Case Report
Successful Treatment of Refractory Wart with a Topical Activated Vitamin D in a Renal Transplant Recipient

Luciano Moscarelli, Filomena Annunziata, Anduela Mjeshtiri, Nunzia Paudice, Aris Tsalouchos, Maria Zanazzi, and Elisabetta Bertoni

Renal Unit, Careggi University Hospital, Viale Pieraccini 18, 50139 Florence, Italy
Correspondence should be addressed to Luciano Moscarelli, moscarellil@libero.it

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1. Introduction

Warts are a benign proliferation of the skin and mucosa caused by infection with human papillomavirus (HPV). HPV is ubiquitous, and renal transplant recipients (RTRs) may never totally clear HPV infections, which are the most frequently recurring infections. This infection is important because of its link to the development of certain skin cancers, in particular, squamous cell carcinoma. Regular surveillance, sun avoidance, and patient education are important aspects of the management strategy. Warts are usually treated by traditional destructive modalities such as cryotherapy with liquid nitrogen, local injection of bleomycin, electrocoagulation, topical application of glutaraldehyde, and local and systemic interferon-β therapy. These treatment modalities often cause pain and sometimes scarring or pigmentation after treatment. We herein report a case with a right index finger wart, which was successfully treated with a topical activated vitamin D.

2. Case Report

A 41-year-old woman with unknown native kidney disease received a renal transplant from deceased donor in January 2009. She was on treatment with immunosuppressive therapy based on tacrolimus, steroid, and mycophenolate mofetil. She presented 19 months after transplantation a wart on right index finger (Figure 1(a)) which obtained partial clearance after 6-month treatment with cryotherapy and electrococagulation but it regrew rapidly. We attempted treatment with simple local application of activated vitamin D (gauze wet with calcitriol 0.5 μg solution) at least two times a day (during the morning and the next night). The patient was advised to reapply a gauze wet with calcitriol 0.5 solution after each handwashing. Three months later, the wart disappeared without pain or other side effects (Figure 1(b)), and it has not recurred within the 9 months since the disappeared. The medication was well tolerated.
No adverse effects or abnormal serum test results, including elevated serum calcium level, were observed.

3. Discussion

Incidence of warts in RTRs varies from 8% to 55% depending on the patient’s characteristics, the time since transplantation, and immunosuppressive protocols [4]. Skin biopsy and identifying the type of HPV are required to diagnose precisely. Unfortunately, we could not take biopsies from the affected lesion. Therefore, our diagnosis was only based on the clinical appearance. The vitamin D system has multiple physiological and pharmacological effects mediated by action of the vitamin D receptors (VDRs). Recently, VDR activators (VDRAs) have been shown to inhibit cell replication and have immunomodulatory properties. An important observation was reported which suggested that toll-like receptor (TLR) activation of human macrophages upregulated expression of vitamin D receptor and vitamin D-1-hydroxylase genes, leading to induction of the antimicrobial peptide [5]. This suggests an association of TLRs and vitamin-D-mediated innate immunity [5]. Previously the topical application of vitamin D derivatives has become a first-line therapy in the routine treatment of chronic plaque psoriasis as well as for palmoplantar keratosis [6]. A combination of isotretinoin and calcitriol has been reported as the most effective therapy for HPV-associated precancerous and cancerous skin lesions [7]. The effect of vitamin D derivatives was speculated to be derived from its potential to regulate epidermal cell proliferation and differentiation and to modulate cytokine production [8]. Our case report demonstrates for the first time to our knowledge that local application of activated vitamin D is an effective and well tolerated supplementary treatment of recalcitrant wart. A new focus of interest is the levels of activated vitamin D to be reached, particularly in relation to local cellular growth regulation [9]. These levels may provide an explanation for the striking effect of the activated vitamin D and the minimal effect of its the simple application seen in this study. In spite of the proposed mechanisms, any treatment of warts may be confounded by a potent placebo effect. Hence, we realize the need for further placebo-controlled studies before any final conclusions can be reached. However, the lack of regression of the wart before being treated by other modality in the same patient seems to suggest a local rather than a systemic or placebo effect.

References

