RASH: ERYTHEMA MULTIFORME

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P-32 CLINICAL AND RADIOLOGICAL FINDINGS OF GRANULOMATOUS APPENDICITIS

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P-33 URGENT SURGERY OR CONSERVATIVE TREATMENT? CHILAIDITI SYNDROME – THE CONFUSION BETWEEN THE IMAGE OF SUBDIAPHRAGMATIC BOWEL AND FREE AIR UNDER THE DIAPHRAGM

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P-34 CARDIAC INVOLVEMENT IN ANOREXIA NERVOSA

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P-35 A RARE REASON FOR ACQUIRED THROMBOTIC THROMBOCYTOPENIC PURPURA: CHRONIC PANCREATITIS

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P-36 AN INTERESTING FOREIGN BODY IN THE ESOPHAGUS: A CLOTHES-PIN COIL

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P-37 PRESENTATION OF A CASE WITH SYSTEMIC COLD URTICARIA

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P-38 ABDOMINAL TUBERCULOSIS IN PATIENT WITH FOCAL SEGMENTAL GLOMERULOSCLEROSIS: CASE REPORT

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P-39 CHILDHOOD ONSET BEHCET'S DISEASE: TWO CASES REPORT

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P-40 A REASON FOR RECURRENT VOMITTING IN NEWBORN PERIOD: PROPIONIC ACIDEMIA, CASE REPORT

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P-41 A NEWBORN WITH UREA CYCLE DISORDER: CASE REPORT

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P-42 A RARE CAUSE OF HYPOTONIC INFANT – CONGENITAL MYASTHENIC SYNDROME 16 (CMS16): A CASE REPORT

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P-43 SECONDARY PSEUDOHYPOALDOSTERONISM

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P-44 THINNING HAIR AND BRITTLE NAILS: BIOTIN DEFICIENCY

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P-45 MICROBIOLOGIC AGENTS THAT ISOLATED FROM PATIENTS WITH FEBRILE CONVULSION

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P-46 UNUSUAL CLINICAL PRESENTATION OF FAMILIAL MEDITERRANEAN FEVER IN TWO CASES

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P-47 DILEMMA IN SYSTEMIC JUVENILE IDIOPATHIC ARTHRITIS: MACROPHAGE ACTIVATION SYNDROME OR SEPSIS?

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P-48 RAMSAY HUNT SYNDROME; CASE REPORT

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P-49 XANTHINE STONE

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P-50 DIFFERENT PHENOTYPES OBSERVED IN SIBLINGS IN THE SAME FAMILY: KRABBE DISEASE AND CUTIS LAXA

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P-51 EEC SYNDROME CASE REPORT

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P-52 A RARE CAUSE OF RECURRENT COUGH AND RESPIRATORY DISTRESS: SCIMITAR SYNDROME

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P-53 GIANT CARDIAC MASS IN LEFT VENTRICLE, RHABDOMYOMA?: CASE REPORT

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P-54 A RARE COMORBIDITY OF GUILLAIN BARRE SYNDROME AND TYPE 1 DIABETES MELLITUS: A CASE REPORT

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P-55 A COMORBITITY OF ACUTE DISSEMINATED ENCEPHALOMYELITIS AND CYSTOPERITONEAL SHUNT DYSFUNCTION IN INCREASED INTRACRANIAL PRESSURE SYNDROME: A RARE CASE REPORT

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P-56 AN IMPORTANT REASON OF SUDDEN VISUAL LOSS OPTIC NEURITIS

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P-57 A CASE OF CARDIAC THROMBOSIS DUE TO HEREDITARY PROTEIN C / S DEFICIENCY

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P-58 IS VITAMIN D DEFICIENCY ASSOCIATED WITH DISBIOSIS IN BOWEL FLORA?

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P-59 MOYAMOYA DISEASE WITH RECURRENT CRANIAL INFARCTION

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P-60 PULMONARY ALVEOLAR PROTEINOSIS ACCOMPANIED BY ATOPY: CLINICAL CASE REPORT

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P-61 SECKEL SYNDROME

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P-62 A RARE CAUSE OF CENTRAL CYANOSIS: PULMONARY ARTERIOVENOUS MALFORMATION

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