Case report / Приказ болесника

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Drug rash with eosinophilia and systemic symptoms syndrome in an adolescent – efficiency of immunoglobulin G in a corticosteroid resistant case

Синдром осипа на лек праћен еозинофилијом и системским симптомима у адолесцента – ефикасност имуноглобулина Г у кортикостероид резистентном случају

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INTRODUCTION

Already we are aware of the mechanisms of the hypersensitive reaction to carbamazepine (IVb/c) in the clinical form titled Drug rash with eosinophilia and systemic symptoms (DRESS) syndrome (sy) [1, 2, 3]. The heterogeneous clinical presentation of DRESS sy [4, 5] is always a challenge, especially if the administration of carbamazepine for five to six weeks was accompanied by the use of Na-valproate and metformin (previous seven months), all preceded by the infection with herpes simplex virus type 1 (HSV1) [6]. There have been questions occurred as to whether (in these circumstances) 1/adverse reaction...
to the drug is purely pharmacological (type B), i.e., pharmacological interaction with the immune receptors (pi) [7, 8] or, is it predominantly immune resulting from the loss of the control mechanism of the immune system due to the previous contact with viruses, and 2/why the efficacy of corticosteroids [9] is unsatisfactory so that the successful control of the clinical picture of DRESS syndrome achieved with the application of class G immunoglobulins (Ig)? In this study, we presented a 16-year-old patient whose DRESS syndrome has been successfully resolved with IVIG.

**CASE REPORT**

A 16-year-old adolescent girl was hospitalized due to the fever up to 38.5oC during the previous day, elevated transaminases, discrete lymphadenopathy, splenomegaly, whereas subjectively in good general condition. In the past few years, the adolescent was treated for Na-valproate epilepsy, but since the seizures were being reported in recent months, carbamazepine was been added five to six weeks prior to this hospitalization. Insulin resistance was determined seven months earlier, which is the reason of her regular using metformin.

During the first seven days of hospitalization, the adolescent fevers up to 38.5oC mainly in the evening, and every day there is an eruption of skin efflorescence (predominantly in the form of erythematous rash (partly maculopapular) and partly in the form of urticates on the trunk, upper and lower extremities, neck, occasionally on the face, followed by itching, without accompanying bullae, and is present clinically until the ninth day of hospitalization. Edema of the face manifests itself on the fifth to seventh day of hospitalization. During the first nine days of hospitalization, lymphadenopathy in the neck is pronounced, bilateral, along the sternocleidomastoid and submandibular, with nodules up to two cm in diameter, arranged in a row, painless and mobile relative to the substrate. During
this period, the adolescent had normal vital functions including adequate diuresis but easily adynamic, which is due to the disease and the use of double doses of antihistamines (cetirizine).

Hematological and biochemical analyzes performed upon admission showed leukopenia (3.9x10^-9/l) with 21% monocytes and 10% eosinophils (380/ml absolute number), CD4/CD8 ratio 1031/63 seven/ul (1.62), total CD3 1 seven/ul, then high values were determined of transaminase AST 164IU/l, ALT 125IU/l, gamma-GT 780IU/l, lactic dehydrogenases 925IU/l, alkaline phosphatases 553IU/l, B-type natriuretic peptide 1395pg/ml, CRP 2 seven/mg/l, and at the same time, reduced values of fibrinogen-C 1.9g/l, activated thromboplastic time 22.4s, serum IgG concentration 6.64g/l, and vitamin D 10. sevenng/ml. The consumption of IgG in addition to normal serum IgE, IgA, IgM values was followed by the increase in C3 complement components 2.8seven/g/l and C4 0.59/g/l.

On the fifth day of hospitalization, the number of eosinophils increased to 12% (seven20/ml absolute number) and on the sixth day of hospitalization to 24% (780/ml absolute number), while at the same time, monocytosis was maintained at 1 seven%. Both disorders in blood cells number are normalized by day 16 of hospitalization. Platelet and erythrocyte numbers are always within normal limits. Splenomegaly is pronounced on the ninth day of hospitalization (134x82mm), and normalized up to the 16th day of hospitalization, and hepatomegaly is pronounced on the 16th day of hospitalization (AP diameter 160mm), and normalized to the 24th day of hospitalization.

Serologic testing revealed an elevated IgG antibody titer for HSV1, while within the normal range there were the titers of other antibodies (HSV1-IgM, HSV2-IgG and IgM, as well as IgG- and IgM- for Epstein-Barr virus, heterophilic antibodies, cytomegalovirus, toxoplasma, parvoB19, hepatitis A, C, HBsAg, Mycoplasma pneumoniae, antistreptolysin titer, lupus antibodies LAC and LAC-SCT). The following biochemical analyzes from serum
were within the reference values: erythrocyte sedimentation, procalcitonin, prothrombin time, international normalized ratio, D-dimer, ferritin, gas analyzes, ionogram, glycemia, hemoglobin-A1c, urea, creatinine, proteins, albumin troponin-hs-I, creatinine kinase, muscle creatinine kinase, valproic acid level, thyroid stimulating hormone, free thyroxine, antinuclear antibodies. No nasal and pharyngeal bruising revealed pathogenic germs, not did the examination of the stool reveal intestinal parasites and Giardia lamblia. Urinalysis, occult bleeding stools, lung, and heart X-rays, heart and kidney and pancreatic ultrasound findings, spirometric findings, pulmonary, cardiac, and infectological clinical examinations were all within the reference values. The neurological finding was unchanged compared to the previous period.

DISCUSSION

Significant clinical, hematologic and biochemical improvement occurred the day after the first dose of IVIG ie. IVIG therapy was a life-saving in this case [10,11]. Actually, we started treatment according to RegiSCAR scoring [5] ie. 6/6 criterias for DRESS sy were established and according to this, the treatment was performed with a corticosteroid at a dose of 1g/kg body weight (methylprednisolone) intravenous (iv) for eight days with continued antihistamine (cetirizine was administered at twice the dose from the regular dose) as well as with the elimination diet, withdrawal of carbamazepine on day sixth of hospitalization and increased doses of Na-valproate at 1500mg/day. As the described clinical picture is maintained until the ninth day of hospitalization, and despite the eight days administration of methylprednisolone (iv), the treatment was continued with iv IgG (IVIG, human normal immunoglobulin for intravenous use), even though the indication was not justified by data from controlled clinical studies but based on case reports [10, 11, 12]. Two doses of IVIG were administered at 0.4 g/kg in eight-day intervals. There was no desired response to
methylprednisolone during the first eight days of neither treatment nor prednisolone during further treatment, with concomitant administration of antihistamines from day one of hospitalization, Na-valproate, metformin hydrochloride, elimination diets, and carbamazepine withdrawal. No adverse effects were observed after administration of two doses of IVIG.

The systemic corticosteroid acts nonspecifically on the tissue by inhibiting the local immune response of the tissue to various stimuli and injuries but does not block the release of mediators [13], which is the rationale for the ineffective administration of corticosteroids in the present case. Whereas, according to modest literature data [6,14], IVIG modulates cytokine production, complement cascades, turnover of B and T cells and neutralization of autoantibodies, thus explaining the effect of treatment and improving the clinical picture the day following administration of the first IVIG dose case.

Fowler indicates clear gaps in our current understanding of DRESS sy, and points to the need for old test (patch) and new diagnostic tests to screen patients before starting the carbamazepine (interleukin 15 and microRNA 122 in the serum) and that certain authors favor IVIG and plasma exchange instead of steroids for the treatment of DRESS sy [18]. Korea authors found approximatively 17% children with adverse skin reactions to antiepileptic drugs (most commonly with aromatic ring in chemical structure) and significant associations with genes encoding the human leukocyte antigen alleles (e.g. HLA-B*15:02) which indicates the need for genetic testing [19]. Korea authors published results of national study and consideration that IVIG monotherapy or the combination of corticosteroids and IVIG might reduce the mortality rate in severe DRESS sy related to antiepileptic drugs [20].

In conclusion, we join the authors [6, 10–18, 20] of who favor the treatment of DRESS sy using IVIG, and we point out to the need to change the order of recommendations in the current treatment recommendations for DRESS sy [5, 9] in favor of IVIG. IVIG should be considered as a second-line treatment as soon as possible in patients in whom
corticosteroid treatment failed. There is not enough experience for recommendation that IVIG should be used as the first line treatment. We advocate for an early introduction of IVIG as early as the third or fourth day from unsuccessful corticosteroid treatment. Further, it is necessary that similar case reports should be collected. Only when new reports on the same topic arrive, corticosteroid shall be considered as a second-line treatment, what would have next consequence - the identification of DRESS sy in children, through pharmacopoeias worldwide, as a proven indication area for IVIG, what we advocate. Finally, we invite the professional and scientific public to specify, through the following case reports and well-designed controlled clinical studies, the dose of IVIG for the DRESS sy indication area in children, especially caused by carbamazepine, and in light of previous and current viral infection. Certainly, the culprit drug should be immediately discontinued.

Informed consent was obtained from the patient’s parent prior to her participation in treatment and was informed of the introduction of each drug throughout treatment. This paper was done in accordance with the institutional Committee on Ethics.

**Conflict of interest:** None declared.
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