A Case of Antithyroid Drug-Induced Agranulocytosis Pre-Covid 19 Era

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ABSTRACT

Agranulocytosis is an infrequent and serious side effect of antithyroid drugs characterized by a noticeable reduction in granulocyte and neutrophil count, it usually occurs within the first 2-3 months of treatment. There is a variety of mechanisms by which ATD can induce agranulocytosis, direct drug effects, and immunological mechanisms. We present 33 years old female attended Atbara teaching hospital who has developed agranulocytosis 2 weeks after starting ATD to treat relapsed Graves’ disease. What was unusual about this patient is that symptoms have occurred in a period less than 15 days of starting treatment and with a dose of 45 mg/day. The physician must educate the patient about the possibility of early onset of serious side effects of ATD and to seek medical advice as soon as possible.

Keywords:
Agranulocytosis, Neomercazole, Hyperthyroidism, Graves’ disease

Abbreviations

Introduction
Basedow’s (Graves’) sickness is an autoimmune disease characterized by hyperthyroidism due to circulating autoantibodies. Thyroid immunoglobulin bind to active thyrotropin receptors causing the thyroid gland to grow and thyroid follicle to increase the synthesis of thyroid hormone [1]. According to the American thyroid association radioactive iodine, thyroidectomy and antithyroid drugs are the standard management for Graves’ disease [2] Carbimazole is a prodrug which is converted into Methimazole (the active form), it’s used for the treatment of hyperthyroidism totaling a dose of 30-60 mg/day, but the maximum dose should be 120 mg/day [3] MZ and CZ are associated with an adverse side effect, one of them Agranulocytosis which is rare but can be fatal [4]. Agranulocytosis is characterized by a marked reduction in granulocyte and neutrophil count, it usually occurs within the first 2-3 months of treatment [5].

Clinical Case
33 years old female patient attended to Atbara teaching hospital who is a known case of Graves’ disease underwent subtotal thyroidectomy 14 years ago she has been on thyroxin 50 mg once daily she remained euthyroid all that time till the last month when she developed palpitation, weight loss despite increase appetite, sweating, menstrual irregularity, and irritability. thyroid function test showed (TSH 0.02 µIU/ml, free T3 5.6 ng/ml, T4 2.1 ug/ml) patient was diagnosed as relapsed graves’ disease and received Neomercazole 15 mg three times a day and B blocker for symptoms control, then 15 days after initiation of ATD patient has developed a severe sore throat and high-grade fever, she sought medical advice at local ENT center. On physical examination, there was tonsillitis and pharyngitis and lab result revealed low TWBCs 1.3 × 10^3/µl count and absolute neutrophil count was 0 cell/ul and CRP was 106. Blood test demonstrated low TWBC 1.3 × 10^3/µl count and absolute neutrophil count was 0 cell/ul and CRP was 106. Urine general was normal. Based on history, examination and lab result CZ induced Agranulocytosis was strongly suspected. Accordingly, CZ was immediately stopped and broad-spectrum antibiotics were prescribed. On the follow up 3 days later her symptoms improved, CBC showed that TWBCs 3.8 × 10^3/µl count and

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Discussion

CT2 and MZ2 have widely used medications for the treatment of Graves disease. They act by inhibiting thyroperoxidase enzyme which is essential in thyroid hormone synthesis CT2 and MZ2 both have the same mechanism of action, they are considered identical [4]. Propylthiouracil was used for a long time until the US Food and Drug Administration (FDA) added a warning that there is an increased risk of severe liver injury and acute liver failure, and it may be fatal so it is only reserved for patients who can’t tolerate other treatment options such as MZ2 or surgery [6] there are several adverse effects associated with MZ2 and CT2 ranging from mild (pruritus and urticaria) to severe (Agranulocytosis and Stevens-Johnson syndrome) [4].

There are a variety of mechanisms by which ATD can induce Agranulocytosis: it may be due to the direct toxic effect of the drug when ATD penetrates the bone marrow; through there, oxide metabolites affect oxygen and glucose utilization of the leukocytes leading to their destruction [7]. However other studies contradict the hypothesis of the direct effect of the drug on leukocytes they suppose that carbimazole-induced agranulocytosis is found to be occurring through a drug mediated reaction rather than the direct toxic effect of the drug [8].

Wall et al. study reported that in vitro peripheral lymphocyte transformation and circulating Antibodies against neutrophils have been detected in a patient with ATD induced agranulocytosis [9].

The immune-mediated destruction of mature neutrophils was the first mechanism to be identified as a cause of ATD induced Agranulocytosis [10]. Sprikkelman et al. described different four immunological mechanisms that can be responsible [10].

Firstly, the destruction of granulocyte due to antibodies formed against ATD and bound to the granulocyte membrane leading to an acceleration of granulocyte destruction. Secondly, the target of the antibodies may be the drug -metabolites complex that has been absorbed into the neutrophil granulocyte in the presence of the plasma component. Thirdly, Autoantibodies may be triggered by the drug, and finally, the production of antibodies may be induced by the interaction between the drug and the granulocyte antigen.

Also, other immunological reactions may occur like induced hypersensitivity reaction mediated by IgE, drug-induced IgG and IgM responses, and associated immune injury by anti- neutrophil cytoplasm antibodies ANCA which may contribute to Agranulocytosis [11-13].

The relationship between the dose of ATD and the incidence of Agranulocytosis is debatable and age-related. a study made by Yasuda et al on pediatric patients with Graves’ disease found that the incidence of a variety of adverse events occurs more often in patients who are on a high dose of methimazole (0.7 or more mg/kg/day) than those on a low dose(<0.7 mg/kg/day) with frequencies of adverse events being 50% and 20% respectively. However, neutropenia was found to manifest independently of dose [14]. As we can see from our case agranulocytosis has occurred with a low dose of ATD 15 mg per day. On the other hand, Cooper et al study concluded that methimazole doses greater than 30mg/day and or age >40, are associated with a higher risk of development of agranulocytosis [15]. Leukopenia in patients with hyperthyroidism may be caused by thyrotoxicosis itself or may be related to ATD [16]. ANC rarely fall below 2000 x 10^3/µl, this is neither associated with an increase in the risk of infection nor necessitate drug discontinuation [17-18].

The majority of ATD induced Agranulocytosis usually occur within 2 months of treatment [15-19]. the clinical presentation is usually infection in the oropharyngeal region, fever and sore throat are the most common presenting symptoms [20-21] our case presented with typical manifestations of ATD induced agranulocytosis; sore throat and fever.

In our case, it occurred within the first 2 weeks of treatment. Usually, it takes two weeks after stopping the offending drug for leukocyte count to return to normal [22]. However, The restoration of TWBCs from 1.3 x 10^3 /µl to 3.8 x 10^3 /µl and Absolute neutrophil count was rapid and dramatic in our case it took only 3 days after discontinuation of the ATD, even without administration of Granulocyte colony-stimulating factor, however, there case report in Taiwan Ming-Tsung et al. [23] the recovery time of TWBCs took only 3 days.

Conclusion

Agranulocytosis is a rare serious side effect of ATD, that can occur at any dose and as early as 15 days, in our case, it is clear that the patient wasn’t vigilant about the onset of serious side effect of her prescribed drugs; we have noticed that she went to the ENT clinic considering her symptoms as symptoms of ordinary throat infection. That could lead to delay of appropriate treatment and raise the risk of the onset of serious complications like sepsis, peritonsillar abscess, or retropharyngeal and consequent upper airway obstruction. Fortunately, she is referred to her original physician early and appropriate management was done. Finally, what we have learned from this case is that when the physician prescribes ATD for a patient he/she should warn the patient in written or verbal means that he or she should seek physician advice and obtain blood analysis as soon as he or she feels sore throat or common cold symptoms.

References