Keywords: Surfactant-skin interactions, Skin impedance, Skin barrier, Skin cleansing.

Abstract
Surfactant-based cleansing products can cause skin damage and/or irritation due to surfactant-skin interactions, which can compromise skin barrier function. Such interactions need to be minimized. In this work, skin impedance measurements were conducted in vitro on porcine skin using vertical Franz diffusion cells to investigate the impact of surfactants, as well as skin cleansing formulations, on skin barrier integrity and function. This method can guide the development of milder cleansing formulations resulting in less or no skin damage/irritation. Examples of some beneficial formulation additives are illustrated and discussed. The study demonstrates that skin impedance is a useful proxy for skin barrier function and can be utilized as a routine approach to screen surfactant containing formulations for their propensity to compromise the skin barrier.

INTRODUCTION
Stratum corneum (SC), the outermost layer of the skin, plays a critical physiological role in protecting the body from stresses in the external environment [1]. However, cleansing the skin with soap bars and other surfactant-based cleansing products can reduce skin barrier function, as frequent exposure to surfactants leads to various degrees of skin damage and irritation, including dry and itchy skin. It is widely established that the cleansing process also causes damage to the skin by denaturing proteins and/or solubilizing or disrupting the organization of stratum corneum (SC) lipids [2-3]. As the result, skin is often left dry and flaky after frequent cleansing as surfactants interact with the skin and weaken its natural defensive SC barrier function. The undesirable effects of surfactants on skin also include poor skin appearance and increased skin tightness. These effects are greater when the ambient temperature and humidity levels are relatively low. An additional negative effect of cleansing is that surfactant molecules may penetrate into the SC and induce further skin irritation; such effects are greater for individuals with sensitive skin. All these undesirable side effects are interrelated. Many methods, both in vivo and in vitro, have been reported for assessing skin barrier properties. [4-5]. Skin’s electrical impedance is a direct measure of its permeability and can be used to evaluate skin integrity and skin barrier damage and recovery [6]. Skin barrier perturbation due to exposure to different chemicals can be quantitatively expressed by measuring changes in skin impedance. When the skin barrier is perturbed, whether by a physical or chemical stress, its electrical impedance decreases since the transport rate of ions flowing through the skin is higher. It has been shown by several research groups, and for different applications, that skin impedance correlates well with skin permeability [6-8].

To mitigate the negative effects, and to develop improved and milder cleansing technologies, it is clearly desirable to modify the cleansing formulations with materials that will reduce or alleviate surfactant related problems resulting from the cleansing process. Emollients and other formulation components such as glycerin are often added to cleansing formulations to enhance their mildness via a variety of direct and indirect mechanisms. Often the biggest challenge with this approach is preventing components intended for skin deposition from being washed away when the cleanser is rinsed off [9].

In this work, in vitro skin electrical current (impedance) measurements were conducted to quantitatively monitor changes in skin barrier integrity as a function of surfactant chemistry and concentration. The beneficial effect of introducing some ingredients to surfactant solutions and skin cleansing formulation were also investigated and demonstrated using skin impedance measurements.

MATERIALS AND METHODS
Materials
Sodium dodecyl sulfate (SDS), phosphate-buffered saline (PBS) tablets, and glycerin were purchased from Sigma-Aldrich (St. Louis, MO). PBS solution was prepared using PBS
tablet and DI water following the instruction. Sodium lauryl ether sulfate (SLES, Standopal ES-3, ~28.5%) solution were purchased from Cognis (now BASF) and diluted with DI water. Diluted NaOH and H₂SO₄ (both from Sigma-Aldrich) were used to adjust the pH of solutions, if necessary. Maleated castor oil (MCO) and lightly cross-linked PVP (LC-PVP) were obtained from Ashland Specialty Ingredients (formerly International Specialty products). A clear shower gel formulation (Table I) was prepared in the laboratory of Ashland Inc. Commercial cleansing products (body wash and soap bars) were purchased.

**Table I. Original formulation of clear shower gel with MCO.**

<table>
<thead>
<tr>
<th>Ingredients</th>
<th>Composition, wt. %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cocamidopropyl Betaine</td>
<td>20.0</td>
</tr>
<tr>
<td>Maleated Castor Oil</td>
<td>0.55</td>
</tr>
<tr>
<td>Deicodol ED1A</td>
<td>0.1</td>
</tr>
<tr>
<td>Glycerin</td>
<td>6.0</td>
</tr>
<tr>
<td>Sodium Lauroyl Sulfate</td>
<td>20.0</td>
</tr>
<tr>
<td>Propylene Glycol, Dioleoyl Eicos,</td>
<td>1.0</td>
</tr>
<tr>
<td>Methyldodecane, and Propylparaben</td>
<td>1.0</td>
</tr>
<tr>
<td>Sodium Chloride</td>
<td>3.0</td>
</tr>
<tr>
<td>Water</td>
<td>49.35</td>
</tr>
<tr>
<td>Total</td>
<td>100</td>
</tr>
</tbody>
</table>

**Skin impedance measurements**

Detailed skin impedance measurement procedures are available elsewhere (10). In short, full-thickness skin samples from Yucatan pigs were cut into small pieces (3 cm x 3 cm), flash frozen, and stored in a -16°C freezer for up to two months. The skin was allowed to thaw at room temperature for one hour before use. The skin samples were secured in vertical Franz diffusion cells serving as a diffusion barrier between the two compartments, with the SC side facing the donor compartment. Both chambers were initially filled with PBS solution (pH 7.25) avoiding any air gaps or trapped air bubbles. The skin sample was soaked in the PBS solution for one hour before measuring the initial skin impedance. An AC sinusoidal signal, 100 mV RMS (root mean square) at a frequency of 10 Hz was then applied across the skin with a wave generator. Sintered Ag/AgCl reference electrodes were placed in the donor and the receptor compartments (through the sampling port) to measure the electrical potential and current across the skin samples. The electric current was measured using a multimeter. After measuring the initial skin impedance (calculated using Ohm’s law), the solution in the donor compartment was replaced with an experimental solution (e.g., surfactant or formulation solutions) and the skin was soaked for 5 hours. After 5 hours the solution in the donor cell was removed and the donor cell and the SC were rinsed thoroughly with the PBS solution three times. Skin impedance was then measured with PBS solution in the donor cell after one hour equilibration. Each experiment was conducted on 4-12 skin samples. Relative normalized impedance is defined and calculated as the ratio of the skin impedance after and before solution treatment and is a direct measure of skin permeability and hence, barrier function. In these studies the higher the relative normalized impedance (RNI), the less the barrier damage is.

**RESULTS AND DISCUSSION**

**Surfactant systems**

Surfactant-skin interactions with different surfactants (SDS and SLES) was first studied using the skin impedance measurement method. SLES is a frequently used surfactant in personal cleansing products and is generally considered to be a milder and less irritating surfactant than SDS.

Figure 1 compares the relative normalized skin impedance of skin exposed to SDS and SLES at similar concentrations and different pH values. The data show that SLES perturbs barrier function to a much lesser extent than SDS. This correlates well with the milder characteristics of SLES reported in clinical studies and agrees with other in vitro results (3,11). This positive correlation regarding the relative harshness of these surfactants supports the use of skin impedance measurements to study barrier function and its disruption by surfactants and cleansing formulations. In addition, Figure 1 shows the impact of the pH of the surfactant solutions on the skin barrier. To investigate the role that pH plays in surfactant penetration into SC, the pH of an SDS solution was adjusted to pH 4 and the pH of an SLES solution was also adjusted from its normal pH 4 to pH 7. It is clear from Fig. 1 that pH plays an important role in regulating surfactant penetration onto the skin. Increasing the pH of a surfactant solution from pH 4 to 7 increases the barrier damage induced by the surfactant and vice versa. The observation that surfactants with higher pH [more alkaline] are less mild is not surprising and is consistent with previous reports (12). It has been proposed that since the SC pH is acidic (generally around pH 5), which helps protect the skin from microorganisms, the attraction and penetration of basic surfactant solutions is more favorable. However, under basic pH conditions, anionic surfactants bind primarily by hydrophobic functional groups to hydrophobic sites on the skin to minimize the repulsion of negative charges (13). Another factor to consider is SC swelling. Higher pH values in cleansing products are often associated with significantly higher swelling and skin irritancy (3,14). Skin impedance measurements also demonstrate the beneficial effects that glycerin (GLY) and other additives provide in reducing skin barrier damage when added to surfactant formulations. The results shown in Fig. 2 demonstrate that addition of 10% glycerin to SDS surfactant solution reduces surfactant-induced skin barrier perturbation, as quantified by the larger relative normalized impedance. This is in agreement with previous reports (15-16). Figure 2 also shows that addition of 1% maleated castor oil (MCO, INCI name: castoryl maleate oil) separately or combined with glycerin can further reduce the harshness of surfactant systems. MCO is a synthetically modified triglyceride molecule and made by the reaction of castor oil with maleic anhydride (17).
The effect of adding lightly cross-linked polyvinylpyrrolidone (LC-PVP) on surfactant-induced skin damage was also studied using the skin impedance method. Figure 4 shows that a significant reduction (Student t-test \( p < 0.05 \)) of skin barrier damage was achieved by the addition of LC-PVP.

### Skin cleansing formulation systems

Having established that the skin impedance approach was useful in evaluating simple surfactant formulations it was further used to evaluate some commercial skin cleansing formulations.

First it was used to compare the harshness of several soap bar products. Basically there are two types of soap bar products: those with fatty-acid soap-based surfactants, which are anionic type and referred to as regular soap “bars”, and those with non-soap-based surfactants and are referred to “syndet” (synthetic detergent-based bars) (3). Soap-based cleansers are alkaline with a pH around 10, while syndets are mostly neutral or slightly acidic (pH 7 or below). Besides the intrinsic harness of the surfactants, the pH difference between these two types of bars makes a significant difference in the extent of cleanser-induced skin damage. Figure 5 confirms that compared to regular soap bars, syndet soap bars compromise skin barrier integrity significantly less (Student’s t-test \( p < 0.05 \)) even at higher concentration.

Skin impedance measurements were also conducted on a fully formulated clear shower gel formulation (Table I) with and without 0.55% MCO. The cleansing formulation with MCO was specifically designed to effectively deliver MCO from rinse-off cleansing formulations. As shown in Fig. 6, and similar to the SLES system in Fig. 3, the inclusion of MCO in the shower gel cleansing formulation dramatically reduces the surfactant-induced skin damage indicating that the clear shower gel formulation with the addition of MCO is significantly milder than the regular formulation (Student’s t-test \( p < 0.05 \)). This is consistent with other in vitro biophysical studies.

### The mechanism of MCO in reducing surfactant-induced skin damage

The mechanism of MCO in reducing surfactant-induced skin damage is not fully understood. From critical micelle concentration (CMC) and micelle size measurements (data not shown), as well as previous biophysical and clinical results, adding MCO to surfactant solutions does not appear to change the CMC. However, it may retard surfactant molecular penetration by increasing micelle size and provide a significant and substantial occlusive lipid/oil barrier by depositing from cleansing formulations.
and in vivo clinical and studies reported previously (18-19). Besides the aforementioned formulations, the impedance method has also been used in our laboratories to study other practical formulations and commercial products such as baby wash products, hand dish washing products and shave preparation formulations (data not shown here). The results correlate well with clinical results and consumer perception.

**CONCLUDING REMARKS**

The examples in the current work on surfactant solutions and skin cleansing formulations demonstrate that skin impedance measurement can be used to characterize skin barrier integrity as a function of a wide range of physicochemical variables. With relatively straightforward in vitro skin impedance studies, it is possible to quantitatively examine in situ skin barrier perturbation induced by surfactants, and other physico-chemical stresses. The data in the current study are consistent with results obtained from a range of other in vitro and in vivo studies and, as such, validate the use of skin electrical impedance to predict the impact a surfactant-based formulation will have on the skin barrier, including damage and irritation.

Using this approach it is possible to screen many formulations and dramatically reduce the time and effort needed for skin-related formulation improvement, including product development and marketing.
Do you have 100% natural? Let’s discover our magical ingredients!

SONGYI MUSHROOM 松茸蘑菇
Extensive research in our USA laboratories has provided a breakthrough in skincare technology. We have discovered that the enzymes found in the Songyi Mushroom are an effective natural skin-lightening agent. Used regularly, the enzymes penetrate the skin to give a fairer, clearer complexion, with wrinkle reduction. CLINICALLY PROVEN! 我们在美国的实验室经过广泛研究后，在皮肤护理领域取得重大突破。我们已发现松茸蘑菇内所含的酶是有效的天然皮肤增白剂。经常使用，松茸酶精会渗入皮肤，使肤色更加白净，还能有效减少皱纹。以上效果经过临床证明。

KINETIN 激动素
Kinetin is from Endospermum Juice. Young Coconut Juice is physiologically extraordinary active in correcting skin wrinkles due to aging. Kinetin is now widely used in cosmetics as a "magical" anti-wrinkle treatment ingredient. 激动素提取自青玉米和青椰子，能有效平复皮肤老化引起的皱纹。现在激动素作为神奇的抗皱治疗成分被广泛地使用在化妆品中。

PEARL EXTRACT 珍珠萃取液
CAMPO Pearl Extract enhances any skin moisturizer, skin whiteners, skin repair and UV protection cosmetic formulations, with active high performance moisturizer action, high performance skin-whitening and repair action and high performance UV repelling actions. CAMPO珍珠萃取液可提高任何肤质的保湿能力，使皮肤更加白皙，修复肌肤，并提供对紫外线的防护。此配方具有显著的保湿、白皙、修复和防晒性能。

ALPHA-LIPOIC ACID 硫辛酸
Alpha lipoic acid is a powerful anti-oxidant that has diverse effects within the cell, due to its unique molecular structure. It is an anti-oxidant that is both lipid and water soluble, and thus has been designated as the universal anti-oxidant. 硫辛酸是有效的抗氧化剂，因其分子结构独特，有多种效果。脂溶性或水溶性，被认为是一种普通的抗氧化剂。

DMAE 二甲氨基乙醇
DMAE is an antioxidant membrane stabilizer that appears to boost the effects of other antioxidants. As age increases, production of acetylcholine declines, leading to sagging wrinkled skin. Applying DMAE to the skin, results can be seen within minutes, continuing to firm the skin over time, more toned appearance & anti-wrinkle reduction. 二甲氨基乙醇是抗氧化膜稳定剂，可增强其它抗氧化剂的效果，随着年龄增长，人体内制造的乙酰胆碱数量减少，导致皮肤下垂，皱褶。将二甲氨基乙醇用于皮肤护理，几分钟就能见效，皮肤紧绷能持续一段时间，还可以减少色素沉着和皱纹。

Discover our magical skin lightening for Asian skin, lightening of age-spot, sun-spot & freckle-spot management for Caucasian skin

CAMPO RESEARCH USA INC, 5 Penn Plaza, 19th Floor, New York, NY 10001, Tel: 1877-329-8449 Fax: 1877-343-4845
International Marketing & Sales HQ: CAMPO RESEARCH PTE LTD, Level 31, Sixth Battery Road, Singapore 049909, Tel: (+65) 63833203 Fax: (+65) 63833362 CAMPO CHINA, Toll Free Tel: 1860-6500276 Toll Free Fax: 1860-6500277
Website: www.campro-research.com Email: techsales@campro-research.com
Distributor in USA (East & West): BIO-ORGANIC CONCEPTS, Tel: (562) 2365730 Fax: (562) 2365736 Email: info@biocorg.com
Distributor in South America: SARFAM COMERCIAL IMPORTADORA LTDA, Tel: (11) 21140414 Fax: (11) 21140400/21140404
Email: ma@sarfam.com.br
The Botanical Preservative

CAMPO PLANTSERVATIVE™ is a series of read-to-use liquids of Natural Plant obtained green Natural Colorless Clear, Odorless Preservatives contains no parabens. from Honeysuckle Flower Buds (Lonicera Caprifolium & Lonicera Japonica).

MIC and Challenge Tests are at low dosage as 0.125% with "No Preservative" and "Preservative Free" Claim.

植物防腐剂 CAMPO PLANTSERVATIVE™ 是使用天然绿色植物的液体系列，无色透明，无味的防腐剂，不含任何基苯甲酸酯，提取自金银花（忍冬）花。

最少抑菌浓度和筛选试验在低用量0.125%下进行，认定“无防腐剂”或“不含防腐剂”

CAMPO PLANTSERVATIVE WS
CAMPO PLANTSERVATIVE Ws-Neutralized
CAMPO PLANTSERVATIVE Ws-FPFree
CAMPO PLANTSERVATIVE (J)
Lonicera Caprifolium (Honeysuckle) Flower Extract
Lonicera Japonica (Honeysuckle) Flower Extract
忍冬（金银花）花提取液
A Novel plant based preservative
(water-soluble) for cosmetic formulations
独创的植物防腐剂（水溶性）适用于化妆品配方

CAMPO PLANTSERVATIVE WM (Jojoba Oil)
CAMPO PLANTSERVATIVE WM (Bio-IPM)
Lonicera Caprifolium (Honeysuckle) Flower Extract
Lonicera Japonica (Honeysuckle) Flower Extract
忍冬（金银花）花提取液
A Novel plant based preservative
[lipo/oil soluble] for cosmetic formulations
独创的植物防腐剂（脂/油溶性）适用于化妆品配方

CAMPO PLANTSERVATIVE POWDER H2O
Lonicera Caprifolium (Honeysuckle) Flower Extract
Lonicera Japonica (Honeysuckle) Flower Extract
忍冬（金银花）花提取液
A Novel plant based preservative
(powder-soluble) for cosmetic formulations
独创的植物防腐剂（粉溶性）适用于化妆品配方

Visit us at the
Cosmetics Trade Exhibition in 2012
27-29 Feb PCHI Shanghai
15-16 Mar HPCI Mumbai
17-19 Apr InCosmetics Barcelona
15-16 May Suppliers Day New Jersey
23-24 May HPCI Istanbul
24-27 Oct InterCHARM Moscow
6-8 Nov In-Cosmetics Bangkok

SIDDHA MEDICO-COLORS

Colorante-Naturel

天然染色剂

Functional Cosmetics Ingredients from Indian Medicinal Herbs

提取自印度药草的功能型化妆品原料

WATER SOLUBLE LIQUIDS 水溶性液体
Brown Red 棕红 Brown 红
Violet (Blue reddish) 深紫 Green 绿
Red Deep 深红 Ultra Sky Blue 天蓝
White Pearlscendent 珍珠白 Red Light 浅红
Dark Ultra Brilliant Yellow 亮黄
Yellow Light 浅黄

LIPO & OIL SOLUBLE LIQUIDS 脂/油溶性液体
Brown 红 Brown Red 自然Red Yellow 纯黄
Green 绿 Red Light / Scarlet Red 鲜红

OIL SOLUBLE POWDER 油溶性粉末
Brown 红 Yellow 黄 Green 绿 Red 红

Distributor in Europe: BREAGLIO & COMPANY Italy: Tel: (39) 039 493096 Fax: (39) 039 2947377 Email: claudio.porrini@bregaglio.it website: www.bregaglio.it
| Distributor in Eastern Europe: KEMI LINK, Moscow: Tel: (+7) 495 5071591 Fax: (+7) 495 7360032 Email: marinakoval@mail.ru | CHEMICAL CENTRE, Belarus: Tel: +375 17 2226341 Fax: +375 17 2226348 Email: kr-nadya@tut.by | Distributors in (ASIA): ROUNDEX CHEMICAL CO. LTD, China: Tel: (8620) 37885158 Fax: (8620) 37885159 Email: nico@roun dex.com.cn | BRINK CHEMICALS PVT LTD, India: Tel: 91 22 2588289 Fax: 91 22 25882819 Email: info@brinkchemicals.com.in | CHEMWEALTH INC, Philippines: Tel: 63 2 7389428 to 317/398444 Fax: 63 2 6315205 Email: abfirom@chemwealth.com
| P.ZENITH CO LTD, Thailand: Tel: +66 0 29937093-5 Fax: +66 0 29937094-7 Email: pzenith@ksc.th | FOOD & CHEMICALS CO LTD, Vietnam: Tel: 84 8 2976880 Fax: 84 8 2976882 Email: hth@foodchemco.vn | P.K. KOLDSAL, Indonesia: Tel: 62 21 4701971 Fax: 62 21 4701976 Email: ksk@cbn.net.id | SHINWOO I.C.T CO LTD, Korea: Tel: +82 31 4234545 Fax: +82 31 4234777 Email: shinwoo@shinchem.com.kr