## Soothing SOS Fluid

## **Application study**

Independent scientific study\* confirming the efficacy of Soothing SOS Fluid

\*In accordance with GCP guidelines (Good Clinical Practice)

| Scientifically<br>confirmed: |  |   |                 |
|------------------------------|--|---|-----------------|
|                              | (5 Min.) Immediate effe  | ect                                     |                 |
| Irritation -24%              | Immediate (within the first 5 minutes) and lasting soothing effect on irritated skin       |   |                 |
| Redness –6%                  | Visible reduction of skin redness  |   |                 |
|                              | <b>r € € €</b> h   |   |                 |
|                              | 14 Long-term effe  | ect                                     |                 |
| <b>TEWL –13</b> %            | Strengthening of the hydro lipid barrier by reducing the TEWL after 14 days of application |   |                 |
| Study design:                | 大大   |   |                 |
|                              | Subjects:<br>2 men + 8 women   | Test area: face, 100%<br>sensitive skin | Age:<br>25 - 62 |
|                              |  |   |                 |
|                              | Dr. med. Christine   |   |                 |
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|                              | derma.cosmetics  |   | Pleas           |

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| Methodology          | Results scientifically confirmed  |  |  |  |
|----------------------|---|--|--|--|
| 5 Min.               | Immediate effect  |  |  |  |
| Irritation:          |   |  |  |  |
| Goal:                | Confirmation of the product's calming potential respectively immediate soothing effect on previously irritated skin   |  |  |  |
| Methodology:         | Stinging test with prior irritation by 10% lactic acid solution   |  |  |  |
| Result :             | Immediate calming effect after one single application (irritation -24%)   |  |  |  |
| Redness:<br>Goal:    | Determination of the visible reduction of skin redness  |  |  |  |
| Methodology:         | Measurement of the reduction of red blood pigment (haemoglobin) in the skin using Antera $3D^{\ensuremath{\mathbb{R}}}$   |  |  |  |
| Result:              | Average reduction     Before     After     Before     After       in haemoglobin level<br>by -6%     Image: Second Se |  |  |  |
| 14                   | Long-term effect  |  |  |  |
| Protective function: |   |  |  |  |
| Goal:                | Confirmation of the strengthening effect on the hydro-lipid barrier by reducing TEWL (transepidermal water loss)  |  |  |  |
| Methodology          | Measurements before product application (D0) and after 14 days of regular application (D 14) by means of Tewameter®™300   |  |  |  |
| Results:             | Average reduction of<br>the TEWL by -13%<br>The lower the TEWL, the stronger the hermion  |  |  |  |
|                      | The lower the TEWL, the stronger<br>the barrier.  |  |  |  |



## Skin tolerance confirmed

Additionally, a dermatological report was prepared as done for every product by Dr. med. Christine Schrammek Kosmetik which confirms the good skin tolerance of the product.

## Dr. med. Christine Schrammek Kosmetik GmbH & Co. KG

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