

Antimicrobial and Anti-inflammatory activity and efficacy of phytosphingosine: An *In vitro* and *In vivo* Study Addressing Acne Vulgaris

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Abstract

In acne different pathogenetic factors contribute to the inflammation process, defect in keratinisation, increased sebaceous gland activity and increased colonisation of *Propionibacterium acnes*. The results of *in vitro* and *in vivo* studies confirm the previous reports on strong anti-microbial effectiveness of phytosphingosine *in vitro* and *in vivo*. In addition, phytosphingosine shows excellent clinical results in the context of skin care in acne, based on the two properties, i.e. anti-inflammatory and anti-microbial activity. These results demonstrate the potential of phytosphingosine to enhance or complement existing acne therapies acting as an active cosmetic ingredient.

Introduction

The earliest subclinical acne 'lesion' is a microcomedone, hyperproliferation of the follicular epithelium being its characteristic feature. Recently significant pro-inflammatory factors, such as interleukin-1, have been identified around clinically normal pilosebaceous follicles from uninvolved skin in acne patients prior to hyperproliferation of the follicular epithelium[1]. This contributes to the concept that acne vulgaris should be classified as an inflammatory skin disease.

Materials and methods

I *In vitro* studies

Methods

Anti-microbial activity

Using a methodology similar to that previously described(2) the inhibitory effect of phytosphingosine on growth of different micro-organisms was tested.

Release of interleukin-1 α by UVB irradiated human skin on culture

The effect of UVB was investigated using human skin explants in culture as a model. Phytosphingosine (0.2% and 1.0%) and

dexamethasone (10M-6) were applied to human skin explants in culture to test their anti-inflammatory potential. The products were applied one hour before and immediately after irradiation (20 minutes of UVB 2 J/cm²). The interleukin-1 α secretion was measured using an ELISA kit at 24 hours.

Effect on the Artificial Human Epidermis after Irritation with SDS
The efficacy of phytosphingosine on a 3D artificial skin model (SkinEthic™) was investigated after damage with the irritant surfactant sodium laurylsulfate (SDS). After thawing of the artificial human epidermis followed by controlling their viability, a 0.25% SDS solution (dissolved in PBS) has been added to the skin models for 40 minutes to induce chemical stress.

Afterwards the skin slides were washed and a cosmetic O/W formulation (vehicle, formulation containing 0.145% phytosphingosine) was applied.

After 24 hours different parameters as cell death represented by lactate dehydrogenase (LDH), viability according to the XTT assay, inflammatory response judging from interleukin-1 α (IL-1 α) expression.

II *In vivo* studies

Topical *in vivo* study on anti-microbial efficacy

The anti-microbial efficacy of topical phytosphingosine within an emulsion-based format was determined in an *in vivo* test. Two products (phytosphingosine and phytosphingosine-salt) were compared against a control formulation, and a frequently used anti-microbial, triclosan, as a positive control. The formulations were tested on the unwashed hands of 12 subjects based on bacterial counts. The total microbial count was redetermined on the skin at zero time, after 1 hour and after 4 hours.