Humoral immunity in children with chronic tonsillitis

Introduction

The aim of our study was to evaluate and compare the humoral immunity in children and adults with chronic tonsillitis.

Subjects

We examined 24 children and 15 adult patients with chronic tonsillitis.

Methods

Immunologic examination included analysis of: total leucocytes; absolute and relative lymphocytes; C reactive protein (CRP), antistreptolysin O (ASO), rheumatoid factor; investigation of humoral immune factors: absolute and relative B lymphocytes, serum IgA, IgM, IgG, circulating immune complexes, interleukin amount - IL-4, IL-5, IL-18 in blood serum.

Results

Increasing level of the absolute lymphocytes amount (2.92 ± 0.24, 2.2 ± 0.11) (p = 0.04), interleukin - 8 (229.09 ± 47.63; 35.05 ± 14.64) (p = 0.047) and interleukin - 1 β level (191.19 ± 70.44; 10.88 ± 4.28) (p = 0.045) were observed in children. Average indices of interleukin-8 and interleukin - 1 β in children is in 6.5 and 17.4 times higher than in adults.

Serum IgA level in adults was significantly higher than in children (1.36 ± 0.16; 1.92 ± 0.19) (p=0.038). An antistreptolysin-O titre in adults in comparison to children (162.5 ± 45.53; 216.67 ± 45.78) is higher in 1.4 times.

Conclusions

Nonspecific defense factors play an important role in children with chronic tonsillitis, while in adults the main role is held by specific immune response. Comparative analysis of anti-inflammatory cytokine synthesis by immune competent cells in chronic tonsillitis proved the fact that in children it is much higher than in adults.

AUTOIMMUNE LYMPHOPROLIFERATIVE SYNDROME IB: IMPROVEMENT WITH RAPAMYCIN

Autoimmune lymphoproliferative syndrome (ALFS) is a disorder on a defect in the apoptosis of lymphocytes with lymphoproliferation and immune dysregulation. Type Ib, defined by mutation in the gene that encodes the protein FAS-ligand, is a rare entity.

Case report

A 9-month-old male infant was referred because a failure to thrive and abdominal distention. Physical examination showed pallor, signs of severe malnutrition, axillary and inguinal lymph nodes, hepatomegaly and giant splenomegaly. Family background: parents, cousins of Moroccan origin, brother died at the age of 4 when he was under study for a giant visceromegaly. Complementary tests highlighted severe anemia (Hb 7.80 g/dl) and thrombocytopenia (platelets 76000/mm3), paraveretbral lymphoid proliferation of 4 cm, elevation of IL10, soluble CD25, 28% T lymphocytes double negative (CD4 - and CD8-) in peripheral blood and cell culture with defect of apoptosis in one of the samples. The genetic study identified the mutation of the gene TNFSF6 which encodes FAS-ligand-protein. After the diagnosis of ALFS type Ib, treatment with rapamycin was started at doses of 2 mg/m2 daily. An optimal evolution was observed, with a reduction of visceromegal size after 30 months of treatment, without adverse events by the time.

Conclusion

We emphasize the importance of the suspicion of this entity in children with chronic visceromegaly, especially with family history. Despite the few existing data on treatment with rapamycin for this disease and children in general, we have seen an appropriate response and a good tolerance in this patient.

A CASE REPORT OF KARTAGENER’S SYNDROME ASSOCIATED WITH NASAL POLYPS

Nasal polyps are benign nasal masses that can cause nasal obstruction, headache and snoring. The overall incidence or prevalence of nasal polyposis is unknown. They are diagnosed more frequently in men and during the third and fourth decades of life. Nasal polyps are rare in children. Most clinical data indicate that there is no greater prevalence of nasal polyps among atopic compared with normal populations.

A 14-year-old male patient was referred by an otolaryngologist to investigate the cause of nasal polyposis. There were otitis media, sinusitis and bronchial asthma to patient’s history therefore he used inhaled corticosteroid for control long term asthma, sometimes bronchodilator to asthma attack and antibiotic treatment for sinusitis. No family member had similar respiratory complaints or any significant systemic illness.

His physical examination was found above 97% for weight and height difficulty breathing through nose, bilateral nasal polyposis, rarely sibilan ronchi on pulmonary auscultation and his heart sound was heard deeply. Laboratory findings; Hemogram was normal, IgA:189mg/dl, IgE 203 IU/L, inhalen sIgE and epidermal prick test were negative, Chest X-Ray showed dextocarcia, A computed tomograph scan of the chest showed situs inversus with lateral segment of right middle lobe tubular bronchiectasis, peribronchial thickening and atelectasis, the right lower lobe tubular bronchiectasis. Pulmonary function testing demonstrated a mixed obstructive and restrictive pattern.

The pathogenic findings of Kartagener Syndrome are dextocardia, bronchiectasis and sinusitis. His findings were consistent with Kartagener Syndrome. This patient were presented due to nasal polyposis associated with Kartagener Syndrome and delayed diagnosis.

CHRONIC RECURRENT SEVERE LIP ANGIOEDEMA IN YOUNG CHILD SECONDARY TO ALLERGIC RHINITIS

Background and Aims

Allergic rhinitis (AR) is a significant disease that affects children and adults and often under-diagnosed which has a deleterious impact on quality of life. Chronic recurrent severe lip angioedema secondary to AR was not previously reported.