Resolution of persistent corneal erosion after administration of topical rebamipide

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Abstract: Rebamipide is an antulcer agent used to treat gastric ulcer and gastritis. Biological effects of rebamipide include cytoprotection, wound healing, and anti-inflammatory properties that are known to be universal for a variety of tissues in addition to gastrointestinal mucosa. The therapeutic effects of rebamipide eye drops are due to its ability to increase corneal and conjunctival mucin-like substances and improve corneal and conjunctival injury in vivo. In this paper, we report a case of Sjögren's syndrome with complete disappearance of corneal erosion after administration of rebamipide eye drops. This was observed even though corneal erosion had not improved for 6 months after punctal occlusion surgery. The patient was a 33-year-old female, diagnosed with Sjögren's syndrome by a salivary gland biopsy. The corneal and conjunctival surfaces were filled with dense erosions, which did not improve with topical drugs. Punctal plugs were applied several times; however, the plugs were repeatedly shed. All four puncta of both eyelids were surgically occluded, and both corneal and conjunctival erosion was clearly improved. However, the erosion in the inferior cornea of both eyes had not improved for 6 months after surgery. We used the newly approved topical rebamipide for treatment of this patient. The corneal erosion gradually improved and completely disappeared 4 weeks after administration of the drug. Dry eye sensation disappeared at the same time. Both membrane-associated mucin and secreted mucin in the ocular surface are thought to be essential for maintenance of the tear film. Induction of mucin from ocular surface epithelium could be an effective treatment in cases of dry eye caused by mucin deficiency. Through its various mechanisms, rebamipide improves ocular surface conditions. To our knowledge, this is the first clinical case report using rebamipide ophthalmic solution. This drug may provide a novel approach to treat drying diseases of the eye.

Keywords: Sjögren's syndrome, dry eye, corneal erosion, rebamipide, Mucosta, mucin

Introduction

Dry eye disease is a multifocal disorder of tears and the ocular surface due to tear deficiency, excessive tear evaporation, and instability of tears.¹ Tear-film instability is due to a disturbance in the ocular surface mucin leading to a dysfunction of mucin, resulting in dry eye. Mucins have several subtypes that are highly O-glycosylated glycoproteins with a high molecular weight. They are present on the apical surfaces of wet-surfaced epithelia, including the ocular surface, respiratory tract, esophagus, and gastrointestinal lining.

On the ocular surface, mucins are classified into two types. One is secreted mucin that is supplied from conjunctival goblet cells, and the other is membrane-associated mucin, which appears on the surface of the cornea and conjunctival epithelium.²
The roles of ocular surface mucin are to maintain tear film, lubricate the ocular surface, form the barrier function of the ocular surface, and remove pathogens.3–5

Rebamipide is an antiulcer agent used to treat gastric ulcer and gastritis since 1990 in Japan.6 Rebamipide has been shown to increase gastric endogenous prostaglandin E2 and I2, to promote gastric epithelial mucin,7,8 to behave as an oxygen–free radical scavenger,9,10 and to have other anti-inflammatory action.11–14 Biological effects of rebamipide include cytoprotection, wound healing, and anti-inflammatory properties that are known to be universal for a variety of tissues in addition to gastrointestinal mucosa.14 The clinical effectiveness of rebamipide for treatment of stomatitis,15 pulmonary,16 renal,17 and liver damage,18 colitis,19 and corneal protection20 have been shown by both basic and clinical research.14 This drug also accelerates healing of experimental gastric ulcers in Mongolian gerbils infected with Helicobacter pylori by improving cell kinetics, reducing apoptosis, and reducing inflammation.21

The therapeutic effects of rebamipide are due to its ability to increase corneal and conjunctival mucin-like substances and improve corneal and conjunctival injury in vivo.20 It is known that rebamipide increases the mucin production in cultured conjunctival goblet cells22,23 and in corneal epithelial cells.24 These reports have indicated that rebamipide increases secretion of both membrane-associated and secreted-type mucins. Rebamipide eye drops (Mucosta ophthalmic suspension UD2%; Otsuka Pharmaceutical, Tokyo, Japan) for dry eye are being developed for approved use in the US. The drug has been approved in Japan since January 2012.

In this paper, we report a case of Sjögren’s syndrome with complete disappearance of corneal erosion after administration of rebamipide eye drops. This was observed even though corneal erosion had not improved for 6 months after punctal occlusion surgery.

Case presentation
The patient was a 33-year-old female, diagnosed with Sjögren’s syndrome at another hospital 5 years previously as a result of Schirmer’s I test and positive anti–SS-A antibody (138.5 U/mL), though with negative anti–SS-B antibody. She had no history of rheumatoid arthritis, systemic lupus erythematosus, scleroderma, or primary biliary cirrhosis. The corneal and conjunctival surfaces were filled with dense erosions, which did not improve with topical artificial tears, hyaluronic acid, or antibiotic ointment (Figure 1). Her best corrected visual acuity was 20/20 in the right eye and 20/60 in the left eye at first visit. Tear secretion was evaluated at 0 mm in the right eye and 2 mm in the left by Schirmer’s I test. Tear breakup time in both eyes was 1–2 seconds. Punctal plugs were applied several times to improve this condition; however, the puncta gradually dilated and plugs were repeatedly shed. Because punctal occlusion by plugs was not effective, all four puncta of both eyelids were surgically occluded by using monopolar electrocautery. The lacrimal puncta had completely disappeared, and recanalization of the lacrimal pathway had not occurred 6 months after surgery. The height of the tear meniscus had elevated, and both corneal and conjunctival erosion was clearly improved due to surgery. However, the erosion in the inferior cornea of both eyes had not improved 6 months after the operation in spite of using topical hyaluronic acid, ointment, and artificial tears (Figure 2). Tear breakup time was slightly improved.
prolonged to 3 seconds; however, it still remained outside of normal ranges. We used the newly approved topical rebamipide for treatment of this patient. The corneal erosion gradually improved and completely disappeared 4 weeks after administration of the drug. The tear breakup time in both eyes increased to 8–10 seconds, and was within a normal range (Figure 3). Dry eye sensation disappeared at the same time, and then her best corrected visual acuity improved to 20/13 in both eyes.

Discussion

Argueño et al reported the tear fluid of patients with Sjögren’s syndrome reduced levels of goblet cell–specific mucin and levels of conjunctival mRNA. Both membrane-associated mucin and secreted mucin in the ocular surface are thought to be essential for maintenance of the tear film. Membrane-associated mucin is necessary for the spread of the tear film. Secreted mucin cannot spread over the site in the absence of membrane-associated mucin, such as occurs at an ulcer site. Drying or keratinization of the corneal and conjunctival epithelium are induced in diseases of the ocular surface, and accompanied by alteration of mucin production via mRNA expression, distribution, and character. Goblet cells are lost at the end stage of these diseases when there is total loss of both types of mucins on the surface of the eye. Induction of mucin from ocular surface epithelium could be an effective treatment in cases of dry eye caused by mucin deficiency. There are reports that rebamipide increases secretion of both membrane-associated and secreted mucins.

Previous studies have demonstrated that rebamipide has various inflammation-inhibitory effects, including suppression of inflammatory cytokine production by monocytes, suppression of free-radical scavenging activity, and suppression of CD4 T cells and T1 cytokines. In addition, rebamipide activates genes encoding such angiogenic growth factors as vascular endothelial growth factor and heparin-binding epidermal growth factor in cultured normal gastric mucosal cells, demonstrating a novel mechanism for its ulcer-healing action. Oral administration of rebamipide also affects all B-cell functions, and suppresses systemic secretion of IgM and IgG1.

Through its various mechanisms, rebamipide improves ocular surface conditions. Rebamipide eye drops immediately improved the patient’s persistent corneal erosion after administration. Rebamipide eye drops may have a different mechanism of action when compared to preexisting drugs. Oral administration of rebamipide increases saliva volume for a mouse model of Sjögren’s syndrome. Although tear secretion didn’t increase in the same research, topical rebamipide may play an important role for the ocular surface of Sjögren’s syndrome through mucin production. To our knowledge, this is the first clinical case report using rebamipide ophthalmic solution. This drug may provide a novel approach to treat drying and keratinizing diseases of the eye.

Disclosure

The authors have no financial interest related to this manuscript.

References


