Corneal and conjunctival intraepithelial neoplasia (CCIN) is a premalignant lesion of the corneal epithelium and limbus and the most common tumor of the ocular surface. Generally it occurs in the interpalpebral fissure, usually at the region of the limbus, although it may be found elsewhere. The etiology of this disease is most likely multifactorial, involving such factors as age, ultraviolet light exposure, human papillomavirus, and human immunodeficiency virus. Management of these lesions includes wide local excision followed by supplemental cryotherapy, and topical chemotherapy with antineoplastic drugs. CCIN has been known to have a high recurrence rate after excision alone. Cryotherapy, radiation,
and chemotherapeutics have been used after excision to reduce recurrence rates. Mitomycin C (MMC) and 5-fluorouracil (5-FU) have been successfully employed against CCIN.\(^3\) More recently, interferon alfa-2b (IFN α 2b) has been shown to be effective in treating CCIN.\(^4\) In the present paper, a successfully treated CCIN patient with IFN α 2b eye drop after clinical and histopathological diagnosis was reported.

## CASE REPORT

A seventy-four year old female patient presented to our clinic with a three-month long history of foreign body sensation, redness, irritation, decreased visual acuity and rising tissue growth in her right eye. She had undergone surgery for retinal detachment in her left eye 10 years before. Best corrected visual acuity was 20/80 on the right eye and 20/1200 on the left eye.

Slit lamp examination revealed a 270-degree (from the 11 to 8-o’clock), gray gelatinous, slightly elevated neovascular tumorous lesion involving the conjunctiva, cornea, as well as the limbus on the right eye (Figure 1A). On examination of the left eye, wide scarring of conjunctiva as a complication of previous retinal detachment surgery, and posterior chamber intraocular lens (PC IOL) were noted.

On the basis of clinical findings, an initial diagnosis of CCIN was made in the right eye and a complete excision of lesion was planned. However, due to the high risk ratio of limbal stem cell deficiency (LSCD) after such a wide local excision for complete lesion removal with no option of left conjunctival autograft because of scarring, we changed our plans and an incisional biopsy was performed from the superonasal of the limbus, which confirmed the diagnosis of CCIN histopathologically (Figure 2A). Therefore, we decided to apply medical treatment instead of surgery. Written informed consent was obtained from the patient according to the Tenets of the Declaration of Helsinki after an explanation of all the treatment options. A single therapeutic agent, topical IFN α 2b 1 million IU/mL was prepared, and was instilled four times daily; this regimen continued until complete disappearance had occurred. She was followed up one day, one week, one month, four months, six months, twelve months, and fifteen months after treatment. Topical therapy was generally well tol-

![Figure 1: A. 270-degree (from the 11 to 8-o’clock) gray, gelatinous, slightly elevated neovascular tumorous lesion involving the conjunctiva, cornea, as well as the limbus. B. Regression of CINN was noted at 4th month of treatment. C. Total resolution of CINN was noted at first year of the treatment. CINN: Corneal and conjunctival intraepithelial neoplasia. (See color figure at http://www.turkiyeklinikleri.com/journal/oftalmoloji-dergisi/1300-0365/)](http://www.turkiyeklinikleri.com/journal/oftalmoloji-dergisi/1300-0365/)
erated with the exception of the development of punctate epitheliopathy developed at the 2nd month of the treatment, which was relieved by preservative-free lubricants. No other side-effects except punctate epitheliopathy were detected. Regression of CCIN was noted during the 4th month of treatment (Figure 1B). Therapy was discontinued after regression of the lesion. Total resolution of CCIN was noted at the first year of treatment, although limbal vascularization was present at 4-o’clock (Figure 1C). Incisional biopsy was performed from the superior, inferior, and nasal limbal region of the right eye to confirm total resolution. No tumor cells were found in the biopsy. No recurrence has been observed yet at the 1.5 years’ follow up, while the limbal vascularization at 4-o’clock remained. Impression cytology was performed including the superior, inferior, and nasal limbal region of the right eye to confirm a total cure histopathologically in the final examination. Benign conjunctival and corneal epithelial cells were reported in the impression cytology (Figure 2B). We suppose that the vascularization depends on LSCD in that area.

**DISCUSSION**

Conjunctival and corneal intraepithelial neoplasia is a precancerous lesion of the ocular surface, which is slowly progressive with low malignant potential. Although CCIN is a slow-growing tumor, appropriate treatment is important due to the potential for local invasion or metastasis to regional lymph nodes. Traditionally, treatment of CCIN is a wide local surgical excision combined with cryotherapy, despite the risk for recurrence. The existence of excision margins at the time of surgery is the most important predicting factor in recurrence rate. In addition, determining the edges and deep margins of the mass during the operation is too difficult. Recent studies have shown that the recurrence rate is approximately 50% when pathological evidence of residual tumor cells in the surgical margins exist, and 5% to 33% when clear margins are found. Due to the high recurrence rates, adjunctive medical interventions for CCIN, such as MMC, 5-FU, and IFN α 2b, have been proposed. These therapeutic agents act as an anti-tumor substance. 5-FU is a pyrimidine analog and works as an irreversible inhibitor of thymidylate synthase. MMC is a chemotherapeutic agent which has an antibiotic activity against the tumor cells. IFN α 2b utilizes intracellular signaling pathway and controls the cell cycle, cell differentiation, apoptosis, and immune response. IFN α 2b eye drops are generally well-tolerated and have minimal side effects. On the contrary, MMC eye drops usually cause epithelial toxicity, reactive conjunctivitis, photophobia and severe discomfort. Several reports have also confirmed the clinical efficacy of topical 5-FU in the treatment of CCIN, although its side-effect profile is worse than IFN α 2b. In this case, we decided to use topical IFN α 2b as a single therapy for primary CCIN based on the reported success of topical IFN.
α 2b for CCIN. This decision was also based on the fact that, as shown in the literature, IFN α 2b has a better side-effect profile and less toxicity. To the best of our knowledge, this is the first reported case of CCIN which was treated only with topical IFN α 2b therapy in Turkey. Interferon therapy was approved by the United States Food and Drug Administration (FDA) for the treatment of many cancers such as hairy cell leukemia, malignant melanoma, hepatocellular carcinoma, cervical neoplasia, multiple myeloma, follicular lymphoma, chronic hepatitis B and C, condyloma acuminata and AIDS related kaposi sarcoma. Despite the off-label use of topical IFN α 2b, it has been reported in several publications to be successful in achieving tumor control with a dosage of one million IU/mL four times daily on the ocular surface. For using this off-label drug, approval was obtained from “The General Directorate of Pharmaceuticals and Pharmacy” in Turkey.

Schechter et al. reported that 28 eyes of 26 CCIN patients had been treated with topical IFN α 2b. Twenty-seven patients had complete resolution; one patient had recurrence after therapy stopped. Likewise, Boehm et al. reported that of the seven patients with recurrent CCIN who had been treated with topical IFN alfa-2b, two patients had a recurrence of CCIN, noted at one year and two months, respectively. In these studies, a wide range of duration-such as a two-to-24-month time frame-is required for complete resolution. Although the reason remains unclear, researchers continued on this anti-tumor medical treatment several months after resolution. This decision was probably dependent upon the subjective experiences of specialists in various cases. In our case, the single IFN α 2b therapy was stopped after we had seen that there was no CCIN did not appear in slit lamp examination and after confirmation of total cure with impression cytology in the final examination.

In conclusion, treatment of CCIN with IFN α 2b is effective, valuable and safe with a low recurrence rate. When compared to surgical excision, this treatment is more reliable because of the non-invasive nature. Furthermore, IFN α 2b is less likely to induce devastating complications such as LSCD on the ocular surface. Additionally, it seems to have less ocular surface toxicity when compared to other topical chemotherapeutic agents. Based on our experience, IFN α 2b is an effective approach in the treatment of CCIN.

REFERENCES