Rapidly Growing and Aggressive Cutaneous Squamous Cell Carcinomas in a Patient Treated with Ruxolitinib

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Ruxolitinib is a Janus kinase (JAK)1 and JAK2 inhibitor approved for the treatment of myelofibrosis and for polycythemia patients who are resistant or intolerant to hydroxyurea. We report a 72 year-old man patient with polycythemia vera who developed multiple cutaneous squamous cell carcinomas (cSCCs) with keratoacanthoma-like histological features while on treatment with ruxolitinib. Similar lesions have been reported in an isolated patient who also received ruxolitinib. Our case confirms that ruxolitinib may induce eruptive cSCCs with characteristic clinical and histological features that differentiate them from conventional non-drug induced lesions. Moreover, we performed a mutational panel analysis of the tumors. The lack of specific mutations in these tumors suggests an impairment of immnosurveillance in the origin of the cutaneous lesions. Frequent and thorough dermatological examinations in patients receiving ruxolitinib with a history of photodamage, skin cancer and/or previous hydroxyurea intake is thus recommended. (Ann Dermatol 31(2) 204~208, 2019)

-Keywords-
Carcinoma, squamous cell, Janus kinase 1, Janus kinase 2, Keratoacanthoma, Polycythemia vera

INTRODUCTION

Ruxolitinib is a Janus kinase (JAK)1 and JAK2 inhibitor approved by the Food and Drug Administration and the European Medicines Agency for the treatment of patients with myelofibrosis (either primary or secondary forms) and polycythemia vera (PV) who are resistant or intolerant to hydroxyurea. We report a patient with PV who developed multiple eruptive cutaneous squamous cell carcinomas (cSCCs) while receiving treatment with ruxolitinib.

CASE REPORT

A 72 year-old man with no significant family medical history and a personal history of hypertension that had been treated for years with amlodipine, fosinopril and hydrochlorothiazide was diagnosed with PV in July 2012. Hydroxyurea and acetylsalicylic acid were prescribed one month later and he was immediately referred to our department to rule out skin cancer in the context of a prospective study of cutaneous manifestations associated with PV and associated treatments. The patient had a Fitzpatrick skin type II and a long sun-exposure history. We detected a facial lentigo maligna, two basal cell carcinomas and one cSCC that had been present for several months. All these lesions were surgically removed with no recurrences. The cSCC was located on the left leg and histologically showed poor differentiation (case 1 in Table 1). In January 2013, five months after hydroxyurea had been prescribed, it had to be withdrawn due to hematological resistance. Hydroxyurea and acetylsalicylic acid were prescribed one month later and he was immediately referred to our department to rule out skin cancer in the context of a prospective study of cutaneous manifestations associated with PV and associated treatments. The patient had a Fitzpatrick skin type II and a long sun-exposure history. We detected a facial lentigo maligna, two basal cell carcinomas and one cSCC that had been present for several months. All these lesions were surgically removed with no recurrences. The cSCC was located on the left leg and histologically showed poor differentiation (case 1 in Table 1). In January 2013, five months after hydroxyurea had been prescribed, it had to be withdrawn due to hematological resistance. Ruxolitinib 10 mg twice a day was then initiated. The dose of ruxolitinib was progressively escalated to 15 mg twice a day, dose that was achieved on August 2013. In May 2014, the patient started developing rapidly growing invasive cSCCs. He developed seven primary lesions and...
Table 1. Clinico-pathological and mutational characteristics of the cutaneous squamous cell carcinomas

| Case number | Area     | Diameter (mm) | Excision            | Reconstruction | Onset during Ruxolitinib treatment | Invasive | Primary/recurrence | WA-like histological features | Desmoplasia | Genetic panel | Gene | cDNA change | Aminoacid change |
|-------------|----------|---------------|---------------------|----------------|------------------------------------|----------|-------------------|------------------------------|-------------|--------------|------|-------------|-----------------|
| 1           | Pretibial left | 10           | Elliptical          | Direct closing | No                                 | Yes      | Primary            | Poorly differentiated         | No          | No           | NP              |             |                |
| 2           | Pretibial right | 25          | Geographical        | Graft          | Yes                                | Yes      | Primary            | Moderately differentiated     | Yes         | Yes          | Yes            | TP53        | c.347C>T p.Ser116Phe |
|             |           |               |                     |                |                                    |          |                   |                              |             |              |                 | TP53        | p.Gly266Arg   |
|             |           |               |                     |                |                                    |          |                   |                              |             |              |                 | TP53        | p.Ser269Asn   |
|             |           |               |                     |                |                                    |          |                   |                              |             |              |                 | TP53        | p.Arg282Trp   |
|             |           |               |                     |                |                                    |          |                   |                              |             |              |                 | TP53        | p.Arg290Q6s   |
| 3           | Pectoral left | 12           | Elliptical          | Direct closing | Yes                                | Yes      | Primary            | Well differentiated           | Yes         | Yes          | NP              |             |                |
| 4           | Occipital | 10           | Elliptical          | Direct closing | Yes                                | Yes      | Primary            | Moderately differentiated     | No          | Yes          | NP              |             |                |
| 5           | Pretibial left | 30          | Geographical        | Graft          | Yes                                | Yes      | Primary            | Well differentiated           | Yes         | Yes          | Yes            | TP53        | c.1024C>T p.Arg342X |
| 6           | Parietal left | 20          | Elliptical          | Secondary intention closing | Yes                                | Yes      | Primary            | Moderately differentiated     | Yes         | Yes          | Yes            | TP53        | c.844C>T p.Arg282Trp |
| 7           | Supra-eye brow right | 5 | Elliptical          | Direct closing | Yes                                | Yes      | Primary            | Well differentiated           | Yes         | Yes          | Yes            | TP53        | c.35G>D p.Gly12Asp |
| 8           | Presternal | 8            | Elliptical          | Direct closing | No                                 | Yes      | Primary            | Well differentiated           | Yes         | Yes          | NP              |             |                |
| 9           | Parietal left | 30          | Geographical        | Flap           | No                                 | Yes      | Recurrence         | Well differentiated           | Yes         | No           | Yes            | TP53        | c.323C>T p.Pro278Ser |
| 10          | Supra-eye brow left | 7 | Elliptical          | Flap           | No                                 | Yes      | Primary            | Moderately differentiated     | Yes         | No           | NP              |             |                |
| 11          | Pretibial right | 10          | Elliptical          | Direct closing | No                                 | Yes      | Primary            | Poorly differentiated         | No          | No           | NP              |             |                |

KA: keratoacanthoma, NP: not performed.
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Fig. 1. Rapidly growing crateriform tumors on the pretibial area (A), presternal (B) and parietal (C).

Fig. 2. Proliferating keratin-filled invagination of the epidermis with areas of deep penetrating strands (H&E, ×10).

Fig. 3. Small nests of keratinizing cells surrounded by a fibromyxoid and keloid-like stroma at the deep invasion front (H&E, ×40).

one recurrence in sun-exposed areas that corresponded to cSCCs over an 18-month period of time (Table 1). All the tumors were rapidly growing crateriform lesions that clinically suggested keratoacanthomas (KA) (Fig. 1). Histologically, all these tumors consisted of cSCCs and six out of seven lesions also showed KA-like features (Table 1, Fig. 2). Areas of fibrosis at the tumor invasion fronts were observed in all the lesions (Fig. 3). All the tumors were treated with surgical excision (Table 1) with no recurrence except for lesion number 6, located on the left parietal area, which showed recurrence 2 months after removal. This lesion (case 9) was excised and irradiated with no further relapses. No decrease in the count number of leukocytes or lymphocytes per mm$^3$ was observed during treatment with ruxolitinib. This treatment was discontinued in December 2015. Four months after ruxolitinib being withdrawn, the patient still developed a new KA-like cSCC (case 10). In a further 12 month follow-up, the patient has only developed a poorly differentiated cSCC with no KA-like features on the leg (case 11).

Mutational analysis was performed on the 6 lesions in which good quality DNA could be obtained by next-generation sequencing with a small targeted panel covering the exonic regions of TP53, KRAS, NRAS, HRAS, BRAF, PIK3CA, EGFR and KIT genes (GeneRead; Qiagen, Hilden, Germany) and sequenced on a MiSeq (illumina, San Diego, CA, USA). The mutational analysis could not be performed in 6 additional lesions due to inadequate DNA quality. Frequent mutations in TP53, previously reported in cSCC, were found in 5 cases and a HRAS mutation in one case (Table 1). The recurrent cSCC (case 9) showed the same mutational profile (mutations in TP53) than that found in the primary lesion (case 6). None of these mutations were observed in granulocytes from this patient, which were mainly characterized by the presence of the p. JAK2V617F mutation. The study was approved by the local ethics committee (Hospital del Mar. CEIC 2010/3913/I). We received the patient’s consent form about publishing all photographic materials.

**DISCUSSION**

We here report a patient who developed rapidly growing cSCCs while being treated with ruxolitinib. Although he had developed several skin cancers before starting ruxolitinib, including a cSCC, the striking clinicopathological
resemblance between all the rapidly growing cSCC and their development over a short period of time, suggests the role of this drug as the inducer of the patient’s tumors. The JAKs are a family of non-receptor tyrosine kinases which can be recruited to cytokine and growth factor receptors where they translate signals triggered by ligand binding into intracellular responses\textsuperscript{1}. The best characterized downstream pathways activated by JAK signaling are the signal transducers and activators of transcription (STATs). STATs are a family of transcription factors that drive the expression of genes involved in proliferation, apoptosis, migration as well as in the production of inflammatory proteins. Inhibition of JAK kinase activity can block the constitutive activation of JAK-STAT pathway that characterizes myeloproliferative neoplasms\textsuperscript{2}.

Extrahematologic side effects induced by ruxolitinib are usually mild and include diarrhea, ecchymosis and headache\textsuperscript{3}. However, as predicted by its in vitro immune deregulation, JAK2 inhibitors may also trigger urogenital tract infections as well as opportunistic infections\textsuperscript{4, 6}. Moreover, some skin adverse side effects such as herpes zoster infection, herpes simplex virus reactivation, disseminated molluscum contagiosum and drug eruption have been reported\textsuperscript{7–9}.

The association of ruxolitinib and skin cancer has also been previously documented in the literature. A higher rate of basal cell and cSCCs was reported in the ruxolitinib arm of the randomized study of efficacy and safety in PV with JAK Inhibitor INCB018424 versus best supportive care (RESPONSE) study\textsuperscript{7}. However, the number of patients with a history of nonmelanoma skin cancer or precancerous skin lesions was also higher in the arm of patients receiving ruxolitinib\textsuperscript{7}. A similar case to ours was recently reported by Fabiano et al.\textsuperscript{10}. They treated a 72-year-old patient with myelofibrosis who developed multiple, eruptive, simultaneous cSCCs with KA-like features two months after starting on ruxolitinib. No additional lesions appeared in the 10-month follow-up period after ruxolitinib had been discontinued. The lesions reported by Fabiano et al.\textsuperscript{10} are clinically and histologically very similar to the ones here described and must be distinguished from KAs. Specially challenging is the histological distinction between KA with malignant transformation and KA-like SCC. They are both exoendophytic lesions showing invaginated infundibulum and pale pink cells with glassy appearance as well as a conventional SCC component. However, in KA-like SCC, the KA and SCC components are mixed and no clear boundaries can be defined. On the contrary, a clear-cut border between the KA and the SCC components can be distinguished in KA with malignant transformation\textsuperscript{11}.

The etiology of cSCC is multifactorial. Ultraviolet radiation is the most relevant etiological factor, inducing numerous C to T mutations (UV signature) in genes such as TP53, HRAS and NOTCH1. However, many other factors, such as genetic susceptibility, inflammation and alterations in local and systemic immunity may also play a role for the development of cSCC. In addition, several drugs may also favor the development of cSCC through different mechanisms. In this regard, the development of rapidly growing cSCCs observed in our patient is similar to those observed in patients taking sorafenib and vemurafenib, both having in common the inhibition of RAF isoforms\textsuperscript{12, 13}. It has been proposed that RAF inhibitor-driven activation of the mitogen-activated protein kinase pathway might unmask oncogenic events in keratinocytes harboring preexisting sun-induced RAS mutations. Of note, RAS mutations are found in 20%–60% of vemurafenib-induced skin tumors\textsuperscript{13}. In contrast, we found HRAS mutations in only 7% of the patient’s tumors, a figure that is similar to that reported in common cSCC\textsuperscript{14}. An alternative explanation would be the impairment of immunosurveillance as a result of JAK inhibition by ruxolitinib. JAK inhibition results in the reduction of pro-inflammatory cytokines such as IL-6 and IL-23 as well as in downregulation of the Th17 cell differentiation programme\textsuperscript{15, 16}. These changes lead to a decrease in natural killer, dendritic and CD4+ T cell populations. Interestingly, a similar immunological microenvironment, favouring cSCC development, is observed in organ transplant patients, who show a Th17 cell differentiation and CD4+ cell decrease due to exposure to immunosuppressants\textsuperscript{17, 18}.

In our patient, the previous ingestion of hydroxyurea, although brief, may have played a synergistic effect in the induction of cSCC. Hydroxyurea is an antimetabolite used in the treatment of myeloproliferative disorders and may predispose to the development of actinic keratosis, Bowen’s disease and cSCC\textsuperscript{19}.

In conclusion, we report a patient with PV who developed multiple cSCC while on ruxolitinib suggesting that JAK1/2 inhibition may trigger the development of cSCCs through immunologic mechanisms. We recommend frequent and thorough dermatological examinations in patients who take this drug and specially in those who have a history of photodamage, skin cancer and/or previous hydroxyurea intake.

**CONFLICTS OF INTEREST**

The authors have nothing to disclose.
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