Bone Marrow Aplasia in a Patient Treated with Cefotaxime

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Description of the first case of aplastic anemia caused by cefotaxime, a third-generation cephalosporin, with lethal outcome is presented. Clinical record of a patient and post-mortem examination results are given. Possible mechanisms of cytopenias induced by cephalosporin antibiotics are discussed.

Keywords: cephalosporin antibiotics, aplastic anemia, agranulocytosis


One of the adverse drug reactions (ADR) during the treatment with cephalosporin antibiotics (CA) are various cytopenias (thrombocytopenias, neutropenias/agranulocytosis, hemolytic anemias and, less often, pancytopenias). Cephalosporin-induced cytopenias can be based on two mechanisms – an immune and a myelotoxic one. Most publications on hematologic ADR in the course of treatment with CA, and ceftriaxone in particular, deal with the cases of neutropenias and agranulocytosis (AC) development. Analysis of these descriptions gives evidence of the possible (antibody-mediated) mechanism of ceftriaxone-induced AC [1]. One of the reports analyzes 14 verified cases of ceftriaxone-induced agranulocytosis. The main mechanism for the development of agranulocytosis is the immune mechanism, since in most cases there was no inhibition of erythropoiesis and thrombocytopoiesis [2]. However, bone marrow (BM) involvement was found in none of the cases, which was taken notice of in other – earlier – studies. Myelotoxic effect of β-Lactam antibiotics was discovered during the experimental studies in the course of in vitro granulopoiesis and monocytopoiesis cell culture, and along with this, a dose-dependent effect, most prominent with ceftriaxone and ceftazidime, was found [3].

In the analysis of cases of neutropenia caused by cephalosporin antibiotics (ceftaroline), the relationship between the duration of drug use and the frequency of neutropenia was established. The bone-marrow (BM) hemopoiesis picture was characterized by the absence of highly-differentiated granulocytic lineage cells, with the presence of numerous granulocyte progenitor cells...
We observed a patient who developed aplastic anemia (AA) during the treatment with cefotaxime. The clinical observation follows.

Patient P., 79 y.o., was delivered to resuscitation and intensive care unit № 2 of S.S. Yudin City Clinical Hospital (Moscow) on 18.10.17 with the complaints about throat-ache and swallowing difficulty, body temperature risen up to 38 °C, loose stool and weakness. The patient fell ill two weeks before when throat-aches and hoarseness started to disturb him. He was diagnosed with pharyngolaryngitis and therefore put on cefotaxime i.m., and Hexoral and Furacilin topically, without effect. The temperature rose up to 38 °C, hemorrhagic brash and loose stools appeared. On the day of hospitalization, the patient had a choking fit due to which he was hospitalized. Condition on admission is severe. The consciousness is clear, the patient is active. The skin is pale, warm and dry. Sporadic skin hemorrhages on the limbs. Small papules rash in the collar zone. Peripheral lymph nodes are not enlarged. Subcutaneous fat tissue is normally developed. Peripheral edemas are absent. Body temperature is 38.6 °C. Meningeal symptoms are negative. The respiration rate per minute is 19. Auscultation findings: harsh breathing, no rales in the lungs. The region of respiration is without changes. Heart borders are not recognized, which corresponded the dependence of reticuloctysis and thrombocytopenia, the most probable AC mechanism should be hemopoiesis depression. The bone marrow cytological screening revealed cellular depletion and considerable decrease of megakaryocyte amount, which, in the absence of blast cells, enabled to exclude acute leukemia and diagnose with AA.

The patient’s state remained very severe. CT of thoracic organs diagnosed multisegmental pneumonia. Antibacterial therapy (Amoxiclav and Metronidazole), Noradrenaline, Sterofundin, Fragmin, Cordarone and erythrocyte transfusions were assigned, artificial lung ventilation was started. Despite the conducted therapy, the patient died on 20.10.17.

Clinical diagnosis. Main diagnosis: Drug-induced aplastic anemia.


BM histology results (Figure) has shown: Lipocytes prevail in the bone marrow; the erythroid lineage is substantially narrowed and represented by sporadic small clusters with prevailing erythrokaryocytes. Sporadic granulocytes, small lymphocytes, separate plasma cells and siderophages are defined. Megakaryocytes are absent. Hemorrhage and sclerosis areas in the stroma.

Thus, hematologic condition of pancytopenia with related clinical signs (fever, hemorrhagic syndrome, aphthous stomatitis, enteropathy) together with the...
Bone marrow hemopoiesis depression and prevailing fat tissue in the BM (results of BM morphological examination) testified of the patient’s having AA. Taking into account acute disease development, absence of anamnestic data about the peripheral blood changes in the past, appearance of clinical symptoms after taking cefotaxime with the absent indications for administering other drugs, cefotaxime should be considered the most probable ethiological agent of AA development. As an AA mechanism in the course of treatment by CA, one can assume the drug’s depressant action on the pluripotent stem cell common for erythroid, granulocytic and thrombocytic hematopoietic lineages [5]. It can be supported by the data about depressing the activity of not only colony-forming units of granulocyte-macrophage progenitor cells, but also those of erythroid lineage in dogs that received high doses of cefazedone (a first-generation CA) for 4 months.

In the literature we managed to find the description of the only case of cefroxadine-induced (a second-generation CA) AA in a 50-year-old patient, who after seven days of urinary tract infection treatment with cefroxadine had a temperature rising up to 38°C, with the AC, anemia and thrombocytopenia found in the blood. The BM histology discovered rapid hemopoiesis depression and prevailing fat tissue, which was regarded as AA [6]. It is notable that this patient had a similar situation with AA development earlier, several years before the current episode, with the following hematopoietic regeneration. The authors are of the opinion that this case is the first description of AA in the course of treatment with CA. However, together with CA, the patient received other drugs – NSAIDs and antihistamines in particular, which does not allow excluding their role in the AA development. Other three cases of AA in the course of treatment with cephalaxin are mentioned in unpublished messages of FDA while analyzing 11301 cases of various ADRs in the course of treatment with cefalexin. It should be noted that these patients did not do without concomitant treatment (NSAIDs and antihistamines) either, which makes the ethiological role of CA in the AA development ambiguous.

Conclusions. As we did not manage to find any publications about the cases of AA in the course of treatment with cefotaxime, it would be justified to consider our observation the first message about cefotaxime-induced AA. This observation confirms the necessity of hematological monitoring of patients receiving CA, especially upon the presence of pancytopenia, which requires a timely BM examination.

Disclosures:
The authors declare no conflict of interest.

References

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