Topical corticosteroids for childhood eczema: clearing up the confusion

What do prescribers need to know?

- Use the lowest potency topical corticosteroid needed to control the patient’s symptoms
- Be clear when prescribing where each product should, and should not, be used and avoid the term “use sparingly” and encourage appropriate use
- Check that patients and caregivers can identify a flare and are able to respond with appropriate treatment
- For patients with persistent eczema, short treatment “bursts”, e.g. three to five days, with higher potency corticosteroids may be preferable to longer courses of treatment with less potent corticosteroids; topical corticosteroids should be stepped down, e.g. from potent or moderate potency to mild potency, as the patients symptoms resolve
- Include descriptions of potency in the prescription so that it is printed on the medicine label to reduce confusion

Navigating common concerns and confusion

Topical corticosteroids are one of the key medicines used in the management of childhood eczema. However, adherence is typically poor, often due to “corticosteroid phobia”. Common themes contribute to the reluctance of caregivers to use topical corticosteroids (Table 1). Addressing these concerns may improve treatment adherence and patient outcomes.

Provide clear information when prescribing and dispensing topical corticosteroids

Avoid “use sparingly”: encourage appropriate use

Advising patients to “use topical corticosteroids sparingly” creates confusion; patients and caregivers are prescribed a medicine but simultaneously warned against using it. This advice may cause patients to only use corticosteroids when symptoms are severe, resulting in inadequate use and poor symptom control. Caregivers should instead be encouraged to “use corticosteroids appropriately”, which will maximise the benefits of use and minimise adverse effects.
Patients should know:
1. **Which** corticosteroid to apply, i.e. using the right potency and formulation
2. **Where** on the body to apply it
3. **When** to apply it, i.e. when to start treatment and how long to use it for
4. **How** much to apply

### Table 1: Caregiver misconceptions and concerns associated with the use of topical corticosteroids in childhood eczema and evidence-based responses.¹⁻⁴

<table>
<thead>
<tr>
<th>Misconception or concern</th>
<th>What does the evidence say?</th>
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| Topical corticosteroids should only be used for severe symptoms | Topical corticosteroids can and should be used for all severities of eczema, including mild symptoms.¹
Products have a range of potencies to treat patients with differing symptom severity. Treatment should be with mildest topical corticosteroid which is able to resolve the inflammation within a short period of time so that the patient is able to have days without using topical corticosteroids. Different potencies are required for different parts of the body depending on the thickness of the stratum corneum. |
| Regular use of topical corticosteroids causes adverse effects such as skin thinning | Topical corticosteroids are unlikely to cause skin thinning or other long-term harm to children if used appropriately.
Skin thinning is one of the most frequently cited concerns reported by patients and caregivers, however, it is very unlikely to occur if patients and caregivers use topical corticosteroids appropriately.²⁻⁵ This consensus is based on research and clinical experience from Australia and New Zealand, including evaluations of children treated with potent topical corticosteroids.⁶ Skin thinning is more likely to occur in areas with a thinner stratum corneum, such as the face and groin. |
| The percentage of a topical corticosteroid is its strength | The percentage value of different formulations of topical corticosteroids does not indicate their potency, e.g. hydrocortisone 1% is a weaker formulation than hydrocortisone butyrate 0.1%. |
| Corticosteroids are confused with anabolic steroids | Clarify the meaning of the word “steroid”.
Inform patients and caregivers that the label “steroids” is a classification used for a wide group of hormones and medicines with different functions, including corticosteroids and anabolic steroids. |
| Topical corticosteroids should not be applied to broken skin | The consensus of paediatric dermatologists in Australia and New Zealand is that topical corticosteroids can be applied to areas of eczema with broken skin.¹
This concern possibly arose as topical corticosteroid absorption will be greater through broken skin. However, this can prevent patients having topical corticosteroids applied to areas of active eczema particularly when severely inflamed or excoriated. All skin with an active eczema flare will have reduced barrier function, and the best way to address this is through appropriate use of topical corticosteroids. |
| Topical corticosteroids are not “natural” | Corticosteroids mimic the effects of hormones produced by the adrenal glands, despite being “man-made”. |

* For further information on symptom severity and recommended treatment escalation, see: www.bpac.org.nz/BPJ/2015/April/eczema.aspx

Arrange to review the patient within two to four weeks of prescribing topical corticosteroids. This gives an opportunity to assess their response to treatment and reinforce education as well as allowing the patient and caregiver to focus on treating the eczema rather than watching for adverse effects.
Which corticosteroid and formulation to apply

There are a range of subsidised topical corticosteroids available for children with eczema (Table 2).

Consider the consistency of the product required:
- Creams, lotions or gels are useful for large areas of skin
- Lotions, solutions or gels are useful for the scalp or other areas with hair
- Ointments are useful for very dry skin and skin with thick scale

Key points when selecting the potency of topical corticosteroids include:
- Use the lowest potency corticosteroid needed to control symptoms, e.g. hydrocortisone 1% daily or twice daily for mild eczema. However, be prepared to increase potency, particularly for eczema on the trunk and limbs, if a mild topical corticosteroid is not working.
- For patients with persistent eczema, short “bursts” with higher potency corticosteroids, e.g. betamethasone valerate 0.1% twice daily for three to five days, may be preferable to longer courses of treatment with less potent corticosteroids. Betamethasone valerate 0.1% twice daily for three days is as effective as hydrocortisone 1% twice daily for seven days. Patients can be treated with a higher potency corticosteroid initially to gain control of symptoms and then stepped down to a less potent formulation, e.g. hydrocortisone 1%.
  - This results in quicker resolution of symptoms and shorter treatment duration
  - If patients are switched to higher potency corticosteroids ensure they understand that the treatment period is shorter
- If a lower potency of corticosteroid is needed, prescribe a weaker corticosteroid rather than diluting a more potent formulation
  - Diluting topical corticosteroids with emollients does not result in a less potent medicine. Potency is related to the affinity of the particular corticosteroid molecule to the receptor.
- Include corticosteroid potency on medicine labels
  - Patients and caregivers may believe that the percentage of a topical corticosteroid determines its strength, e.g. that hydrocortisone 1% is stronger than hydrocortisone butyrate 0.1%, without realising that different corticosteroids have differing potencies. Labelling a topical corticosteroid as mild, moderate, potent or very potent (Table 2) or similar terms that will be clear to patients, e.g. low, medium, strong, very strong, on medicine labels, reduces confusion and the risk of inappropriate use.

- Very potent topical corticosteroids, i.e. betamethasone dipropionate 0.05% (in propylene glycol base) and clobetasol propionate 0.05%, should not be initiated in children without prior discussion with a dermatologist.

Provide a written plan for the patient and caregiver to take home. This can help to remind them which topical corticosteroid to apply where. For an example, see: [www.starship.org.nz/for-health-professionals/new-zealand-child-and-youth-clinical-networks/child-and-youth-eczema-clinical-network/family-information-and-handouts/]

Where on the body to apply the topical corticosteroid

When prescribing, be clear where each product should be used, e.g. lower potency topical corticosteroids for the face, and specify the treatment duration and any areas of the body where use of the corticosteroid would be inappropriate. For example:

- Methylprednisolone aceponate 0.1% cream
  Potent (strong) corticosteroid – apply once daily to eczema on the limbs and trunk until the flare has cleared. Seek medical attention if symptoms persist after 7 days.
  Mitte 15 g and 2 repeats

For further information on applying topical corticosteroids to different body areas, see: [www.bpac.org.nz/BPJ/2016/February/eczema.aspx]

Caution is required when applying a topical corticosteroid to the face, periorbital or perioral regions and flexural or groin areas.

The face, flexural and groin areas are more susceptible to adverse effects such as striae or skin atrophy and systemic absorption is increased in these areas compared to other sites. For children with eczema on the face, use mild potency or short courses of moderate potency corticosteroids. In flexural or groin areas moderate or potent topical corticosteroids should be used only for short periods, e.g. 7–14 days. Topical calcineurin inhibitors (unsubsidised) are an alternative treatment option for application to these sites. They are more likely to cause a burning sensation and pruritis than topical corticosteroids.

In periorbital regions potent or very potent topical corticosteroids should not be used.

In perioral regions the use of even mild topical formulations has been associated with the development of periorificial dermatitis or “steroid rosacea.” Ongoing use of topical corticosteroids may aggravate these conditions.
Table 2: Topical corticosteroid potency and currently subsidised formulations, sizes and brands.\textsuperscript{16, 17}

<table>
<thead>
<tr>
<th>Potency</th>
<th>Active ingredient</th>
<th>Formulations available</th>
<th>Potency</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Mild</strong></td>
<td>Hydrocortisone 1% (Stat dispensing, three or six month quantities)</td>
<td>Lotions or liquids: Useful for large areas of skin, the scalp or areas with hair.</td>
<td>30 g, 100 g, 500 g&lt;br&gt;DermAssist, Pharmacy Health</td>
</tr>
<tr>
<td><strong>Moderate</strong> (2–25 × as potent as hydrocortisone)</td>
<td>Clobetasone butyrate 0.05%</td>
<td>Cream: Useful for large areas of skin.</td>
<td>30 g&lt;br&gt;Eumovate</td>
</tr>
<tr>
<td></td>
<td>Triamcinolone acetonide 0.02%</td>
<td>Ointment: Useful for skin with thick scale.</td>
<td>100 g&lt;br&gt;Aristocort</td>
</tr>
<tr>
<td><strong>Potent\textsuperscript{*}</strong> (100–150 × as potent as hydrocortisone)</td>
<td>Betamethasone dipropionate 0.05% * †</td>
<td>Lotion 50 mL&lt;br&gt;Betnovate Application 100 mL&lt;br&gt;Beta</td>
<td>15 g, 50 g&lt;br&gt;Diprosone</td>
</tr>
<tr>
<td></td>
<td>Betamethasone valerate 0.1%\†</td>
<td>Lotion 50 mL&lt;br&gt;Betnovate Application 100 mL&lt;br&gt;Beta</td>
<td>50 g&lt;br&gt;Beta</td>
</tr>
<tr>
<td></td>
<td>Diflucortolone valerate 0.1%</td>
<td>Topical emulsion: Useful for skin with thick scale.</td>
<td>100 g&lt;br&gt;Locoid ointment</td>
</tr>
<tr>
<td></td>
<td>Hydrocortisone butyrate 0.1%</td>
<td>Lotion 100 mL&lt;br&gt;Locoid Scalp Topical emulsion&lt;br&gt;100 mL&lt;br&gt;Locoid Cielo</td>
<td>30 g, 100 g&lt;br&gt;Locoid Lipocream</td>
</tr>
<tr>
<td></td>
<td>Methylprednisolone aceponate 0.1%</td>
<td></td>
<td>15 g&lt;br&gt;Advantan</td>
</tr>
<tr>
<td></td>
<td>Mometasone furoate 0.1%</td>
<td>Lotion 30 mL&lt;br&gt;Elocon</td>
<td>15 g, 50 g&lt;br&gt;Elocon</td>
</tr>
<tr>
<td><strong>Very potent\textsuperscript{*}</strong> (up to 600 × as potent as hydrocortisone)</td>
<td>Betamethasone dipropionate 0.05% (in propylene glycol base) * †</td>
<td>Application 30 mL&lt;br&gt;Dermol</td>
<td>30 g&lt;br&gt;Diprosone OV</td>
</tr>
<tr>
<td></td>
<td>Clobetasol propionate 0.05% \† (Stat dispensing, three or six month quantities)</td>
<td></td>
<td>30 g&lt;br&gt;Clobetasol (BNM)\‡ Dermol</td>
</tr>
</tbody>
</table>

| Non-prescription topical corticosteroids | \* Betamethasone dipropionate is available as a potent formulation (Diprosone) or modified formulation with increased potency (Diprosone OV; very potent), both containing 0.05\% active ingredient<br>† Not approved for use in children aged under 12 months\textsuperscript{17}<br>‡ Note that the face, flexural areas, genitals and the groin are more prone to irritation and skin atrophy than other sites; treatment of these areas is usually limited to mild or moderate potency topical corticosteroids.\textsuperscript{7, 9}<br>\‡ To be delisted in March, 2017<br>\(\textcircled{O}\) Fully subsidised; \(\textcircled{\triangleright}\) Partly subsidised; \(\textcircled{\triangledown}\) Unsubsidised |
When to apply topical corticosteroids
Check that patients and caregivers understand when to initiate treatment with topical corticosteroids and when treatment should be stepped down or withdrawn.\textsuperscript{1, 8, 12}

- Emollient use should continue during flares
- Topical corticosteroids should only be applied to areas of active eczema, unless during “weekend treatment” (see below)
- Initially, once daily application of topical corticosteroids is often sufficient. As symptoms improve treatment can be stepped down by either applying a lower potency corticosteroid with the same frequency, or the same potency corticosteroid applied less frequently

How long should topical corticosteroids be applied for?
Topical steroids should generally be effective in clearing inflammation so that long-term treatment is primarily with emollients.

- Flares should typically resolve within seven to 14 days of treatment. If treatment is not effective then consider whether the diagnosis is correct or if treatment should be changed.
- For patients in whom treatment is effective but they have frequent flares, “weekend treatment”, also known as maintenance treatment, should be considered. This consists of applying topical corticosteroids for two days a week during remission.
  - Clinical trials in children with frequent recurrences have shown that once daily application of a potent topical corticosteroid for two days a week to areas of recurrent flaring in the absence of symptoms results in a 55–65% reduction in flares.\textsuperscript{13} Less potent topical corticosteroids have not been studied.

How much topical corticosteroid should be used?
Calculate how much corticosteroid to prescribe and consider offering an indication of timeframe between product repeats if the patient’s history is known. Caregivers can use the fingertip unit (FTU) to estimate the amount of topical corticosteroid to apply (Table 3 and Figure 1). A fingertip unit is the amount of product which covers the tip of the caregiver’s index finger to the distal skin crease from a standard 5 mm tube.\textsuperscript{14} This is a sufficient quantity for an area of skin equal to the palms of two adult hands. One FTU is approximately 0.5 g.\textsuperscript{15}

For example, a child aged five years with eczema mainly affecting both arms will require approximately four FTU of topical corticosteroid per application (Table 3). If this is applied once daily during flares, and flares last approximately seven days in total during a month, this would equate to:

\[4 \times 0.5 \text{ g} \times \text{once daily } \times 7 \text{ days} = \text{approximately } 14 \text{ g}\]

Usage may vary depending on the extent of flares, how quickly they resolve, whether topical corticosteroid use is tapered or stepped down, and whether patients are also using topical corticosteroids during “weekend treatment”.

![Figure 1: Fingertip unit](image)

The adverse effects of topical corticosteroids are mild and reversible
Shortly after application of a topical corticosteroid some patients may experience local irritation or a change in skin colour caused by corticosteroid-induced vasoconstriction.\textsuperscript{5} Hypopigmentation typically clears when the topical corticosteroid is stopped.\textsuperscript{15} Changes in pigmentation usually occur due to the eczema itself or another dermatological condition, e.g. pityriasis alba.\textsuperscript{5, 18}

There is little evidence as to what percentage of a topical corticosteroid dose is absorbed systemically. Studies investigating systemic effects do not measure how much

<table>
<thead>
<tr>
<th>Description</th>
<th>3–6 months old</th>
<th>1–2 years old</th>
<th>3–5 years old</th>
<th>6–10 years old</th>
</tr>
</thead>
<tbody>
<tr>
<td>One entire arm and hand</td>
<td>1</td>
<td>1.5</td>
<td>2</td>
<td>2.5</td>
</tr>
<tr>
<td>One entire leg and foot</td>
<td>1.5</td>
<td>2</td>
<td>3</td>
<td>4.5</td>
</tr>
<tr>
<td>Torso (front)</td>
<td>1</td>
<td>2</td>
<td>3</td>
<td>3.5</td>
</tr>
<tr>
<td>Back and buttocks</td>
<td>1.5</td>
<td>3</td>
<td>3.5</td>
<td>5</td>
</tr>
<tr>
<td>Face and neck</td>
<td>1</td>
<td>1.5</td>
<td>1.5</td>
<td>2</td>
</tr>
</tbody>
</table>

* Note that these values are a guide and will be influenced by the size of the child
of the corticosteroid is in the blood, but instead focus on measuring cortisol as a marker of hypothalamic-pituitary-adrenal (HPA) axis suppression. After a few weeks’ treatment with potent or very potent topical corticosteroids temporary HPA axis suppression does occur. However, this resolves upon cessation of the topical corticosteroid, without the need for dose tapering.\(^5, 19\) HPA axis suppression is more marked when topical corticosteroids are applied under occlusion, e.g. with wet wraps.

**Inappropriate or prolonged use may cause more serious adverse effects**

More serious adverse effects include clinically significant HPA axis suppression, skin atrophy or striae or withdrawal symptoms upon stopping the corticosteroid, such as erythema and aggravation of cutaneous symptoms.\(^5\) These are rarely seen with normal prescribing patterns.

The risk of these adverse effects is increased:\(^5, 20\)

- With a higher potency of corticosteroid
- With application to a greater area of skin or a larger quantity of application
- When corticosteroids are applied under occlusion or to flexural or groin areas, which increases absorption
- If patients are also taking oral or high-dose inhaled corticosteroids
- When potent topical corticosteroids are applied to striae-prone areas, e.g. axillae or groin areas, during growth phases of puberty

If patients request repeat prescriptions earlier than expected consider whether they may be using a topical corticosteroid inappropriately; case reports of adverse effects typically involve patients who have used the product for longer than it was prescribed for.\(^5, 20\)

Ask patients to bring their topical corticosteroids with them to appointments so you can more accurately assess the quantities used

**Patient information on the use of topical corticosteroids is available at:**

- [www.healthnavigator.org.nz/medicines/t-topical-steroids/](http://www.healthnavigator.org.nz/medicines/t-topical-steroids/): Information on topical corticosteroids, how to apply them and potential adverse effects
- [www.nhs.uk/Conditions/Corticosteroid-preparations-(topical)/](http://www.nhs.uk/Conditions/Corticosteroid-preparations-(topical)/): Information on what conditions topical corticosteroids are used to treat, different potencies and formulations of corticosteroids, how to use these medicines and potential adverse effects

**Additional information for general practitioners on the use of topical corticosteroids is available from the Goodfellow Unit:** [www.goodfellowunit.org/podcast/topical-steroids-paul-jarrett](http://www.goodfellowunit.org/podcast/topical-steroids-paul-jarrett)

**Acknowledgement:** Thank you to Dr Diana Purvis, Dermatologist, Auckland DHB for expert review of this article.

**References:**