Ketoprofen, an emerging photoallergen

Ewelina Bogumiła Zuba¹, Agnieszka Osmola-Mankowska², Dorota Jenerowicz², Maciej Stawny³, Magdalena Czarnecka-Operacz²

¹ Heliodor Święcicki University Hospital, Poznan University of Medical Sciences, Poland
² Department of Dermatology, Poznan University of Medical Sciences, Poland
³ Department of Pharmaceutical Chemistry, Poznan University of Medical Sciences, Poland

ABSTRACT

Introduction. Ketoprofen, which belongs to the group of non-steroidal anti-inflammatory drugs (NSAIDs), is an emerging photoallergen. Especially its topical use may be a cause of drug-induced photosensitivity.

Material and methods. We report two cases of photoallergic and one case of phototaggravated contact dermatitis due to topical ketoprofen application, confirmed by photopatch testing.

Results. All patients presented positive reactions to ketoprofen at an irradiated site. Only one patient demonstrated a positive reaction to ketoprofen both at an irradiated and non-irradiated site.

Conclusions. Photosensitive reactions due to topical application of ketoprofen being of significant clinical importance need to be properly diagnosed. It is crucial to provide patients with a detailed instruction how to protect photoexposed areas during therapy with ketoprofen.

Keywords: ketoprofen, photoallergic contact dermatitis, photoallergy, photosensitivity, photopatch testing.

Introduction

Ketoprofen belongs to the group of non-steroidal anti-inflammatory drugs (NSAIDs). It is widely used because of its well-known analgetic and anti-inflammatory properties. Photoallergic dermatitis is one of possible side effects of ketoprofen. In fact ketoprofen is considered to be one of the main photocontact allergens. The first case report of photoallergic reaction due to topical ketoprofen was published by a Spanish dermatologist in 1985 [1]. Since then, photoallergy to ketoprofen has been reported mostly by researchers originating from Mediterranean countries [2] as well as from Japan [3, 4], Belgium [5], Sweden [6] and recently also from Poland [7].

In this study, two cases of photoallergic contact dermatitis and one case of phototaggravated contact dermatitis due to topical use of ketoprofen have been presented.

Material and methods

Case 1

A 48-year-old female patient was directed to the Department of Dermatology, Poznan Medical University of Sciences in May 2011, with initially suspected erysipelas. The patient demonstrated itchy, erythematous skin lesions localized within the right calf and erythematous as well as exfoliating skin lesions within the left palm. There was neither fever nor malaise and new eczematous skin lesions on both lower extremities, neck and decollete appeared. Therefore diagnosis of erysipelas has been excluded. The patient demonstrated itchy, erythematous, papular skin lesions localized within the right calf and erythematous as well as exfoliating skin lesions within the left palm. There was neither fever nor malaise and new eczematous skin lesions on both lower extremities, neck and decollete appeared. Therefore diagnosis of erysipelas has been excluded. The patient reported on applying ketoprofen gel on her legs due to myalgia and previous exposure to the ultraviolet light while working in a garden during sunny weather, wearing short trousers. Our treatment was composed of clemastine 2 mg iv, dexamethasone 8 mg iv, 10 mg oral cetirizine, and topical betamethasone combined with gentamycin.
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in a cream formulation. Further on 1% hydrocortisone cream has been applied. Within the time period of nine days of the treatment significant improvement has been obtained.

Case 2
A 38-year-old female patient with a history of erythromelalgia related to idiopathic trombocythemia, experienced erythematous and papular eruption on both lower legs, where she had previously applied ketoprofen gel to reduce burning pain associated with erythromelalgia. Further exacerbation of skin lesions has been reported by the patient after the sun exposure.

Case 3
A 22-year-old male developed itchy, erythematous and papular skin lesions with exudation, restricted to the site of application of ketoprofen. It has been prescribed by the general practitioner in order to reduce the contusion-related pain within the neck and left upper thorax. Highly itchy, erythematous and papular skin lesions with significant exudation, restricted to the site of the drug application appeared after 24 hours. A few hours after the first application of the medicine, the patient visited an outdoor swimming-pool.

Patch and photopatch testing
In case of all patients after remission of skin lesions diagnostic testing has been performed. The diagnostic set consisted of standard patch tests (European Standard Patch Test Series, 28 haptens) provided by Chemotechnique Diagnostics. According to the International Contact Dermatitis Research Group recommendations evaluation of results has been performed after 48 and 72 hours.

Minimal erythemal dose (MED) was determined using mono-chromator irradiation simulator test and photopatch tests have been performed according to the European Guideline [8]. Medications tested included: diclofenac 1%, ibuprofen 5%, ketoprofen 2.5%, naproxen 5% (all in white petrolatum, which was also the negative control). Two photopatch series have been applied symmetrically on the back of patients for 48 hours, after which both were removed and one site has been irradiated with UVA at the dose of 5 J/cm². Test reactions have been evaluated 24 and 48 hours after irradiation. Results have been assessed according to the recommendations of International Contact Dermatitis Research Group.

Results
In case of all patients response to UV light was normal in MED testing. Patch tests’ results with the European Standard Set of contact allergens were all negative. In all cases after 24 hours results of photopatch tests with ketoprofen were positive (erythema and papules). In cases one and two, positive reactions presen-
ted the ‘crescendo’ pattern with an increasing response (erythema, papules and vesicles) 48 hours after irradiation. Patch tests result with ketoprofen (at a non-irradiated site) was positive only in the case of patient three (++) in both readings. Based on results of our diagnostic procedure, patients one and two were diagnosed with photoallergic contact dermatitis and patient number three – as phototaggravated contact dermatitis. Results of photopatch testing have been presented in Figures 1–3.

Discussion

The two most common agent groups currently responsible for photoallergic contact dermatitis are organic ultraviolet absorbers in sunscreens and topical NSAIDs.

Figure 2. Photopatch testing of a 38-year-old woman

Figure 3. Photopatch testing of a 22-year-old man
Ketoprofen is considered as the most common cause of NSAIDs-induced photosensitivity. It may promote phototoxic, photoallergic as well as phototaggraved reactions, where photoallergy is much more rare than phototoxic phenomenon [11]. Both topical and oral forms of ketoprofen are commonly prescribed by doctors of all professions.

Foti et al. [12], presented two cases of photodermatitis due to oral ketoprofen in patients with a history of photocontact reactions to topical ketoprofen. Both patients developed eczematous skin lesions after oral intake of ketoprofen and subsequent sun exposure. The authors highlight the risk of photoinduced reactions to systemic ketoprofen in patients with previous sensitisation to topical form of this drug.

Conti et al. [13] reported a case of a patient who developed acute eczematous reaction affecting the right lip commissure, and extending to the upper and lower lip and chin region after administration of oral granulated ketoprofen (OKI) and subsequent sun exposure. The patient never experienced any cutaneous reaction due to systemic ketoprofen, but only when the granules came in contact directly with the perioral skin followed by exposure to the sun. A positive reaction to only photostimulated ketoprofen was observed.

Recently published studies [14, 15] demonstrated that photoallergy to ketoprofen is often associated with photoallergy to octocrylene, and benzophenone-3, which are compounds of sunscreens. Therefore, patients, who have experienced a photoallergic reaction to ketoprofen, should be informed about importance of avoiding sun blockers, which contain octocrylene and benzophenone-3.

Although photocross-reactivity to ketoprofen, octocrylene and benzophenone-3 is well-known, such a cross-reactivity between ketoprofen and butyl methoxydibenzoylmethane, a compound of sunscreen preparations, was not reported. Recent research [16] has demonstrated that the addition of a UVA absorber to topical ketoprofen formulations may be effective to reduce the ketoprofen photosensitivity. The authors presented that butyl methoxydibenzoylmethane has a strong potential to reduce ketoprofen photosensitivity. From the viewpoint of cross-reactivity, it is considered that butyl methoxydibenzoylmethane may be one of the best choices as a UVA filter added to topical ketoprofen formulations to prevent photosensitive reactions to ketoprofen.

According to recommendations of the European Medicines Agency’s Committee for Medicinal Products for Human Use (CHMP) [17], ketoprofen is not available over the counter in Poland due to significant risk of severe photosensitivity reactions and cross allergy with octocrylene. Medical professionals should inform their patients about photosensitive reactions being a highly possible side effect of ketoprofen. Patients should also protect treated skin areas against UV during topical therapy with ketoprofen and two weeks after termination of the treatment.

On the basis of the results of the European Multicentre Photopatch Test Study [9], and on the presence or absence of photosensitizing agents in consumer products within the European market, a recommended European Baseline photopatch test series has been changed. Ketoprofen, etofenamate, piroxicam, benzodamine, promethazine, and 15 compounds of sunscreens, including octocrylene, and benzophenone-3, are included in the new standard European Baseline photopatch test series [18].

**Conclusion**

Medical specialists should be aware, that avoiding prescriptions as well as selective use of topical ketoprofen formulations during sunny weather on photoexposed areas may contribute to the reduction of risk of severe photosensitive reactions or cross-sensitizations.

**Conflict of interest**

The authors declare no conflicts of interest concerning this article.

**References**


Correspondence address:
Ewelina Bogumiłłłał Zuba
Heliodor Święcicki University Hospital
Poznan University of Medical Sciences
49 Przybyszewska Str., 60-355 Poznan, Poland
phone: +48 601158212
fax: +48 618691572
email: ewelina.zuba@interia.pl