IN VITRO & EX VIVO TESTING



Atopic Dermatitis

Skin barrier disruption in a Th2-driven inflammation

Atopic Dermatitis is a very common skin disease affecting 2 to 20% of the general population. Dermatitis is characterized by a Th2 inflammatory response associated with epidermal barrier defects. Intense pruritis and colonization by *Staphylococcus aureus* exacerbate the inflammatory process and therefore the lesions.

StratiCELL has developed 2D and 3D skin models displaying atopic dermatitis features, allowing to study the efficacy of dermocosmetic active ingredient and skin care products to restore the skin barrier and reduce inflammation of atopic and sensitive skins. *In vitro* efficacy tests related to staphylococcal infections are also available at StratiCELL.



2D & 3D models

NHEK-Th2: Normal Human Epidermal Keratinocytes stimulated with Th2-type interleukins RHE-AD: Reconstructed Human Epidermis stimulated with Th2-type interleukins



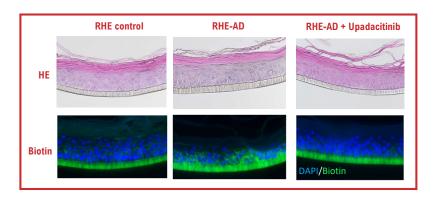
Positive References

- Upadacitinib (JAK/STAT inhibitor)
- GW3965 (LXR agonist)



Testing Methods

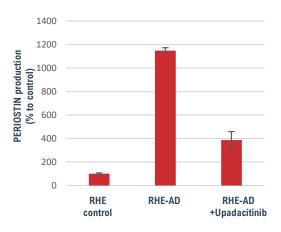
1. Barrier function analysis based on histological Hemalun/Eosin (H/E) images and trans-epidermal in/out Biotin diffusion assay.

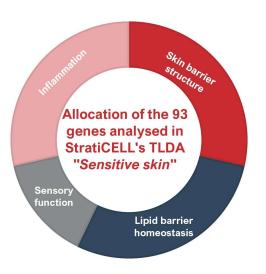


Disrupted epidermal barrier induced in RHE control, treated with Th2-type interleukins alone (RHE-AD) or in the presence of Upadacitinib, after Hemalun/Eosin staining (HE) or biotin diffusion assay (Biotin).

3. Gene expression analysis by RT-qPCR using StratiCELL 's TaqMan Low Density Array (TLDA) studying the expression of 93 genes playing key roles in sensitive skins (inflammation, barrier, lipids, pruritus and sensory function).

2. Quantification of Periostin by ELISA.





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