Topical administration of mometasone furoate- a combined impedance spectroscopy and in vitro drug diffusion study

Abstract
Mometasone furoate (MF) is a potent steroid for treatment of e.g., eczema and psoriasis. It exerts its function by binding to the glucocorticoid receptors in viable epidermis and dermis. The aim of the current project was to estimate if two clinically equivalent 0.1% MF creams, one w/o (A) and one o/w (B) cream, might impose different systemic load, which is related to adverse side effects. We approached this question by combining analysis of drug diffusion in flow through cells and membrane perturbation detected by impedance spectroscopy using excised porcine skin as membranes. We also analyzed the amount of drug that accumulates in the membranes following topical application of the two creams. The results show that both creams generate about the same amount of MF in skin, while Cream A generates an order of magnitude higher drug flux through skin. Cream A also caused a twofold increase in the skin dielectric constant (ε), which may be attributed to an increased fluidity of the extracellular lipid matrix in the stratum corneum corresponding to a higher skin permeability. No significant change in skin ε was seen with cream B. To conclude, Cream B appears to be the safer alternative as it does not seem to perturb the skin and imposes less systemic burden without sacrificing clinical efficacy.

Keywords: topical glucocorticoid, in vitro drug diffusion, impedance spectroscopy, mometasone furoate, water-in-oil emulsion, oil-in-water emulsion

Introduction
Topical Glucocorticoids (TGs) show vasoconstrictive, anti-inflammatory, immunosuppressive and anti-proliferative effects, and are therefore frequently used to treat psoriasis, atopic dermatitis, asthma and chronic rhinitis. After application to skin, TGs penetrate the stratum corneum and bind to glucocorticoid receptors in viable epidermis and dermis.

Although TGs have been shown to be efficient in treatment of various skin diseases, their use can result in both local and systemic side effects. Some of the local side effects are skin atrophy, rosacea, and corticoid acne, while the major systemic side effects are hypothalamic-pituitary-adrenal-axis suppression along with diabetes mellitus and hypertension. Mometasone furoate (MF) is a potent steroid (according to the European Union classification system) with relatively low risk of systemic side effects. The low risk of side effects is largely due to the high lipophilicity (lgP 4.5), resulting in low solubility in aqueous fluids such as blood. Systemic uptake is further reduced by the ability of MF to degrade in aqueous solvents at pH values above 4.9.

For topical formulations, factors such as their ability to hydrate the skin, drug solubility, and release rate of the drug can also affect the bioavailability of the steroid and are therefore important to consider. In this study we evaluate topical application of two different types of MF creams, both containing the same amount of drug (i.e. Elocon® 0.1% cream, Merck, Sharp & Dohme BV Haarlem, the Netherlands (Cream A) and Ovixan® 1mg/mL cream, Galenica AB, Malmo, Sweden (Cream B). According to SWEDIS 2007, Cream A is water-in-oil (w/o) emulsion with white soft paraffin (34wt%) as the main component of the continuous oil phase. Only 3wt% of the cream consists of water, and hexylene glycol (12wt%) is used as co-solvent for MF. In contrast, Cream B is an oil-in-water (o/w) emulsion with coconut oil (8wt%) dispersed in the continuous aqueous phase. This cream contains approximately 50wt% water and propylene glycol (25wt%) is used as a co-solvent for MF. A previously reported clinical study showed that use of these two different types of formulations gave similar vaso-constrictive properties and equal clinical response.

The aim of the current project was to estimate if these two 0.1% MF creams might differ in systemic load and corresponding risk for adverse effects. We combined impedance spectroscopy and flow through diffusion cells to evaluate barrier perturbation and penetration of MF through excised skin. We also analyzed the amount of drug that accumulates in the membranes following topical application of the two creams. The effects of the creams on the dielectric properties of the skin and the possible correlation between the dielectric properties of the skin and the drug penetration rate were investigated. The combination of impedance and penetration measurements also provided further information on potential pathways for MF through skin.

Materials and methods
Materials
Cream A, Elocon® 0.1wt% cream (Merck, Sharp & Dohme BV Haarlem, the Netherlands), was purchased from a local pharmacy and Cream B, Ovixan® 1mg/mL cream (Galenica AB, Malmo, Sweden), was provided by Galenica AB. Citric acid, sodium citrate dihydrate and sodium chloride were obtained from Sigma Aldrich. Ethanol (99.5%) was purchased from Kemetyl AB (Jordbro, Sweden). Ultra-purified water (ρ=18.2Ωcm; PURELAB UHQ II system; ELGA

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Labwater, High Wycombe, UK) was used for preparation of all aqueous solutions.

Methods

The experimental design comprised membrane characterization with impedance spectroscopy in Franz diffusion cells immediately before and after (within 10min) the drug (MF) penetration study, which was conducted in flow through cells with sampling every second hour. The formulations were removed from the membranes prior to the second impedance measurement by washing 3 times with water and gently wiping with tissue and cotton swabs. Any drug retained in the membranes after terminating the experiment was extracted and MF was assayed with HPLC-UV. To minimize possible damage to the membranes during transport between the impedance and the penetration equipment, a 12.3×2.6mm x-ring, made of Nitrile rubber (Tatingen Teknika AB, Haninge Sweden), was attached to the dorsal side of the membranes using cyanoacrylate glue (Loctite Super Glue Precision, Henkel Norden AB, Stockholm, Sweden).

Membranes

Porcine ears were obtained fresh from a local abattoir and stored at -80°C until use. The ears were then thawed at room temperature and rinsed with cold water. Hair was carefully removed from the inner side of the ear with a trimmer and skin membranes were produced with a dermatome (TCM 3000 BL; Nouvag, Goldach, Switzerland) to a thickness of approximately 500µm. Before initiating the experiment, the skin was allowed to hydrate for one hour at 32°C in the diffusion cells with the dermal side exposed to the receptor media (a mixture of 1.5mL/h and collected as 3-mL aliquots. The collected aliquots of 1.5mL/h and collected as 3-mL aliquots. The collected aliquots were placed in sealed glass vials and stored at −20°C until analysis. Penetration of MF through skin and silicone membranes was determined using flow-through diffusion cells mounted in a heat block according to the system designed by Clowes et al. A water bath with circulation (TechnE TE-10A thermoregulator; Bibby Scientific, Staffordshire, UK) was used to keep the temperature at 32±0.3°C. The test formulations were applied either as semi-finite dose (25mg/cm²), resembling the set-up used in the clinical study by Berg et al. or as infinite dose (260mg/cm²). When the higher dose was used the cells were also occluded with Parafilm to maintain fixed boundary conditions. Furthermore, MF has very low water solubility and degrades at pH values above 4.9. To overcome these restrictions, we decided to use isotonic 20mM citrate buffer (pH 4.5) and ethanol (70:30 v/v) and with the stratum corneum side exposed to the ambient air. Silicone membranes with a nominal thickness of 127mm were obtained from Specialty Manufacturing Inc. (Michigan, USA) and rinsed with distilled water prior to use.

Impedance spectroscopy

The membrane impedance spectroscopy set-up comprised a Franz cell (orifice diameter=9mm, Vsingle=6mL; Permegear Inc, Hellerstown, USA) with four electrodes connected to a potentiostat from Ivium Technologies (Eindhoven, the Netherlands). The electrodes were connected as working electrode (Pt) and sensing electrode (Ag/AgCl) in the donor chamber and as counter electrode (Pt) and reference electrode (Ag/AgCl) in the receptor chamber of the Franz cell. For impedance measurements, an alternating voltage of 20mV amplitude was applied. The system was controlled with Ivium software, which performed a frequency sweep between 1Hz and 6MHz with frequency steps of 5 points per decade (35 points per scan).

Data from the impedance measurements, Z of the membranes were modeled as an equivalent circuit consisting of a constant phase element (CPE) in parallel with the resistance of skin, Rskin:

\[ Z = R_{\text{skin}} + \frac{R_c}{1+(\omega \tau)^\kappa} \]  

Where j is the complex number √(−1), the parameters Q and α define the CPE, and Rskin is the resistance of the electrolyte. The values of α and Q for the different experimental conditions are summarized in Tables S1, S2 (supplementary material). The capacitance, C (Equation 2), was calculated by exploiting the power-law formula proposed by Hirschon et al. and later reexamined by White et al. The approach assumes monotonic, nonlinear (power-law) distribution of the resistivity within the membrane where the power of this distribution is accounted for by the parameter g (Equation 3). As can be seen from Equation 3, g is related to the parameter α, which is a characteristic of CPE used in modeling the frequency dependence of the membrane capacitance.

\[ \frac{1}{\varepsilon} = \frac{L}{\varepsilon_0} g^\alpha \left( \frac{L}{\varepsilon_0} \right)^{1-\alpha} \]  

\[ g = 1 + 2.88 \left( 1 - \alpha \right)^{2.375} \]

Where A is area of skin membrane (0.64cm²), ρδ is the resistivity at position δ perpendicular to the surface (at the dermal side of the skin membrane), L (10µm) is the assumed thickness of the main barrier, the stratum corneum (SC). The dielectric constant is related to the capacitance according to Equation 4.

\[ \varepsilon = \frac{C}{A L} \]  

Where, ε is the permittivity of vacuum (8.8542*10⁻¹²F/m). Hence, the dielectric constant of the skin membrane is given by

\[ \varepsilon = \frac{L}{\varepsilon_0} g^\alpha \left( \frac{L}{\varepsilon_0} \right)^{1-\alpha} \]

The lowest value of the membrane resistivity rd was assumed to be equal to the resistivity of the solution in the receptor chamber of the Franz cell.

Drug penetration

Penetration of MF through skin and silicone membranes was determined using flow-through diffusion cells mounted in a heat block according to the system designed by Clowes et al. A water bath with circulation (TechnE TE-10A thermoregulator; Bibby Scientific, Staffordshire, UK) was used to keep the temperature at 32±0.3°C. The test formulations were applied either as semi-finite dose (25mg/cm²), resembling the set-up used in the clinical study by Berg et al. or as infinite dose (260mg/cm²). When the higher dose was used the cells were also occluded with Parafilm to maintain fixed boundary conditions. Furthermore, MF has very low water solubility and degrades at pH values above 4.9. To overcome these restrictions, we decided to use isotonic 20mM citrate buffer (pH 4.5) mixed with 30% (v/v) ethanol as receptor media. In this way, degradation of MF was avoided and sink conditions were maintained (addition of 30% ethanol increased the solubility of MF to 26µg/mL). The receptor solution was continuously pumped through the cells with a flow rate of 1.5mL/h and collected as 3-mL aliquots. The collected aliquots were placed in sealed glass vials and stored at ~20°C until analysis.

Analysis

MF was extracted from the membranes according to a procedure proposed by Şenyiğit et al. The membranes were gently dried with tissue wipes and the areas which had been exposed in the diffusion cells were placed in sealed glass vials and stored at −20°C until analysis.
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Figure 1 shows penetration of MF through silicone membranes after application of an infinite dose of the two formulations. Interestingly, we found that the penetration rate of MF was much higher from Cream A than from Cream B. The steady state flux of MF after application of the infinite dose was 0.25±0.02µg/cm²/h and 0.05±0.01µg/cm²/h, respectively. Similar results were obtained when semi-finite doses of the two formulations were applied without occlusion during the experiment, with a steady-state flux of 0.24±0.05µg/cm²/h for Cream A and 0.07±0.03µg/cm²/h for Cream B. The absorbed amounts of MF in silicone were proportional to the steady state MF flux from the respective creams (Table 1). The alternative experimental conditions did however not affect the results obtained with any of these two creams. Therefore it was decided to go for infinite dose (260mg/cm²) and occlusion in the subsequent penetration experiments with skin as the membrane.

**Barrier perturbation and mometasonefuroate penetration through skin**

Figure 2 shows penetration of MF through skin following application of Cream A and Cream B. The steady-state flux was calculated over the interval from 6-12h, which gave 0.04±0.02µg/cm²/h for Cream A and 0.02±0.01µg/cm²/h for Cream B. The ratio in steady state flux between the two formulations (A/B) did however differ when silicone and skin data were compared. The difference was significantly higher for skin, 13.9 vs 5.5 (Table 1).

In order to ensure that the observed difference between the penetration rates of MF through skin from Cream A and Cream B (Figure 2) was not due to accidental selection of more (or less) permeable membranes for one of the formulations we measured the electrical impedance of the skin membranes after terminating the penetration experiment. The steady-state flux of MF was plotted vs. the reciprocal resistance (1/R, Figure 3A) and the apparent dielectric constant e, (Figure 3B) of each membrane. After reviewing these data we may conclude that the distribution of skin membranes for studying Cream A and Cream B, respectively, did not result in a systematic difference based on impedance-derived R and e values. Furthermore, there was an apparent correlation between the MF steady-state flux and e(R» 0.6-0.8, Figure 3B), while no such correlation was found between the steady state flux and 1/R (R» 0.2; Figure 3A).

Impedance data were also used to evaluate whether the two creams investigated affect the dielectric properties of skin membranes to different extents. Changes in the dielectric properties of skin upon application of the creams would indicate that excipients from the formulations probably interact with and alter the structure of the skin.
extracuticular lipid matrix in the stratum corneum. The apparent dielectric constant measured after exposure of the membrane to the formulations ($\varepsilon_{\text{after}}$) was therefore compared to that measured before application of the formulation ($\varepsilon_{\text{before}}$). The resulting ratio $\varepsilon_{\text{after}}/\varepsilon_{\text{before}}$ was 1.94±1.38 for Cream A and 0.71±0.31 for Cream B. These values indicate that Cream A has a greater tendency to increase the fluidity of the skin lipids, and thereby the diffusion of lipophilic substances such as MF, than Cream B. It should however be mentioned that the absolute values of dielectric constants for skin membranes, calculated following the formalism described by Equation 2–5, are too high. This is a known problem though attempts are being made to improve interpretation of impedance data, e.g.\textsuperscript{15}

There was a large difference between MF release during the last 2 hours (0.17±0.09 µg/cm\textsuperscript{2}/h) compared to the steady state flux obtained when Cream A remained on the skin ($J_{ss}=0.34±0.15$ µg/cm\textsuperscript{2}/h). A deviation from the linear regime is also clearly visible in Figure 4A, indicating depletion of MF from the skin. For Cream B no significant change in MF release was observed from removing the formulation (0.02±0.018 µg/cm\textsuperscript{2}/h over the last 2 hours) compared to continuous exposure ($J_{ss}=0.02±0.016$ µg/cm\textsuperscript{2}/h). Figure 4B illustrates the relative decrease in individual membranes drug load after removing the creams, calculated by subtracting the release data (Figure 4A) from the amount of MF absorbed in the skin after 6h exposure (Table 2).

Table 2 shows the results from extracting MF from skin after each penetration experiment. After 6h application of Cream A, the skin membranes contained 1.74±1.36 µg/cm\textsuperscript{2} MF, while the skin membranes treated with Cream B contained 0.42±0.35 µg/cm\textsuperscript{2} MF (see Table 2). No significant difference was seen when the membranes were exposed to the creams for 12h instead of 6h (0.04±0.48 and 0.95±0.53 µg/cm\textsuperscript{2}, respectively), as evaluated by the two-sided t-test with a significance level of 0.05. This indicates that equilibrium is reached within 6h exposure, which is in agreement with the observed lag time for skin-penetration (Figure 2). As shown in Figure 2, the amount of MF in skin decreased significantly during the subsequent 6h after removing Cream A (by 57%±10%, to 0.93±0.37 µg/cm\textsuperscript{2}), while only a small decrease was seen for Cream B (by 14%±11%, to 0.62±0.19 µg/cm\textsuperscript{2}, c.f. Figure 4). The ratios in absorbed amounts of MF from the two formulations (A/B) after 12h were 4.3 vs 2.1 when silicone and skin data were compared (Table 1).

Table 1 Clinical and biochemical variables of individuals with overweight-obesity

<table>
<thead>
<tr>
<th>Variable</th>
<th>Cream A</th>
<th>Cream B</th>
<th>Cream A/Cream B</th>
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<tbody>
<tr>
<td>Silicone</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>$J_{ss}$ (µg/cm\textsuperscript{2}/h)</td>
<td>0.25±0.02</td>
<td>0.046±0.014</td>
<td>5.5</td>
</tr>
<tr>
<td>Amount of MF in membrane (µg/cm\textsuperscript{2})</td>
<td>2.70±0.05</td>
<td>0.63±0.12</td>
<td>4.3</td>
</tr>
<tr>
<td>Skin</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>$J_{ss}$ (µg/cm\textsuperscript{2}/h)</td>
<td>0.34±0.15</td>
<td>0.024±0.017</td>
<td>13.1</td>
</tr>
<tr>
<td>Amount of MF in membrane (µg/cm\textsuperscript{2})</td>
<td>2.04±0.48</td>
<td>0.95±0.53</td>
<td>2.1</td>
</tr>
<tr>
<td>$\varepsilon_{\text{after}}/\varepsilon_{\text{before}}$</td>
<td>1.94±1.38</td>
<td>0.71±0.31</td>
<td>n.a.</td>
</tr>
<tr>
<td>$R_{\text{after}}/R_{\text{before}}$</td>
<td>0.16±0.12</td>
<td>1.14±0.93</td>
<td>n.a.</td>
</tr>
<tr>
<td>Amount/$J_{ss}$</td>
<td>6</td>
<td>40</td>
<td>n.a.</td>
</tr>
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Discussion

In the penetration experiments with silicone sheets as membranes, application of 25mg/cm\textsuperscript{2} or 260mg/cm\textsuperscript{2} formulation gave rise to similar MF steady-state flux. It shows that MF penetration though this membrane is not limited by the amount of formulation applied within the range investigated. When 25mg/cm\textsuperscript{2} of Cream A or Cream B was applied, 18% and 4% of the MF, respectively, had penetrated after 12h. Thus, even in the case with 25mg/cm\textsuperscript{2} formulation on silicone membranes the applied amount can be considered to be close to infinite. As the steady-state flux of MF through skin and silicone membranes is in the same order of magnitude, addition of 25mg/cm\textsuperscript{2} and 260mg/cm\textsuperscript{2} would be expected to give similar flux on skin membranes also.

Furthermore, when 260µg/cm\(^2\) was applied, the formulations were occluded while no occlusion was used for 25µg/cm\(^2\). As both set-ups gave similar penetration through silicone membranes, it is likely that any potential evaporation does not affect penetration through the membranes. We chose to apply 260mg/cm\(^2\) of formulation on all skin membranes, in order to increase the reproducibility during application and to ensure excess MF in the donor compartment, which is important to obtain steady-state flux. Furthermore, to avoid changes in the content of the formulation due to evaporation, all skin membranes were occluded during the experiments.

Although Berg et al.\(^6\) reported equal clinical effect from using the two creams, our study has shown that Cream A gives rise to an order of magnitude higher penetration of MF through skin (Figure 2), compared to Cream B. Higher penetration of MF through stratum corneum indicates that increased systemic availability can be expected. As for all topical Glucocorticoids, the target of treatment is viable skin. Systemic availability and uptake in the blood is therefore not desirable, and could possibly lead to severe side effects such as hypothalamic-pituitary-adrenal axis suppression.\(^1,7\)

There was also a slight difference between the two creams regarding the amount of MF that accumulated in skin. After a 6h application, we found that the skin had absorbed 20×10\(^6\) and 5×10\(^6\)MF molecules per cm\(^2\) from Cream A and Cream B, respectively. Smith and Shuster have previously determined that human skin contains 12×10\(^6\) glucocorticoid receptors per cm\(^2\). Thus, after a 6h application of either of the two creams skin contains four orders of magnitude more MF molecules than the number of available receptors. A constraint in comparing our in vitro results to clinical data is however that we ensured that sink conditions applied for the receptor media, resembling a situation where normal blood flow is maintained in vivo. Application of Glucocorticoids does generally result in vasoconstriction with reduced blood flow in the dermis and a reduced ability of molecules to be removed by the systemic circulation. It is therefore possible that higher accumulation of MF in the dermis occurs in vivo where vasoconstriction takes place.

Table 2 Amount of MF accumulated in the skin membranes after application of Cream A and Cream B for 6 or 12 h (n = 4±6, error bars show SD)

<table>
<thead>
<tr>
<th>Time (h) for Removal of Cream</th>
<th>Amount MF in membrane(µg/cm(^2))</th>
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<tbody>
<tr>
<td></td>
<td>Extraction of MF</td>
</tr>
<tr>
<td></td>
<td>6</td>
</tr>
<tr>
<td></td>
<td>12</td>
</tr>
<tr>
<td></td>
<td>6</td>
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<td></td>
<td>12</td>
</tr>
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</table>

Previous studies have shown a direct correlation between the reciprocal of the skin resistance and penetration of hydrophilic molecules through the skin.\(^20\) MF is, however, a very lipophilic compound (logP = 4.5), and therefore it is not surprising that such a correlation between penetration and skin resistance was not observed (Figure 3A). It has been suggested that non-polar lipophilic molecules penetrate the skin by diffusion through the lipid-rich parts of the skin.\(^21\)

The structural features of the lipid layers in the stratum corneum are reflected by the effective dielectric constant, ε.\(^21\) It is therefore more likely to find a correlation between the dielectric constant of the skin and the steady-state flux of MF. This is shown in Figure 3B as linear dependencies between J\(_0\) and ε, with R\(^2\) values of 0.6 and 0.8 for membranes exposed to Cream A and Cream B, respectively. A higher ε value indicates that the extracellular lamellar lipid matrix in the stratum corneum has an increased fraction of lipids in the fluid state, and in many studies it has been shown that molecular diffusion in lipids is facilitated by a transition from the crystalline to the liquid crystalline state.\(^22-24\)

We also compared the dielectric properties of the skin, measured before and after application of the formulations. It is clear that the two formulations affect the permeability of the skin to MF to different extents. Table 1 illustrates how Cream A and Cream B affect the penetration of MF through skin relative to silicone. For penetration through silicone membranes, the ratio between Cream A and Cream B had a value of approximately five. However, when it came to skin, Cream A generated a steady-state flux that was 14 times higher than that for Cream B. This indicates that Cream A promotes the diffusion of MF in skin to a greater extent than Cream B. Because MF is a lipophilic substance, it could be expected to penetrate the skin through the lipid pathway.\(^20\) Increased diffusion would therefore be expected to correlate with increased fluidity of the skin lipids and a corresponding increase in dielectric constant. The resulting ratio ε\(_{\text{cells}}\)/ε\(_{\text{crystalline}}\) had a value of 0.71±0.31 for Cream B and 1.94±1.38 for Cream A. These values confirm that Cream A has a greater tendency to increase the fluidity of the skin lipids, and thereby the diffusion of lipophilic substances, e.g. MF in this case, than Cream B.

Furthermore, a substantial decrease in skin resistance was seen on application of Cream A (R\(_{\text{skin}}\)/R\(_{\text{bead}}\) = 0.16 ± 0.12), which also indicated that the lipid matrix in skin became more fragmented/disturbed. For Cream B, no significant difference in resistance before and after application of the cream could be detected (R\(_{\text{skin}}\)/R\(_{\text{bead}}\) = 1.14±0.93). This, together with the fact that the change in ε on application of Cream B was also not considerable, suggests that the initial structure of the skin membrane is preserved on application of Cream B. In summary, the impedance results indicate that Cream A strongly affects the lipid structures of skin membranes, which probably become more fluid and fragmented. This might be a reason for the much higher penetration of MF from Cream A than from Cream B.

Conclusion

MF from the two creams absorbs to the same extent in excised skin, while Cream A generates an order of magnitude higher steady state drug flux through skin. In corresponding experiments with silicone sheets as membrane the ratios in absorbed amounts of MF and drug flux are similar, which is expected when neither of the creams affects the membrane. Cream A also caused a two-fold increase in the skin dielectric constant (ε) and a decrease in resistance (R). The increase in ε may be attributed to an increased fluidity of the extracellular lipid matrix in the stratum corneum corresponding to a higher skin permeability for lipophilic molecules like MF. No significant change in ε was seen with cream B. The ratio between absorbed amount of MF in skin (after 12h exposure) and drug flux over skin was 6 for Cream A and 4 for Cream B. The relative depletion of MF from skin after removing the formulations was also higher in skin treated with Cream A. To conclude, Cream B appears to be the safer alternative as it does not seem to perturb the skin and imposes less systemic burden without sacrificing clinical efficacy.

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Conflicts of interest

The author declares that there is no conflict of interest.

References
