Quetiapine Induced Autoimmune Hemolytic Anemia in a Child Patient: A Case Report

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Autoimmune hemolytic anemia (AIHA) is a disease characterized with destruction of erythrocytes as a result of antibody produce against patient’s own erythrocytes and anemia. AIHA can be roughly stratified into two groups according to serological features and secondary causes including drugs induced hemolytic anemia. Drugs induced autoimmune hemolytic anemia is very rare in pediatric patients. Even though hematological side effects such as leucopenia, agranulocytosis, eosinophilia, thrombocytopenic purpura and aplastic anemia might occur due to psychotropic drug use; to the best of our knowledge there is no autoimmune hemolytic anemia case due to quetiapine, an atypical antipsychotics, in literature. We hereby describe the first child case of autoimmune hemolytic anemia during quetiapine treatment. We also are pointing out that one should keep in mind serious hematological side effects with atypical antipsychotic drug use with this case report.

KEY WORDS: Autoimmune hemolytic anemia; Antipsychotic agents; Quetiapine fumarate; Child; Hematological side effects.

INTRODUCTION

Autoimmune hemolytic anemia (AIHA) is a disease characterized with destruction of erythrocytes as a result of antibody produce against patient’s own erythrocytes and anemia. AIHA can be roughly stratified into two groups according to serological features and secondary causes including drugs induced hemolytic anemia.1) All drugs, including psychotropic drugs, have potential to cause hemolysis and hemolysis in patients might present earlier and become fatal.2,3)

Hematological side effects of psychotropic drugs such as atypical antipsychotics are rare, but important. These hematological side effects are usually leucopenia, agranulocytosis, eosinophilia, thrombocytopenic purpura and aplastic anemia. Uncommonly, leukocytosis, thrombocytosis, thrombocyte function disorders and coagulopathies might occur.4-6) Side effects of atypical antipsychotics over erythrocytes are limited. To the best of our knowledge there is no AIHA case due to atypical antipsychotics in literature.

Hereby, we report a 12-year-old boy with mental retardation (MR) and attention deficit hyperactivity disorder (ADHD) who developed AIHA with quetiapine, an atypical antipsychotic, use. Even though drug related hemolytic anemia is rare in pediatric cases, we wanted to point out that quetiapine has such side effect and hemolysis persevered a long time even after discontinuation of the drug.

CASE

A 12-year-old male patient who has been followed-up with MR and ADHD in our clinic for 6 years was receiving risperidone for his aggression, impulsivity, insomnia and hyperactivity complaints for 5 years. Patient experienced a convulsion even though he has no history of previous neurological disease and presented to pediatric neuro-
logical outpatient clinic. Discontinuation of risperidone was recommended because there was no other pathology except disturbance in electroencephalogram (EEG) and these EEG disturbances and convulsion were attributed to risperidone. His laboratory test results showed these results; hemoglobin (Hb) 12.4 g/dl, red blood cell (RBC) 4.42 M/μL, white blood cell (WBC) 7.77 K/μL, platelet 183 K/μL, total bilirubin 0.7 mg/dl, direct bilirubin 0.3 mg/dl, indirect bilirubin 0.4 mg/dl, lactate dehydrogenase (LDH) 296 U/L. After discontinuation of risperidone, quetiapine 25 mg/day was initiated for complaints of the patient. Patient’s complaints of insomnia got better after changing evening dosage of quetiapine to 50 mg/day at the 4th day of treatment. Patient presented to pediatric emergency department with weakness and jaundice at the 8th day of treatment. In his laboratory tests these results were observed; Hb 9.6 g/dl, RBC 3.21 M/μL, reticulocyte (RET) %2.01, WBC 6.81 K/μL, LDH 307 U/L, total bilirubin 3.8 mg/dl, direct bilirubin 0.8 mg/dl, indirect bilirubin 2.88 mg/dl. In his physical examination general medical condition was moderate, his sclera was icteric and his abdomen was slightly sensitive. Patient was referred to pediatric hematology department for research of the etiology with initial diagnosis of hemolytic anemia after discontinuation of quetiapine. Patient was diagnosed as hemolytic anemia and treatment was initiated in hematology department after test results of Hb 9.6 g/dl, RBC 3.21 M/μL, WBC 6.81 K/μL, LDH 307 U/L, RET %2.01, total bilirubin 4 mg/dl, direct bilirubin 0.8 mg/dl, indirect bilirubin 3.2 mg/dl. Patient was hydrated and metilprednisolone (2 mg/kg/day), nerox tablet (TB), and folbiol TB 2×1.5, folbiol TB 3×1 week, nerox TB 1×0.5 and discharged with a recommendation of weekly follow-up. Steroid treatment was reduced and discontinued at 145th day due to negative result of direct coombs test. At 145th day of treatment his laboratory test results were; Hb 12.4 g/dl, RBC 3.72 M/μL, total bilirubin 0.6 mg/dl, direct bilirubin 0.3 mg/dl, indirect bilirubin 0.7 mg/dl, direct coombs test (+++) positive. Because clinical findings were improved, his treatment changed to oral metilprednisolone 16 mg 2×2. At 21st day of treatment his laboratory test results were; Hb 12.4 g/dl, RBC 3.72 M/μL, total bilirubin 0.43 mg/dl and indirect bilirubin 1.29 mg/dl. Steroid treatment of patient was reorganized to oral metilprednisolone 16 mg 2×2. At 21st day due to negative result of direct coombs test. At 145th day of treatment his laboratory test results were; Hb 12.4 g/dl, WBC 8.67 K/μL, total bilirubin 0.6 mg/dl, direct bilirubin 0.14 mg/dl, indirect bilirubin 0.46 mg/dl, direct coombs test (−).

**DISCUSSION**

Even though psychotropic drug-related hematological side effects are so rare, they shouldn’t be overlooked because they might threaten the life. Most of hematological side effects caused by psychotropic drugs were explained with different pathophysiological mechanism such as bone marrow suppression, destruction of immune-associated cells, active or toxic metabolite formation and direct bone toxicity.

Hematological side effects of psychotropic drugs including atypical antipsychotics effect mostly thrombocytes and leucocytes. The effects on erythrocytes are limited. In literature, there are pancytopenia cases of clozapine, quetiapine and risperidone use, while there was also a case of hemolytic uremic syndrome related with
thrombocytopenic purpura due to quetiapine use. However, we didn’t find any atypical-antipsychotic-related hemolytic anemia in our literature review.

Quetiapine, which is an atypical antipsychotic, is dibenzo thiazepine structure and has dopamine-2 (D2) receptor antagonism just like any other antipsychotic-active drug. It has a greater affinity for 5-HT2 receptors than for D2 receptors; also has an affinity for H1 and alpha 1 and alpha 2 adrenergic receptors. Quetiapine that shows structural similarities to clozapine and olanzapine, rank 3rd in terms of hematological side effects after these two typical antipsychotics after considering case reports in the literature. There are case reports that suggest development of agranulocytosis, neutropenia and thrombocytopenic purpura due to quetiapine. Our patient didn’t have any abnormal clinical or laboratory test results except EEG disturbances before initiation of quetiapine treatment. Patient’s hematological and biochemical parameters were in normal range before quetiapine treatment. Because our patient didn’t use any other medication except quetiapine and other causes of hemolytic anemia were excluded with history, physical examination, clinical findings and laboratory tests, hemolytic anemia of patient was considered due to AIHA related with quetiapine use.

Drugs induced hemolysis may occur as immune-related or non-immune-related. Three main mechanisms were explained in drug-related immune hemolysis; First mechanism, which is also known as hapten, is a type of hemolysis due to absorption of drug into erythrocyte membrane and gain immunological features, thus antibodies against this complex causes hemolysis. It is mostly extravascular, progresses slowly, rarely causes severe disease and regresses after discontinuation of drug. Second mechanism, which is also known as complex type, is a type of hemolysis caused by destruction of erythrocyte membrane with immunological and non-immunological paths after formation of a complex of antigens, which include drug and plasma proteins, and antibodies. Hemolysis is intravascular and usually ends 1-2 days after discontinuation of the drug. Third mechanism, which is also known as autoimmune type (alfametildopa), is a hemolysis caused due to immune response to erythrocyte antigens by drug (stimulated autoimmune). Hemolysis occur slowly with long-term use of drug and regresses slowly after discontinuation of the drug. Hemolysis type of AIHA in our patient might be considered as alpha methyl-dopa because of long-lasting erythrocyte destruction. In this type, drug should be discontinued when hemolysis is clinically apparent. Even though it usually takes a couple days to resolve, it might also last a few months. Laboratory tests and clinical findings of our patient improved at 21st day of treatment however his direct coombs test turned negative at 145th day of treatment.

Hematological side effects usually occur at the start of the therapy. Our patient also had hematological side effects at the first week of quetiapine treatment. If drug-related hemolytic anemia as have mild clinical findings, discontinuation of the drug might be enough but in serious cases such as ours, it is important to initiate supportive treatment with corticosteroid therapy. We used corticosteroids in line with supportive treatment in our case.

In conclusion, we believe quetiapine was responsible for erythrocyte destruction in our case. However, it is unclear whether quetiapine or its metabolites cause this clinical picture and whether hemolysis occurred with or without dose-dependent manner. To our knowledge, this case is the first case to report quetiapine-related AIHA and one should always keep in mind that atypical antipsychotics might cause several hematological side effects including hemolytic anemia. In addition, we present only a child patient of quetiapine-induced AIHA in this case report, therefore there is a need for further clinical studies to determine the relationship between atypical antipsychotics such as quetiapine and AIHA.

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