Successful Therapy with intravenous gamma globulin in two children with postinfectious bronchiolitis obliterans

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Bronchiolitis obliterans (BO) is an infrequent clinical syndrome characterized by the chronic obstruction of small airways due to fibrosis [1]. Intravenous immunoglobulin (IVIG) could be used for treatment while underlying immune mechanisms in the pathogenesis of BO exist [2]. Here, we present two children with BO due to adenovirus infection whose complaints resolved after IVIG replacement.

CASE 1

A 21-month-old boy was admitted with the complaints of hacking cough, wheezing, difficulty in breathing for 4 months. From the history; he needed mechanical ventilation due to a severe adenoviral pneumonia (adenovirus PCR was positive on nasal swab), was diagnosed as bronchiolitis obliterans (BO), 1mg/kg methylprednisolone was continued for 4 weeks, was discharged to home with oxygen replacement, and hospitalized 5 more times in a 4 month period. His weight was 10600gr (10-25p), height was 83.5cm (25p). On physical examination, there were tachpnea, central cyanosis, bilateral rales and rhonchi. Chest computerized tomography demonstrated mosaic pattern and ground glass opacities. Immunodeficiency, cystic fibrosis, allergic hypersensitivity and tuberculosis were excluded (Data not shown). Methylprednisolon treatment (1mg/kg/day) was restarted, but he had to be hospitalized for three times during reduction period of steroid. Steroid dosage had to be increased during hospitalizations. Azithromycin prophylaxis was started 3 times a week during reduction period of steroid, and continued for 5 months, but clinical improvement could not be achieved. Angiotensin converting enzyme inhibitor was started while his pulmonary artery pressure was 29 mmHg. The family was convinced to move to another house due to intense moisture and mould. While clinical improvement couldn’t be achieved, intravenous immunoglobulin G (IVIG) replacement every 3 weeks for a dose of 400 mg/kg was started when he was 30 month-old. During 6 doses IVIG replacement, only one hospitalization was needed at winter time, and his need for oxygen was disappeared. The replacement of IVIG was stopped when he was 35 month-old. After termination of IVIG replacement, he was hospitalized for 2 times when he was 38 and 40 month old, so IVIG replacement was given for 3 times more. He is now 72-month old, takes inhaled corticosteroids, azithromycine prophylaxis for 6 months in a year. He has no cough, tachypnea, exercise intolerance, rales, and need for oxygen anymore. The last pulmonary artery pressure was detected as 20mmHg. His body weight: 16.5 kg (50-75p), height: 104 cm (75p). Radiological imagings were shown in figure 1.
CASE 2

A 4 month-old boy was admitted with the complaints of cough, wheezing, difficulty in breathing for 2 months after Influenza-A and Respiratory syncytial virus-B bronchiolitis. From the history; maternal smoking was present. On physical examination, there were tachypnea, central cyanosis, wheezing, bilateral rhonchi. Immunodeficiency, cystic fibrosis, allergic hypersensitivity and tuberculosis were excluded (Data not shown). Chest computerized tomography demonstrated mosaic pattern and ground glass opacities. Inhaled budesonide, oral methylprednisolone (1 mg/kg/day), gastroesophageal reflux treatment were started. Oral steroid was continued for two weeks and stopped after a reduction period. But the symptoms reappeared. He had totally 2 months oral methylprednisolone treatment until he reached 7.5 months old and had to be hospitalized for after every cessation period of steroid. Subcutaneous immunoglobulin G (SCIG) replacement was started (0.12 mg/kg/dose every week) when he was 7.5 month-old. He is now 10 month-old and he never needed systemic corticosteroid treatment and hospitalization after SCIG replacement. His complaints revealed totally and he needed inhaled corticosteroid intermittently.

Bronchiolitis obliterans (BO) is a severe form of chronic obstructive lung disease that results from an insult to the lower respiratory tract [3]. The clinical scenario consists of chronic respiratory symptoms for more than 4 to 8 weeks in a previously healthy child that occurs after an episode of a severe acute respiratory infection. Patients present with decreased pulmonary compliance, increased airway resistance, elevated air trapping, and, in general, poor or no response to bronchodilators. Adenovirus has been identified as a cause of post infectious BO (PIBO) in up to 50% of the cases [3].

The treatment for BO has not been definitely established in the literature. The treatment protocol for PIBO included oral prednisone and azithromycin [4]. Li et al showed that treatment with corticosteroids and azithromycin was effective in 85.7% of cases [3]. Oral corticoids can be given in divided doses of 1-2 mg/kg/day, four weeks later the dose is gradually decreased to a single early morning dose of 0.5 mg/kg/day, and the total course of steroids is 6 months. There are reports of beneficial effects of pulse therapy with methylprednisolone (10 mg/kg/day or 25-30 mg/day, maximum 1 gr, IV bolus 3 days/month totally 6 cures) in the initial phases of the PIBO, before fibrosis is established [5]. High doses of inhaled steroids can also be administered. In the most severe cases of lung damage, a lung transplant may be the best recourse [1,3].

Azithromycin is a modulator of neutrophil activity. It suppresses chemotaxis in inflammatory region, acts as an anticytokine by inhibiting IL-8, IL-1β, IL-6, TNFα, prostaglandin, and even improves pulmonary function [6,7]. Azithromycin is given
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orally at 5 mg/kg/day, once daily for 3 consecutive days per week, or 10mg/kg/day, every other day (3 times a week) for 6 months [6]. Based on these recommendations we also administered azithromycine 10 mg/kg 3 times a week in winter time to our patient.

Some current treatments were based on the aim to control the inflammation because of the role of the immunological response in the physiopathology of this disease. Mycophenolate mofetil, sirolimus, IL-2 receptor antibodies, inhaled cyclosporine A, cyclophosphamide, plasmapheresis, statin, thalidomide, alentuzumab (anti-CD52 antibody), tumor necrosis factor antagonists are other treatment choices specifically used for BO developed after bone marrow transplantation which were reported usually in case reports [8]. Almost all the treatment modalities tried till now can cause secondary immunodeficiency and none of them could achieve satisfying results. So we tried an alternative treatment modality which can be supposed as a safe one. The empirical monthly administration of IVIG has been used as a treatment option only in one case who had a favorable response to IVIG infusions and a 3-day regimen of pulse IV methylprednisolone (30 mg/kg per daily dose), both given monthly for 6 months [2]. The pathological findings of our patients also resolved after IVIG replacement treatment, too. Although it is hard to decide for which patients IVIG is an optimal treatment choice, we suggest that it can be used before the other immunosuppressive treatment modalities other than corticosteroids.

The precise mechanism by which IVIG suppresses harmful inflammation has not been definitively established. The massive quantity of antibodies may stimulate the host’s complement system, leading to enhanced removal of all antibodies, including the harmful ones. Intravenous immunoglobulin G also blocks the antibody receptors on immune cells (macrophages), leading to decreased damage by these cells or regulation of macrophage phagocytosis. Immunoglobulin binds to the Fc receptors on surfaces of B and T cells, prevents entrance of antigen to the cell and antibody response, so immunomodulation effect occurs [9]. Anti-inflammatory effect is by decreasing the cytokines and other proinflammatory mediators released from monocyte/macrophages. IL-6, IL-8, IL-1Ra, TNF-alfa, IL-2, IL-10, interferonγ (IFNγ) ve tumor necrosis factor (TNFβ) levels decrease significantly after IVIG infusion [10]. Improvement can be achieved in our patients with one or more of these mechanisms. While only three cases reported including ours are not enough to insist on the effectiveness of IVIG for BO, further studies are necessary in order to decide whether IVIG could be an alternative choice for BO and for which patient groups it can be chosen.

SUMMARY

In conclusion, early diagnosis and treatment, and avoidance of repeated respiratory tract infection may be helpful to improve the prognosis. We suggest that intravenous or subcutaneous immunoglobulin G treatment may offer some benefit for PIBO patients by decreasing the number and severity of infections, and its immunomodulatory effects.

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AUTHORSHIP AND CONTRIBUTORSHIP

Sevgi Pekcan is responsible for writing the article. Bahar Gokturk is responsible for writing and correcting the article. Ismail Reisli is responsible for correcting the article.

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