ACECLOFENAC-INDUCED STEVENS-JOHNSON SYNDROME AFTER ONE SINGLE DOSE: A MAIDEN CASE REPORT

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INTRODUCTION

Medical treatment is very challenging as every day the treating physician experiences new challenges in treating various conditions as well as encounters new adverse drug reactions to medications. Adverse cutaneous drug reactions are very common in clinical practice, but it often goes unreported. Adverse cutaneous drug reaction is defined as any reaction that causes damage to skin, skin appendages, and mucous membrane and all adverse effects related to drug eruption are included irrespective of the etiology [1]. Nonsteroidal anti-inflammatory drugs (NSAIDs) are known to cause many adverse reactions which include nausea, vomiting, dyspepsia, gastric ulcers, and cutaneous reactions pruritus, morbilliform rashes, urticaria, and photosensitivity [2]. Sometimes, NSAIDs cause life-threatening conditions such as Stevens-Johnson’s syndrome and toxic epidermal necrolysis which are very rare and only few cases have been reported so far in the literature [3]. Hence, we report a rare case of Aceclofenac-induced Stevens-Johnson’s syndrome in a tertiary care patient in Southern India.

CASE REPORT

Informed consent was taken from the patient. The patient was treated by a general practitioner with paracetamol and Aceclofenac on December 25th, 2016 for severe fever and body pains with myalgia. Later, December 26th, 2016 patient came to our hospital with complaints of oral lesions with fluid-filled blisters all over the body with skin peeling over the upper chest and trunk. Multiple purpuric macules were also noted all over the trunk with additional fluid-filled bullae over the face/nails and scalp. The patient also had complaints of burning sensation in eyes associated with lacrimation and redness along with burning sensation of palms and soles. History also revealed one episode of vomiting. No history of similar episodes in the past or any allergy. The patient was not on any other medications. Immediately, tablet Aceclofenac was suspected as the cause of adverse drug reaction ruling out other causes. Diagnosis of Aceclofenac-induced Stevens-Johnson syndrome was made and the patient was treated with fusidic acid ointment and fluticasone ointment thrice daily to be applied locally till the lesions subside. He was also started with ciprofloxacin eye drops along with other supportive measures. Moreover, the patient was asked to review after 2 weeks. The patient showed significant improvement on his next visit.

DISCUSSION

Stevens-Johnson syndrome is very rare and is said to be a life-threatening skin condition. Incidence ranges from 2.6 to 7.1 persons per million populations per year in the United States [4]. Various drugs are implicated in causing Stevens-Johnson syndrome and incidence ranges from 75 to 90% for drug-induced etiology [5]. The most common drugs implicated include NSAIDs, anti-epileptics, and antimicrobials. Based on the body surface area involved Stevens-Johnson syndrome (SJS) and toxic epidermal necrolysis (TEN) is classified with former involving < 10% and later involving > 30%, respectively [5]. Sometimes, both these conditions overlap together which is called SJS-TEN overlap. It is diagnosed usually with the help of clinical signs and histopathological diagnosis. Usual mechanisms include reactive drug metabolites and immunological reactions. Interactions between FAS (CD 95) and CD95 ligand are said to play a major role. This interaction takes place in the epidermal layer of the skin. TEN is more common in women compared to men while the reverse is true in case of SJS. Antibiotics have an incidence of 34% in causing SJS followed by analgesics with an incidence of 33% [6].

Aceclofenac is a NSAIDs mainly used in the management of pain in rheumatic disorders. It acts by inhibiting the cyclooxygenase enzyme system thereby inhibiting prostaglandins synthesis. It helps in reducing inflammation as well as in pain control. It is associated with side effects which include gastrointestinal (GI) disturbances, paresthesias, vertigo, and tremor. It is also known to cause cutaneous adverse reactions ranging from rashes, pruritus, urticaria, and hypersensitivity reactions [7]. There are only few case reports on Aceclofenac-induced Stevens-Johnson’s syndrome/toxic epidermal necrolysis [8,9]. According to severe cutaneous adverse reactions to drugs, study group piroxicam and tenoxicam had the highest risk of SJS/TEN with diclofenac and ibuprofen having the lowest risk. According to a study by Lapeyre-Mestre et al., Aceclofenac caused many adverse reactions including GI disturbances and cutaneous reactions, but liver toxicity was found to be the most common reported adverse reaction [10]. There are many prognostic factors for SJS, but those of paramount importance include age at diagnosis, extent of skin necrosis, and serum urea level [9]. Mortality with SJS was found to be around 1 to 3% compared to higher mortality rates observed with TEN [9]. A large multicenter study also revealed early administration of glucocorticoids together with withdrawal of offending agent helps in better treatment prognosis and patient survival rates [9].
In our patient, the lesions appeared within 1 day of treatment with Aceclofenac. Other causes of SJS/TEN were ruled including history of intake of drugs known to cause SJS/TEN. Aceclofenac was stopped immediately and the patient was started on Wysolone 20 mg along with proper treatment of skin lesions along with supportive measures which included electrolyte management. Causality assessment was done using Naranjo's scale [11], and a probable causal relationship was established. The adverse drug reaction was found to be moderately severe and preventable as per Hartwig's scale and Thornton's scale, respectively [12,13].

CONCLUSION
Since SJS/TEN is a life-threatening cutaneous reaction leading to mortality, proper care of the patient along with stopping the offending agent after ruling out other causes should be of foremost importance. Further, clinical trial can be done in India to find out the incidence of SJS/TEN associated with NSAIDs.

CONFLICTS OF INTEREST
The authors declare that they have no conflicts of interest.

AUTHOR'S CONTRIBUTION
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REFERENCES