Synergistic effect of combined IFN-alpha2b and IFN-gamma treatment for periocular basal cell carcinoma

Tratamiento del carcinoma basocelular periocular con una combinación sinérgica de interferones Alpha-2b y gamma

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Abstract

Purpose: To assess the results of the application of perilesional HerberPAG® on periocular basal cell carcinoma. Methods: An experimental investigation was carried out on a group of 7 patients who received the same treatment regimen with Heber-PAG® 10.5 × 10⁶ IU 3 times a week for 3 consecutive weeks. Demographic variables were evaluated as well as duration, initial lesion size, clinical subtype, histological subtype, clinical response, objective response and adverse events. Results: The average time of lesion duration was 7.0 ± 1.6 months, with a large diameter mean of 17.4 ± 2.3 mm. The clinical nodular-ulcerative subtype was present on 42.9% of the subjects and the solid and superficial histological subtypes were present with an equal proportion of 42.9%. Conclusions: The use of HerberPAG® was an effective and safe alternative to conservative treatment in subjects with periocular basal cell carcinoma when other therapies are not available.

Key words: Basal cell carcinoma. Interferons. HeberPAG®.

Resumen

Objetivo: Evaluar los resultados de la aplicación del HeberPAG® perilesional, en carcinoma basocelular periocular. Método: Se realizó una investigación experimental, en un grupo de siete pacientes a los que se les aplicó un mismo esquema de tratamiento con Heber-PAG® 10.5 × 10⁶ UI perilesional 3 veces a la semana durante 3 semanas consecutivas. Se consideraron variables demográficas, así como tiempo de evolución, tamaño inicial de la lesión, subtipo clínico, subtipo histológico, respuesta clínica, respuesta objetiva (RO) y eventos adversos. Resultados: El tiempo promedio de evolución de la lesión fue de 7.0 ± 1.6 meses, con un diámetro mayor medio de 17.4 ± 2.3 mm. El subtipo clínico nódulo ulcerativo se presentó en el 42.9% de los casos y los subtipos histológicos sólido y superficial se presentaron con igual proporción, en un 42.9%. El diámetro mayor de la lesión mostró un comportamiento decreciente en función del tiempo, y a las 16 semanas se alcanzó una respuesta clínica completa y objetiva en el 85.7% de los casos. Todos los pacientes presentaron eventos adversos, que se resolvieron con medicación oral sin abandonar el tratamiento. La totalidad de ellos ofrecieron un alto grado de satisfacción con el tratamiento. Conclusiones: El empleo del HeberPAG® resultó ser efectivo, por lo que se podría considerar como una alternativa útil y segura como tratamiento conservador en pacientes con carcinoma basocelular periocular cuando otras terapias no son posibles.

Palabras clave: Carcinoma basocelular. Interferones. HeberPAG®.

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Introduction

Basal cell carcinoma (BCC) is a malignant neoplasm derived from non-keratinized cells that originates from the basal layer of the epidermis. It is the most common cancer type in humans. There are 2.8 million BCC cases diagnosed each year in the US and 700,000 new cases in Europe. Skin cancer in the US population exceeds the statistics of other types of cancer; it is estimated that 1 in 5 Americans will develop skin cancer during their lives. It represents between 80 to 90% of all malignant skin conditions. In the last decade, an increase in its incidence and a tendency to appear at younger ages has been detected.

Initially it presents as a slow-growth small tumor, with the appearance of an ulcerated lesion, with telangiectasias and rounded margins; however, the signs may vary according to the clinical subtype. The local destruction it causes can be significant in relevant structures such as the eyelids, and can generate deformities or loss of function of the affected organ.

Selecting the appropriate treatment will depend in each case on the size of the lesion, location, BCC subtype, general condition, age and aesthetic needs of the patient. Surgery continues to be the first therapeutic option. Traditionally, BCC has been completely removed with 3 to 4 mm of surgical margin combined with primary reconstruction, and excellent results have been obtained. Alternatives such as radiotherapy, cryotherapy, laser ablation, chemotherapy and immunotherapy have been described as useful variants in inoperable cases or extensive lesions. Topical treatment with imiquimod may be an alternative to surgery, but long-term results are not as successful as other modalities. Photodynamic therapy is not generally recommended for BCC management due to low success rates in head and neck.

Conservative treatment of BCC is important in the periocular region, because when surgery is performed respecting the oncological margin at this level, it may involve the removal of large areas of the visual apparatus. The reconstruction of the affected area can cause serious aesthetic and functional alterations. In addition, for aesthetic reasons, there is a tendency in facial surgery to eliminate as little tissue as possible and consequently, tumor margins are not completely resected. This significantly increases the risk of recurrence. Immunotherapy is a valuable tool in the conservative management of BCC. Interferon (IFN) intralesional injections were reported as effective for the treatment of BCC in a preliminary study in 1986. Other studies agree that the use of IFN in the treatment of BCC and squamous cell carcinoma (SCC) has shown a wide range of response (60-100%), with a recurrence rate of up to 17%.

Although the literature on this topic is not comprehensive, encouraging results have already been published regarding the administration of the synergistic combination of IFN alpha-2b and interferon gamma (HeberPAG®, Heber Biotec SA, Havana, Cuba). It was registered in 2008 by the Center for the State Control of the Quality of Medicines [CEDMED], the regulatory authority for medicines in Cuba, and is indicated in the treatment of all BCC subtypes by intratumoral and/or perilesional injection in BCC skin lesions, inducing total regression of the tumors. This alternative enhances the pharmacokinetics of the drug by the combination of two active ingredients that can act synergistically. In this way, the same advantages are obtained with a new distribution mechanism that includes an increase and prolongation of the pharmacological activity without additional toxicity, a decrease in the frequency of injections, together with a greater compliance and quality of life of the patient.

The objective of the authors of this study is to evaluate the results of the application of perilesional HeberPAG® in periocular BCC in a series of treated patients.

Methodology

An experimental study (prospective phase IV clinical trial) was conducted in patients with a diagnosis of periocular BCC, attended in the oculoplastic consultation at the Hospital Universitario Arnaldo Milián Castro de Villa Clara, Cuba, in the period from February 2014 to February 2015.

We included 7 patients of both sexes, over 18 years of age, with a diagnosis of any stage and/or clinical subtype of periocular BCC, with a lesion greater than 1 cm and in which surgical treatment was not recommended, as well as lesions with incomplete resection by previous surgical treatment or relapsing. Exclusion criteria were pregnant women, patients who were receiving onco-specific treatments and patients with hypersensitivity to IFN at the time of inclusion.

A positive diagnosis was established by the clinical characteristics of the lesion, histological and dermoscopic results. Incisional biopsy was performed using the 4 mm punch technique, with fixation of the tissue in a
5% formalin solution and proceeding to its inclusion in paraffin to observe the sections under a conventional light microscope. A specialist in Dermatology performed dermoscopy to establish the presence or absence of signs suggestive of tumor. Both tests were performed at the beginning and 16 weeks after treatment. All the patients in the series were treated with the same treatment scheme, using a stabilized pharmaceutical formulation containing a synergistic combination of IFN-alpha2b and IFN-gamma, sodium hydrogen phosphate, dextran 40, sodium chloride, and human albumin (HeberPAG®, Heber Biotec SA, Havana, Cuba) at a dose of 10.5 x 10⁹ IU/1 ml, by perilesional infiltration 3 times a week for 3 consecutive weeks. All the treatments were administered on an outpatient basis and reevaluated in consultation at 4, 8, 12 and 16 weeks after the treatment was initiated, where the size of the lesion, clinical changes and adverse reactions were recorded.

Demographic variables were considered, as well as duration, initial size of the lesion, clinical subtype, histological subtype, clinical response, OR, and adverse events.

The initial size of the lesion was measured before treatment using a millimeter ruler; the histological subtype was classified according to the anatomopathological report. The clinical response was determined according to the characteristics of the lesion 16 weeks after starting the treatment. Considering this criterion, the following categories were obtained:

- Complete response (CR): total disappearance of the lesion.
- Partial response (PR): reduction of at least 30% of the sum of the largest diameters, taking as a reference the sum of the baseline largest diameters.
- Stable response (SR): not enough reduction to qualify as partial response.
- OR was expressed by the sum of CR + PR.

Hematological and biochemical determinations were made (hemoglobin, hematocrit, leukocyte and platelet count, transaminase, creatinine, urea) before and after treatment.

The long-term follow-up continued quarterly during the first year, on a biannual basis during the second and third year, and annually from the fourth to the tenth year.

The information of each patient was registered in a data collection notebook and processed using the statistical package SPSS 20.0. The results were presented in tables and graphs. Statistical techniques were applied according to the types of variables. Bayesian confidence intervals were estimated for the proportions of clinical response, with a 95% confidence. The ethical principles contained in the declaration of Helsinki for human experimentation were fulfilled and written informed consent of each patient was obtained. This study was carried out with the approval of the ethics committee of the institution. The clinical trial is registered in the Cuban Public Registry of Clinical Trials.

**Results**

In the period described, a total of 7 patients who received the treatment were evaluated. They had a mean age of 75 years (Table 1).

The mean time of tumor duration was 7 months and its largest diameter had an average value of 17.4 mm. The clinical subtype of ulcerative nodule predominated in 42.9% of the cases, and the rest of the subtypes presented in the same proportion, accounting 14.3% for each case. From a histological point of view, the superficial and solid subtypes affected 3 patients, respectively, and only 1 presented an infiltrative subtype (Table 2).

In our series, it was observed that in general there was a decreasing, regular and sustained behavior of the large diameter of the lesion as a function of time (Fig. 1). In 85.7% of the cases, CR was obtained; only one patient presented a stable response, representing 14.3%. These values showed the same behavior in the case of the histological and dermoscopic response (Table 3).

All patients evaluated had adverse events related to the administration of the medication. The most frequent was perilesional erythema, which occurred in 100% of the cases, and fever, in 71.4% (Table 4).

**Discussion**

Periocular BCC continues to lead the list of ocular tumors worldwide. Its management depends on factors related to the patient and the characteristics of the tumor. Surgical resection and reconstruction continues to be the main standard treatment; however, in some conditions it becomes an option to evaluate. Reconstruction of the periocular region may cause eyelid retraction, ectropion or cicatricial entropion, ptosis, dry eye, tumor recurrence, trichiasis, infections, graft rejection, unsightly scars, hyper or hypopigmentation. Currently, there are authors who show increasing interest in the use of immunotherapies as a non-surgical option in the treatment of non-melanoma skin cancer. As a starting point, results of the application of synergistic combinations of IFN, which offers invaluable advantages, begin to be registered.
In this study, the group of patients treated were people who were on average in the seventh decade of life. Some articles indicate that oncoproliferative processes are more frequent in advanced ages, although recently it has been described that there is an increasing tendency for these entities to appear more frequently in young women, possibly associated with excessive skin tanning and sun bathing.

Most patients presented skin phototype type II, according to the international Fitzpatrick classification, which coincides with the literature, where white skin is described as a risk factor for the development of BCC, mainly with the cutaneous phenotype I, with freckles, red or blond hair, and light eyes. The history of exposure to solar radiation, particularly ultraviolet radiation, in inverse correlation to the decrease in skin pigmentation, is generally considered the main risk factor for BCC.

In the group studied, there were no significant differences in terms of sex, although there are authors who suggest that its presentation is more frequent in men, with an approximate ratio of 1.5-2:1.

Mean diameter of the diagnosed lesions was 17.4 mm. It is suggested that metastasis due to BCC is extremely rare; however, there is a risk of 0.55% after years of diagnosis without treatment. Metastasis involves regional lymph nodes, bones, liver and lung, mainly when the lesion has a diameter > 2 cm, the resection is incomplete and has perineural and/or perivascular compromise.

In this study a patient was diagnosed with periocular BCC with a diameter of 2.3 cm; however, it had an excellent response to treatment with a surprising clinical and aesthetic evolution.

The treatment with the synergistic combination of IFN achieved an OR in 85.7% of the cases (CR 85.7% + PR 0); in each case, CR coincided with the histological and dermoscopic results. Only one patient presented SR, since his lesion modified but without reaching a...
30% reduction, and he underwent surgery with a wide-margin excision and aesthetic reconstruction. The OR obtained in these cases was similar to that described by García Vega, et al. 19 in a recent study on the application of HeberPAG® in the periocular region using the same dose of this study.

Figures 2, 3 and 4 show examples of CR and figure 5 shows a case of SR. In all of them, clinical-histological correlation of the lesion with hematoxylin and eosin staining is represented.

Only the patient represented in figure 5 had SR to treatment. The authors propose that, despite the excellent results that HeberPAG® has shown, there is the possibility of not achieving a clinical response, mainly when the characteristics of the tumor prevent the correct diffusion of the drug. Anasagasti, et al. 20 have recorded impressive and very stimulating results using a combination of HeberPAG® with chemotherapy and/or radiotherapy, but this was not the objective of our work and it was not due to ethical reasons.

Recently, the use of vismodegib was approved for the treatment of advanced BCC, which acts through a mechanism of inhibition of the Hedgehog pathway. Garcia Vega compared the effectiveness of vismodegib vs. HeberPAG® in the treatment of non-melanoma skin cancer, and found an evident greater clinical effect with HeberPAG®, with a longer response duration.

The possible mechanisms involved in the clinical effects observed with the IFN combination could be the following: intralesional IFN induces apoptosis of BCC cells through the CD95 ligand-receptor interaction, a mechanism that may be reinforced by IFN through the increase of the CD95 receptor. IFN, by stimulating the expression of the IFN-receptor, could contribute to reverse the observed low levels of this membrane receptor in BCC cells. Additionally, it has been shown that in the presence of IFN-, intracellular IFN signaling is stronger. Both IFNs have anti-angiogenic activity and significantly suppress the expression of the CXCR-4 gene, which in turn can decrease the vascularity surrounding non-melanoma skin cancer and affect the migration induced by SDF-1 and also the migration of CXCR-4 positive cells, as has been demonstrated in squamous cell carcinomas of the head and neck for IFN-. All these properties of IFN, among others, probably contribute to increase the antitumor activity of this new formulation of IFN.

The most frequent adverse events with the treatment were perilesional erythema (100%), fever (71.4%), asthenia, anorexia, chills (57.1%) and, less frequently,
nausea, vomiting and muscle aches. There were no changes in hematological or biochemical tests. These results coincide with other studies. All the reported events were mild in intensity and responded well to symptomatic treatment, without having to discontinue the administration of HeberPAG®. This indicates that the synergistic administration of IFN has a safety profile similar to other marketed pharmaceutical presentations of IFN, which, together with the clinical effects of the combination of IFN, suggest that its use is safe and possible in similar therapeutic designs and prolonged treatment schemes, with the purpose of offering these patients an effective and safe therapeutic option.

The adverse effects reported in patients with vismodegib exceeded 30% of the cases, and included muscle spasms, alopecia, taste alterations, weight loss and fatigue, and were serious in 25% of them. However, HeberPAG® is considered a safe drug, with transient adverse reactions, such as catarrhal symptoms, which can be managed with pre-medication without having to discontinue the administration of IFN. In a recent investigation, García Vega, et al. conducted a compassionate study in a small series of patients with high-grade inoperable brain gliomas, and they noted that the combination of IFN showed signs of clinical improvement and quality of life of these patients, with an acceptable safety profile.

Mohs micrographic surgery continues to be the standard option for the cure of all BCCs located on the face, with an approximate recurrence rate of 6.5% at 5 years. However, due to time and cost limitations, it is reserved for cases with high risk of recurrence. Therefore, treatment with HeberPAG® is useful before surgical treatment in patients with coagulation disorders, including the use of anticoagulants. Also in cases with high risk of scarring defects, such as in diabetic patients, older adults and those where surgery or scarring could generate functional alterations. An additional benefit of this technique is the conservation of the integrity of the skin, important in the clinical recognition of recurrences.

The main objective of the use of HeberPAG® in periocular BCCs is the preservation of the eye and the structures that surround it, as well as the conservation of vision, avoiding reconstructive surgery and obtaining a satisfactory cosmetic effect.

Although the impact of the results of HeberPAG® treatment promise to make it a new and powerful therapeutic option for non-melanoma skin cancer, the behavior and long-term follow-up of this series will be published in future articles, and will provide evidence-based information to demonstrate the stability and sustainability of the clinical response obtained in the periocular region. The inclusion of more representative sample sizes in future research will be taken into account.

Conclusions

The results of this study suggest that the use of HeberPAG® could be considered as a useful and safe alternative as a conservative treatment in patients with periocular BCC when other therapies are not possible.
Ethical disclosures

Protection of human and animal subjects. The authors declare that the procedures followed were in accordance with the regulations of the relevant clinical research ethics committee and with those of the Code of Ethics of the World Medical Association (Declaration of Helsinki).

Confidentiality of data. The authors declare that they have followed the protocols of their work center on the publication of patient data.

Right to privacy and informed consent. The authors have obtained the written informed consent of the patients or subjects mentioned in the article. The corresponding author is in possession of this document.

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Conflict of interest

The authors declare no conflicts of interest.

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