A PDF of this entire information pack is available to download from our website at www.eczema.org/eczema-at-school.

The checklists on pages 17–22 are available to download as both PDF and Word documents.

You can also download the activities on pages 24–39 in different formats for planning and as classroom resources.
Why do schools need to know about eczema?

Starting school is an exciting yet anxious time for any parent and child. For a family where the child has eczema, this new phase of independence represents an even greater challenge.

In most cases, this will be the first time their son or daughter has been responsible for managing their eczema themselves, without a parent or carer on hand to oversee them. Also, while experienced teachers are likely to have previously taught children with mild eczema, few will have had to support a pupil at the severe end of the spectrum for whom the condition – rather than being a minor irritation – can make life extremely difficult. In the case of newly qualified teachers, it may be the first time they have come across eczema in the classroom environment.

We recognise how daunting this can be for everyone involved, and that’s why we’ve designed this new information pack, which aims to:

1. Provide an overview of the condition, with advice on how teachers can help a child with eczema to integrate into both their class and the school routine.
2. Equip teachers with tools to teach their class about eczema, in order to encourage understanding and compassion amongst a child’s peers.
3. Form the basis of an informed, ongoing dialogue between teacher and parent regarding a child’s condition.

Over the years we have seen first-hand the pivotal role that teachers can play in ensuring a child’s experience of school – whether they have mild, moderate or severe eczema – is as rich, positive and opportunity filled as that of their classmates. We therefore hope that this new resource proves to be both informative and useful and one that you refer to time and time again.

With one in five children now living with the condition, we need the support of the teaching profession more than ever to ensure that every child is able to achieve their full potential. On behalf of those with eczema I would therefore like to say a big thank you for your time today and would welcome any feedback you may have. You will find a short online survey to complete at www.surveymonkey.com/s/eczemaschoolpackfeedback

Margaret Cox
Chief Executive of the National Eczema Society

About us

The National Eczema Society was established in 1975. We support and represent the six million patients with eczema in the UK.

As well as providing information, advice and support through our free* Helpline 0800 089 1122 and popular website www.eczema.org the Society also holds regular training sessions for healthcare professionals and is committed to taking the eczema message out into the local community by attending events across the UK.

*Calls are free from BT landlines. Other providers and mobile services may charge for calls.
What is eczema?
Eczema is a chronic dry skin condition that is accompanied by a ferocious itch.

What causes it?
Eczema is caused by a combination of genetic, immunological and environmental factors that are the subject of ongoing research. Put simply, healthy skin keeps moisture in and external irritants out. When someone has eczema, the skin does not provide the same level of protection as normal skin since the skin barrier is defective. So moisture is lost – causing the skin to become very dry – and external irritants can penetrate the body more easily and cause inflammation.

How common is the condition?
Six million people live with eczema in the UK – 1 in 5 children and 1 in 12 adults.

Why do some people get eczema and not others?
Eczema is one of a group of atopic conditions that often run in families. These include eczema, asthma and hay fever. While a family history of these conditions indicates that a child might be at risk of developing eczema, this doesn’t always happen. Equally, many children develop the condition with no family history at all.

Is it contagious?
Eczema is not contagious and it’s vital that everyone who comes into contact with the child – teaching staff, support staff, fellow pupils – understands that. It can help if the child is taught to respond directly to negative comments with something simple like ‘It’s only eczema, you can’t catch it’.

What are the main symptoms?
Eczema varies in severity from person to person. Some people might have small patches of red, dry, itchy skin. Others might be covered in eczema from head to toe. In severe cases the skin may appear raw, bleed and weep.
Eczema is not a static condition. It can change from one minute to the next and many people are taken aback by how quickly a bout of itching can come on and by the intensity of the urge to scratch.
Eczema is at its most active during ‘flare-ups’, when the skin takes a sudden turn for the worse. For example, the eczema might cover a bigger area than normal or be more inflamed, more itchy, or painful – or all of the above.

Do children grow out of it?
In a large proportion of children eczema clears by the time they reach puberty. However, people who had eczema as a child will often go on to have ‘sensitive’ skin as adults and, in some cases, the eczema will come back in later life. Some children do not grow out of eczema.

Is there a cure?
There isn’t currently a cure for eczema so the aim is to ‘manage’ the condition to minimise its impact on daily life. There is a range of different treatments available. Eczema is treated on a continual basis with medical emollients. Eczema flares are generally treated first with topical steroids and – if the eczema is infected – with antibiotics. There are other options for treating more severe eczema.
Scratching – the agony and the ecstasy!

One of the most frustrating things that people can say to someone with eczema is ‘Don’t scratch’!

We all know that scratching is counter-productive as it tears at delicate skin causing it to become red and sore and even to bleed. However, the short-term pleasure and sense of relief it gives to someone with eczema cannot be underestimated.

To give you an idea of the challenge that not scratching presents, why not try the following?

Next time YOU have an itch, do NOT scratch it. No matter how great the urge, leave it alone. How does that make you feel? Uncomfortable? Irritable? Angry?

Now, imagine that itch was ten times as ‘itchy’ and that it wasn’t located in just one area of your body but in three or five or ten. How would that feel? How would you cope if you couldn’t scratch to relieve your suffering?

This is a great exercise to get the whole class to do over the course of a day or week.

When an itch strikes, tell them to think how the itch and being unable to scratch it makes them feel.

Everyone’s feelings can then be recorded on the board or on a chart to be discussed at the end of the day or week depending on how long you decide to run the experiment for.

Alternatively, why not set it as a weekend task and ask the children for feedback on Monday morning?

One of the reasons it is vital to avoid scratching is that it can lead to a vicious ‘itch–scratch cycle’ (see below).

Another common problem is habitual scratching where the child will scratch even when they aren’t actually itchy. For example, as itching makes them frustrated and the scratching helps to alleviate the itch, they then come to associate scratching with alleviating frustration and will start to use it, unconsciously, whenever they feel this emotion.
Monitor them
Parents will often tell you that their child gets itchy at a certain time of day. As they have identified this pattern they know to look out for it and to be prepared. See if you can identify a similar pattern in the child’s itching at school. Share what you’ve found with the parents so you can work together to identify what’s causing it – tiredness, difficulty understanding a particular subject, a classroom that gets very hot in the afternoon, etc. Equally, if the child is especially itchy one day or one week, inform the parents in case they need a different cream, or need to moisturise more during school hours or are reacting to a substance or situation at home or at school.

Use distraction
As well as openly scratching, a child may generally become fidgety and restless when they are feeling itchy. If the children are working individually, go up to the child and ask how they are getting on and chat with them and do the same with other children on their table. If you’re talking to the class as a whole, give them a role in the talk such as holding up a prop or taking part in a scene. Get them up and moving around by asking them to fetch you something from a cupboard. Busy hands and minds don’t have time to scratch so involve them and some other children in a fast-moving task such as solving problems as a team against the clock.

Eliminate possible triggers
Unfortunately eczema is, by its very nature, ‘itchy’ so we cannot alleviate the symptom completely. However, there are lots of things that make the itching worse which can be addressed, as we describe in our next section.
Triggers are environmental factors that might cause a child’s eczema to worsen. These vary from individual to individual and it can be difficult to identify them as they might not trigger an immediate or noticeable reaction.

Below are some of the most common ‘triggers’ for a child with eczema along with suggestions of ways in which these can be addressed.

**Soap and water**

Normal soap and plain water can further damage the already defective skin barrier in those with eczema, causing the skin to become dry and irritated. Parents should therefore supply a soap substitute for use during school hours. Generally, this will be the child’s emollient, which could be provided as an additional pump dispenser to be kept by the classroom sink, or in a small container to be kept on their person. To ensure that the child always has access to it, discuss with the parents where the most appropriate location(s) would be. It’s also important that the other children understand that the soap substitute is part of the child’s treatment regime – and that only they should use it.

**Sweat**

Whether as the result of break time games in the playground or PE, sweat can be incredibly irritating to sensitive skin. While we would always encourage participation, if the child’s skin is especially bad, the teacher on playground duty could suggest a less active form of entertainment to the group the child is playing in, while a PE teacher might split the class into groups and give each an activity with a different level of intensity, so that the child isn’t singled out.

**Temperature**

Being too hot or too cold or suddenly moving from one temperature to another – e.g. from a warm classroom to a cold school gym – can all trigger a bout of itching.

> Young children might not realise they are overheating so you might need to make suggestions such as ‘If you’re getting a bit warm, take your jumper off’.

> Be aware of the weather. If it’s very hot or very cold outside and the child is already feeling itchy and irritable, it might be an idea for them to play indoors at break time.

> Ensure that a child with eczema doesn’t sit near a draughty door or near a radiator or near a window that the sun pours through.

> On a lovely summer’s day it is tempting to teach part of a lesson outside. If you do decide to venture out, be sure to pick a spot with some shade and ensure the child sits in it – don’t assume they will know to stay out of the sun.

**Wet and messy play**

In the early years these are regular learning activities, which the child with eczema should participate in. However, eczema may be triggered by sand, water, paint, clay and some foods. Sometimes items on the nature table (e.g. plants, leaves and tree bark) may be a problem. The child may benefit from wearing PVC gloves with a cotton liner. Hands should be moisturised before messy or wet activities, then washed with a soap substitute and moisturised with emollients afterwards. Give parents advance notice of such activities so they can bring in any items that the child might need.

**Clothing and uniform**

Clothing can irritate the skin and those with eczema respond best to items that are 100% cotton and that don’t have seams and labels that constantly rub and chafe. Not all schools have uniforms but, if yours does, ask the appropriate people if uniform substitutions for health reasons are acceptable. For example, can the child wear a red jumper like the other children but one that is made from a different material? A child might also be able to wear a thin cotton or silk layer under their school uniform as a barrier to synthetic fabric or wool.
Pollen

If the pollen count is high and the child’s eczema is bad or triggered by pollen, indoor play at break time should, again, be made an option. If the child does go outside, remind them to play in areas of the playground with a manmade surface (i.e. not on the grass).

Dust

With the best will in the world it is impossible to eradicate every speck of dust from a classroom on a daily basis. In fact, it’s not the dust itself that can be irritating to those with eczema but the house-dust mite droppings which it contains. Damp dusting (using a slightly wet cloth) is a particularly effective way of removing dust from a room and it may be worth recommending this to the cleaning team.

Chairs

These can be a problem as inflamed skin can ‘stick’ to them, especially in warmer weather, and plastic can rub and catch. Ask the child’s parents if they can supply an additional cotton sitter or a thin synthetic cushion with a cotton cover that can be placed on the seat to overcome these issues. Again, some children will question this behaviour so the child needs to be taught to say, ‘The chair makes my skin sore. That’s why I have a special cushion to sit on’.

Animals

The other children may love ‘Bob the rabbit’ but he could be responsible for a number of potential irritants, such as his fur, saliva and hay bedding. If moving him to another class is not an option, then he needs to be located at the opposite side of the class from the child and not be allowed to ‘roam’. Cleaning out should take place in another room because of the volume of irritants it will release into the atmosphere, and all the children must be taught to wash their hands immediately after handling Bob in case they then go on to have physical contact with the child with eczema. It is probably best for the child with eczema not to handle ‘Bob’ and definitely not to volunteer to look after him for the weekend.

Food

A few children will have food triggers for their eczema, with cow’s milk and eggs being the most common. As a result, these children will probably bring in a packed lunch that is tailored to their needs. Ask parents if there are any foods their child must avoid as this may also have a bearing on cookery lessons and snacks provided by the school.

Carpets

Carpets are a prime location for house-dust mites and their droppings and can also chafe exposed skin. Ideally, a child with eczema should be provided with a cotton sitter (towels will be too rough) so that their skin does not come into direct contact with the carpet. Also, a child with very dry, sore eczema will have less flexibility in their flexures (the insides of their elbows and behind their knees) so sitting cross-legged on the carpet with their arms folded will be at best painful and at worst impossible. As the rest of the class may view this as preferential treatment, make sure you point out from the first carpet time of term that it is because the carpet will make the child’s skin very sore.

Damp and mould

Older school buildings can have problems with damp and mould, the spores of which generate a reaction in some children with eczema. Unfortunately, cleaning, ventilating and repair work can only do so much.

Fragrance

Fragrance, in the form of liquid, powder or paste, or airborne, can be a trigger for eczema. Scent particles can emanate from cleaning products used in the classroom, an air freshener or your own perfume or aftershave. Ask the cleaning team to use unscented products where possible and to remove air fresheners from both your classroom and adjoining rooms and corridors.

Swimming

Swimming lessons can be a particular challenge as they combine the drying effects of water (it strips oils from the skin) with exposure to chemicals, the most common being chlorine.

All children with eczema will need extra time after the lesson to rinse themselves carefully with clean water – using washing emollient – gently pat themselves dry and then apply leave-on emollient before getting dressed.

If the child’s eczema is moderate to severe, they may also need extra time to apply emollient prior to entering the pool. It is preferable for the child to have some privacy when applying emollient (e.g. a separate cubicle in the pool area) and younger ones may need someone to oversee the process.

Time constraints and the ratio of staff to pupils will determine how flexible you can be in accommodating the child’s needs. It would be valuable to discuss what can and can’t be achieved several weeks before lessons start so that parents can work with you to investigate the best possible plan of action.
Emollients – the first line of defence

The foundation of eczema management is to use emollients (often several times a day) to rehydrate the skin, help prevent further moisture loss and stop external irritants from penetrating the body and causing inflammation.

Emollients are special medical moisturisers that are generally prescribed by a healthcare professional or sometimes bought over the counter at a pharmacy. These are not the same as the cosmetic moisturisers that you would purchase at a beauty counter.
There are many different emollients to choose from and these are available as lotions, creams and ointments.

There is no such thing as one emollient for all children with eczema – choosing a suitable emollient is normally a process of trial and error since what is soothing to one person’s skin may irritate another’s.

The National Eczema Society recommends that emollient is applied gently, by stroking downwards in the direction of hair growth. Even for mild eczema, emollients often need to be applied three times a day. Daily ongoing emollient therapy is extremely important for all severities of eczema, to repair the skin barrier and try to prevent the itch–scratch cycles, which will inevitably lead to eczema flares. In more severe cases of eczema, emollients will need to be applied far more frequently as the skin dries out much faster.

Most children with eczema will therefore need to apply emollient at least once during the school day. Because of the ‘no touch’ policy employed by schools, we recommend that parents begin teaching their child to apply their own emollient well in advance of starting the first term. However, it is likely that most children will still need reminding about what time to apply their creams and some degree of supervision during the process.

What can I do to help?

Ask parents what works best with their child. For example, some children might need to be talked through the process: ‘First your right arm, that’s it, now your left arm…’ while others might be extremely self-conscious and prefer if you to talk about anything but them applying their cream!

It might also be helpful to have the parent oversee the first application on school premises so you can observe the approach they take and the child can understand that, during school hours, you will be taking on the role.

Ask parents to measure out how much cream the child should use so you have a visual guide. This will vary from child to child and will also depend on the product being used.

The process of the child dipping their hand in and out of a tub of cream as they apply it will lead to the remaining emollient becoming contaminated.

Ask parents if it would be possible for the child to have a pump dispenser instead. Alternatively, they could provide a clean spoon or spatula that the child can use to spoon out the cream from the tub, rather than using their hands.

Discuss with the parents when would be the best time to apply the cream. Some children may be able to manage with just a lunchtime application; others might need to apply cream at morning break, lunchtime and afternoon break.

If this is the case, you may need to involve some of your colleagues so that each of you oversees one application slot.

It is important to leave a gap between applying emollients and putting on sun screen – ideally 30 minutes – so that emollient is absorbed properly and does not dilute the sun screen; also, to prevent the child from ‘frying’ in the sun.

Discuss with the parents how long it takes their child to apply the cream. If this proves to be longer than the duration of a break time, you might need to explore alternative solutions with them.

Decide where the child will go to apply the cream. A private location should be provided, but toilets are not appropriate due to the risk of infection. Most children will also view this as a very private process that they want to undertake well away from their classmates.

Decide where the emollient is going to be stored in-between applications. This needs to be somewhere cool and dry, away from direct heat and light. Contamination of skin-care products is an issue as eczema can easily become infected, so the tub or dispenser needs to be kept in a place where it won’t be inadvertently opened or tampered with.
Ensure that all members of staff who might be called upon to oversee the moisturising process know where the emollient is stored, how much the child needs to apply and the best strategy to take in terms of guiding them through it.

Ask the parents for a smaller container of emollient for ‘top-ups’ following wet play, arts and crafts, outdoor activities and so on, when the skin on their hands will have been exposed to various chemicals and irritants. Make sure the other children know not to touch it. Lower down the school, suggest that a responsible adult looks after the container.

Another important reason for having this small tub accessible to the child is that they will need to use it as a ‘soap substitute’ to wash their hands (soap must be avoided, as it irritates eczema).

Agree on a simple line that the child can use with their peers if asked about the process. For example, ‘I have to put cream on my skin or it gets very dry and sore’.

If the child is reluctant to apply their cream on school premises, but is happy to do so at home, suggest to the parents that they use a reward chart. The chart is on display at home and the stickers are handed over by you at the end of each day – one for every successful application at school. In this way there will be a clear link between school (the stickers) and home (the chart) to encourage consistency of behaviour. The parents then offer a small reward after a certain number of stickers have been achieved.

Finally, ask parents to keep you updated on any changes (for example, if they decide to use a new emollient, if the amount to be applied changes, or if the frequency of application increases or decreases).

Maximising the school nursing team contribution to the public health of school-aged children.

Department of Health. DoH (2014)

From September 2014, new Department of Health (DoH) legislation means that supporting children with medical needs in schools is mandatory. This document replaces the current ‘Managing Medicine in Schools’ DoH/Department for Education document. There are several layers of care or intervention for children with medical needs; eczema fits into the ‘Universal Plus Intervention’.

Universal Plus Intervention

The ‘Maximising the school health nursing team contribution to the public health of school-aged children’ document clarifies the vision and model for school nursing and clearly maps out the different levels of intervention.

The school nursing service is for every child in the UK aged 5–19 years. Every education setting will have a named school nurse and every GP practice will have a named school nurse linked to the practice.

As a child approaches school entry, transition to the local school nursing service will happen in line with local policy. The school health check will identify the healthcare needs of a child entering school.

A child starting school with eczema would fit into the Universal Plus Intervention, which defines that a swift response is required from the school nursing service when a need for expert help is identified through a school health check or accessible services (the GP, health visitor or specialist dermatology clinic). This means the school nurse with the school will set up a health needs assessment to address the individual needs of the child with eczema and ensure that appropriate support is available to meet the health needs identified.

The Universal Plus Intervention will therefore provide teachers with an additional support system when managing the specific needs of a child with eczema in their class. Combined with the information, advice and resources provided by the National Eczema Society in this pack, this should ensure that every child with eczema is able to fully participate in the academic experience and achieve their true potential.

The DoH document can be found at: www.gov.uk/government/publications/school-nursing-public-health-services
The building blocks of eczema treatment

**Emollients**

Emollients (medical moisturisers) are the foundation of all eczema management – to prevent moisture loss, rehydrate the skin and stop external irritants from penetrating the body. Emollients are required on a daily basis for all children with eczema. For more information, see pages 8–10.

**Topical treatments for eczema flares**

For children with mild to severe eczema there is a range of other treatments that will be prescribed for use alongside their emollients to help treat eczema flare-ups.

As eczema is such an individual condition, sometimes identifying the most effective combination of treatments can take time and is a complex process, especially if a child has severe eczema. Most children with mild and moderate eczema will have a standard skin-care routine and treatments prescribed by their healthcare professionals for flare-ups.

While you will not be expected to oversee the administration of the following treatments, it is useful to understand how each works and any implications for the child during the school day.

**Topical steroids**

Topical steroids are usually prescribed if emollients alone are not enough to manage a child’s eczema. They are used for short periods during flare-ups to reduce inflammation and itchiness and help the skin to heal.

Topical steroids are available as ointments, creams and lotions and come in four different strengths – mild, moderately potent, potent and very potent. Parents will be instructed to apply them once or twice a day, depending on the steroid prescribed and the severity of the eczema. Unlike emollients, which are used daily, topical steroids will normally only be used during a flare-up.

While a child can be taught to apply their own emollients, topical steroids must be applied by an adult in very precise amounts and only to active and recovering areas of eczema. Children should therefore have their topical steroids applied at home. As they are prescribed for use only once or twice a day, application will not be required during school hours.

**Topical immunomodulators**

Topical immunomodulators help to calm down the skin’s overactive immune system, reducing inflammation and redness. One of their advantages is that they can be used on very delicate skin, such as the face and eyelids, and for prolonged periods.

Topical immunomodulators are applied thinly to active eczema once or twice a day. Again, children must not apply topical immunomodulators themselves, but since application is only once or twice daily this can happen outside of school hours.

Patients using topical immunomodulators are advised to limit their exposure to the sun and to use sunscreen to protect their skin. If a child is using this form of treatment, you will need to agree with the parents how much sun exposure they can have over the course of the day – taking into account break times, lunchtime play, outdoor PE lessons and any other outdoor lessons – and the options for limiting this; for example, a child may need to stay indoors over lunchtime when the sun is at its most intense.

You will also need to ask the parents how often their child should apply sunscreen, if they know how to apply it, how much supervision they need and how this will fit in with their emollient routine during school hours. Remember, it is important to leave a gap between applying emollient and sun screen (see page 9).
Paste bandages and wet wraps

Bandaging the skin not only helps it to absorb emollient better but also prevents further damage being caused by scratching. Paste bandages are impregnated with a paste containing zinc oxide or zinc oxide and ichthammol, which are soothing and cooling ingredients that help to relieve irritation. Wet wrapping involves a layer of emollient on the skin, a wet inner bandage and a dry outer bandage. The inner layer will require spraying with water during the school day to prevent it drying out. While most children will have bandages and wet wraps applied at night, sometimes a child may wear a wrap or bandage on a small area, such as a wrist or ankle, to school. Extensive bandaging and full body wrapping look unusual and can therefore result in a negative reaction from peers, so it is best if the child only wears bandages and wraps to school if absolutely necessary.

It is important that the child is taught to say, ‘The bandages help my skin to get better’, but further intervention from you may be required to reassure the class that the bandages are nothing for them to be alarmed about.

Therapeutic clothing

A range of silk garments are now available on prescription. Because the fibres are soft, natural and ‘breathable’, they help to regulate body temperature and humidity, absorb sweat and reduce irritation. As they are worn under a child’s school uniform (providing a useful barrier against synthetic materials and wool), they should not be visible to classmates.

Tablets and hospital-based treatments

A small number of children have severe atopic eczema (and often other atopic and allergic conditions). These children are often under the care of a dermatologist and may be prescribed additional treatments to ones that are applied to the skin.

Azathioprine

Azathioprine is a potent immunosuppressant drug that reduces the body’s normal immune response. In eczema it helps to reduce the inflammation associated with the condition, allowing the skin to heal, and will be taken for several months at a time. Only a consultant dermatologist can prescribe and oversee its use. All patients taking azathioprine have regular blood tests to monitor for bone marrow suppression, which can lead to severe anaemia and an increased risk of infection. Liver function tests are also carried out regularly.

Given the nature of the treatment, you can expect to have several weeks’ notice of a child starting a course of azathioprine as they will need to undergo tests beforehand to assess their suitability. During school hours you should monitor for nausea, diarrhoea and loss of appetite as these are all possible side effects. Azathioprine can cause photosensitivity, so extra diligence with sun protection is required.
Ciclosporin

Like azathioprine, ciclosporin is a potent immunosuppressant drug that reduces the inflammation associated with eczema. In children it is used as a short-term treatment under the supervision of a consultant dermatologist.

The main side effects of ciclosporin are high blood pressure and reduced kidney efficiency. Blood pressure and kidney function will therefore be checked before treatment and monitored closely throughout treatment using blood and urine tests.

As the risk of bacterial, fungal and viral infection is greater when a child is on this medication, it may be advisable to discreetly seat the child away from other children who are exhibiting signs of becoming unwell, or who are unwell or who are recovering from illness.

Oral steroids

Oral steroids are anti-inflammatory drugs that help to bring about a rapid reduction in the redness, weeping and irritation associated with eczema. A typical course will last up to two weeks and medication is taken once a day. **It is important that the school knows if a child is having a course of oral steroids, and the child should carry a steroid card at all times.**

Oral steroids can suppress the immune system, which means that the risk of bacterial, fungal and viral infection is increased. Therefore, where possible, you should try to minimise direct contact between the child and pupils who are exhibiting any signs of illness.

**In particular, the child’s parents will need to know as soon as possible if their child has potentially been exposed to the chickenpox or shingles virus or measles.**

Phototherapy

Some people with eczema find that their condition improves with phototherapy, a type of UV light treatment that is available at a specialist clinic or hospital. Treatment is usually given two or three times a week for up to ten weeks and helps to reduce inflammation. Generally, treatments are arranged outside school hours but sometimes the child may need time off school to attend a session.

Some patients experience redness and itching after this treatment and, as a result, the emollient used during school hours or the frequency of application, or both, may need revisiting during the treatment process.

After treatment the child should avoid further exposure to UV light. You will need to discuss with the parents if this means they must stay inside the building during school hours or whether they can have a limited amount of time outside – perhaps morning break only – as long as they apply sunblock beforehand.

Time off school due to eczema

Sometimes children may miss a number of lessons due to hospital or GP appointments, and this will be a key concern for parents.

It may help to sit down with them and discuss the dates of these appointments and ways in which you can work together to prevent the child from falling behind. For example, a series of short activities based on what their classmates will be learning while they are away can be used by Mum and Dad during the journey to and from the hospital or while sitting in the waiting room.

Occasionally, children with severe eczema need to be kept at home or hospitalised due to infection. If the child is well enough, you can provide some activities so that they do not fall too far behind with schoolwork.

If a child with severe eczema is admitted to hospital for a stay, it is important to liaise with the parents about provision of hospital school.

A teacher may need to liaise directly with the hospital teacher to give information on the child’s educational needs and current school work and curriculum.
Infection and eczema

When someone has eczema the skin does not provide an adequate level of protection as the skin barrier is defective, so moisture is lost - causing the skin to become very dry - and external irritants can penetrate the body more easily and cause inflammation. Inflammation leads to scratching, which leads to more broken skin, leaving the body vulnerable to a range of infections.

Bacterial skin infections

Infection is generally caused by the bacterium *Staphylococcus aureus*, which is the most common skin infection. Unfortunately, people with eczema are more susceptible to *Staph. aureus* skin infections.

You can recognise this type of infection as the area becomes redder, itchier and will weep. There is often yellow ‘crusting’ and sometimes small red blisters may also be present. Eczema infected with bacteria takes longer to recover, as the body has to take control of both the eczema flare-up and the infection. Infected eczema may cause the child to have a raised temperature (they may complain of feeling ‘fluey’). If the infection is severe, the child will need to have time off school (although they are not infectious to other children, they will feel generally unwell for a couple of days until the antibiotics have treated the infection).

It is important to note that even though infected eczema is caused by the same bacteria as impetigo (a common skin infection that mainly affects children), infected eczema is not the same thing as impetigo. Like any other child, if a child with eczema has a diagnosis of impetigo, they should be kept off school for 48 hours, from commencement of antibiotics, as impetigo is highly contagious.

Once an infection has been identified, a child’s treatment regime will need to be stepped up. This may include:

- Applying emollient more frequently and/or changing to antimicrobial emollients.
- Using topical steroids – as this is an eczema flare.
- Taking a course of oral antibiotics, which will need to be administered during the school day.
- Application of a topical steroid or antibiotic cream (for localised infection) – applied at home.

In addition, as infections thrive in warm, moist conditions, the child will have to stop bandaging and wet wrapping, if this is a normal part of their daily treatment regime, until the infection has completely gone.

Fungal skin infections

Fungal infections generally affect small parts of the body and are treated with twice daily antifungal creams. They do not make eczema worse or cause a child to be ill. Children with eczema are more prone to fungal infections.

**Candida**
A yeast infection that thrives in warm, moist folds of skin, such as the groin.

**Ringworm**
A fungal infection that can often look like eczema.
Viral skin infections

**Chickenpox**
A viral infection caused by the Varicella zoster virus, chickenpox is very common in all children. This is a particularly unpleasant condition for a child with eczema because eczema often gets worse when the chickenpox is crusting and resolving and there is additional itch due to chickenpox.

**Molluscum contagiosum**
This is a very common childhood virus that any child can get, although children with eczema are more prone to it. It is seen as small clusters of pearly lumps; some children have many crops all over their body while others have only one or two. It is contagious but is generally not treated. Molluscum will simply disappear when the child’s immune system rejects the virus.

**Herpes simplex**
Better known as the ‘cold sore virus’, herpes simplex is a high-risk infection for those with eczema as it can lead to the development of eczema herpeticum, which can be life-threatening. If **eczema herpeticum develops**, onset is rapid and the condition needs to be treated with antiviral drugs and will usually require hospitalisation. It is vital that a child with eczema does not have close contact with anyone with herpes simplex.

---

**What can I do to help?**

- Ensure that the child does not come into contact with anyone with herpes simplex.
- If there is a case of chickenpox anywhere in the school, forewarn the parents at the first opportunity.
- Ask the parents to alert you if they suspect the child’s eczema has become infected as, if it has, the child will be itchier and more uncomfortable than usual and you may need to amend your lesson plan accordingly.
- If an infection is diagnosed, ask the parents to update you on the new treatment regime and any implications for care during school hours, especially the administration of antibiotics during the school day.

- A child might already use a pump dispenser at school for their emollient to prevent it from becoming contaminated – this is especially important if the skin is infected.
- If the infection is particularly bad and the child is generally feeling unwell so needs to stay home, provide parents with activities that they can do with their child to prevent them falling too far behind with schoolwork.
- Ask the parents to confirm when the infection has completely cleared and if the child’s treatment regime during school hours has changed (e.g. a child might not simply revert to their pre-infection treatment regime, as their GP may decide that changes need to be made to help prevent another infection).
Emotions

As well as the physical effects of eczema, the child and their family may also face huge emotional challenges.

‘Why me?’ is a common question parents face – especially if the child doesn’t know anybody else with the condition – closely followed by ‘When will it go away?’

A child might feel...

Anger at the perceived injustice of having eczema when others don’t.

Frustration when none of the treatments they try ‘cure’ the eczema completely.

Self-conscious because they look different to their peers.

Left out when they can’t take part in the same activities as their friends (sleepovers, etc).

Equally, remember that some children actually cope well with eczema and that the parents can be more anxious and worried than the child.

Meanwhile parents might feel...

Guilt for the fact the child has eczema, either because the child has inherited their atopic tendency or because they feel they could have done something ‘differently’ prior to conception, during pregnancy or in the early years of life to prevent the condition from developing. (There is currently nothing that parents can do to prevent eczema from developing.)

Frustration at being unable to ‘cure’ their child’s eczema and at the lengthy ‘trial and error’ process sometimes associated with identifying the best treatment regime, especially for children with severe eczema.

Overwhelmed by the amount of care that a child with eczema can require, such as:

- spending extra hours each day applying emollients and other treatments;
- constantly cleaning the house to eliminate triggers;
- scouring the Internet and other information sources for specialist items such as seam-free clothing, special washing powder, 100% cotton school uniforms, etc;
- scheduling and attending GP, hospital and clinic appointments and organising prescriptions; and
- getting up several times in the night to soothe and cream a distressed child – broken sleep can be very draining for the whole family.

Grief for the family life they had imagined during pregnancy, which was no doubt very different to the one they find themselves living. This is especially true when the child has a sibling without eczema as the parents will be very aware of what they are missing out on – decisions around trips out and holidays, etc, will automatically be driven by the needs of the child with severe eczema.

What can I do to help?

Talk, talk and talk some more. Encourage a good two-way communication with parents so that you know how the child is doing at home and are aware of any relevant issues and so that the parents know how they are getting on at school.

In addition, make sure that the child knows that you are always there to talk to. Children with eczema are often more aware than their parents might realise of the demands and constraints that their condition places on the family and can, as a result, shy away from telling Mum and Dad things as they don’t want to upset them.
Parent–teacher meeting checklist

While we’ve tried to keep this information pack as short as possible, we appreciate that there is a lot to digest, especially if you’ve never come into contact with eczema before.

That’s why we’ve put together this handy list of questions to go through with parents before their child joins your school or class.

Where relevant, we have also listed the page numbers you can refer to for additional information on a particular subject.

It might be helpful to include such questions in the child’s ‘New starter’ induction pack for parents to complete so that you have the opportunity to go through the information before meeting them for the first time. Copies are available for download on our website at www.eczema.org/eczema-at-school-eczemainformation

### Checklist

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<th>Name of child</th>
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<td>Class</td>
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1. **When did the child’s eczema first develop? How has the condition changed since it was first diagnosed? Has it improved? Has it got worse? Has the eczema spread?** What is the eczema like when at its best and when at its worst?

2. **Do they have any other health conditions? If so, do these have a bearing on their eczema?**

3. **How does the child feel about their eczema? Do they accept it as part of them? Are they very self-conscious about it? Do they get frustrated? Does how they feel change over time? (See page 16.)**
4 Is the child under the care of a consultant dermatologist? How often do they see them?

5 Is there a pattern to the child’s eczema (e.g. worse in winter and better in summer)? How often do they suffer from flare-ups? (See pages 6–7.)

6 What treatments does the child currently use? Emollients? Emollients and a topical steroid and/or other additional treatments? Are there any side effects that you should be aware of or special precautions that you must take (e.g. keeping the child out of the sun, etc)? (See pages 8–10 and 11–13.)

7 What is their current treatment routine (i.e. when are treatments applied during a normal day)? How will this routine translate in terms of the school day? (It might be that the child’s entire routine – during and outside of school hours – will need to be amended. For example, some parents apply emollient more frequently outside of school hours than they normally would to compensate for less being applied during the school day.)

8 Ideally, how much emollient should the child apply during the school day, when and to what parts of their body? Can the child apply their own emollient? Are they confident with it? Are they happy to do it or is refusal common? What is the best approach to getting the child to co-operate? What if they won’t and are getting very distressed? Confirm where the child will be applying their emollient, who will be overseeing it and where the emollient will be stored between applications. Suggest to the parents that it might be beneficial if they come in and oversee the first application on school premises. (See pages 9–10.)

9 Ask the parents if they will be supplying a soap substitute (or using emollient as soap substitute) and whether the child needs their own towel to dry their hands on. If so, ask how often a fresh one will be supplied. (See page 6.)
10 Ask if the parents can supply a pump dispenser for emollient application and a travel-sized pump dispenser or small, separate pot that the child can keep in their tray for quick ‘top ups’ during the day (e.g. after wet play and for hand washing). (See pages 6, 9, and 10.)

11 Does the child’s treatment routine change seasonally (i.e. do they use a heavier emollient during winter because of the drying effects of the biting cold outside and central heating indoors)?

12 What are the child’s main triggers? How do the parents propose overcoming these in a school environment? Will they be supplying additional items – cotton gloves for example – for use during school hours? If so, is the child familiar with them and happy to use them? (See pages 6–7.)

13 If clothing is an issue, confirm how flexible the school is in terms of uniform. (See pages 6 and 12.)

14 Ask how itching and scratching are handled at home (e.g. do the parents use distraction techniques or does the child squeeze a ball or pinch their skin to prevent themselves from scratching and making their skin bleed)? Does the child get itchier at particular times of the day? (See pages 4–5.)

15 How often will the child need to take time off school to attend medical appointments? How much notice of these can the parents give?

16 Does the child suffer from disturbed sleep? How often does this happen? How do the parents feel this will affect the child’s performance (i.e. do they bounce back quite quickly)? (See page 16.)
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<td>17</td>
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<tr>
<td>Does the child’s eczema get infected? Does this happen often? (See pages 14–15.)</td>
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<tr>
<td>If food is an issue for the child, confirm if the school can provide lunches for children with special dietary requirements. If not, discuss the arrangements for packed lunches. Discuss possible implications for things like cookery lessons. (See page 7.)</td>
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<td>Ask the parents if the child will be participating in PE and swimming lessons and discuss the implications in terms of needing to keep cool, sweat, chlorine, applying emollient and having an eczema-friendly PE kit. For example, some children will need to wear trousers rather than shorts to protect their skin from further damage. (See pages 6–7.)</td>
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<td>If the child’s condition or medication – or both – require sun cream to be applied, ask how this will fit in with their emollient regime, at school, how much should be applied and when and whether the child is able to apply their own sun cream and is comfortable doing so. (See pages 9, 11–13.)</td>
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<td>Inform the parents of any planned school trips during term time so that they can think about potential issues and solutions in advance.</td>
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<td>As parents will worry about bullying, it would help to alleviate their concerns if you can explain the school’s approach, so they know in advance the support that is available and how any issues would be addressed.</td>
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<tr>
<td>Ask the parents to keep you informed of any changes in the child’s medication, treatment routine, forthcoming medical appointments and any other issues that may have a bearing on their condition.</td>
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Eczema – planning checklist

If you’ve no or little previous experience of having a pupil with eczema in your class, it can be difficult at first to keep on top of their treatment regime and to remember all their individual triggers. That’s why we’ve included this at-a-glance checklist for your planning folder so that when putting together a lesson plan you can easily identify possible issues for the child, which can then be highlighted to all the adults involved with the lesson.

Copies are available for download on our website at www.eczema.org/eczema-at-school-eczemainformation

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### Treatment...

- When does the child need to apply emollient during the school day?
- When does the child need to be reminded to ‘top up’ their emollient (e.g. after wet play, etc)?
- Do they also need to apply sun cream? If so, when and where?
- Who will oversee each of these applications (assuming the staff members rotate)?
- Is the child using their soap substitute, if one has been supplied?
- Are there any side effects from the child’s current medication that the team needs to look out for?
## Potential triggers...

*Tick any that apply*

- Being too hot or too cold or sudden temperature changes
- External conditions (e.g. very sunny, bitterly cold, high pollen count, etc)
- Natural environment (e.g. contact with grass, plants, trees, etc)
- Animals and birds
- Soap
- Wet play
- Chemicals (e.g. glue, paint, clay)
- Clothing
- Sweat
- PE
- Swimming
- Dust
- Carpet
- Certain foods

## Visual clues for the team to watch out for...

How does the child convey that they are itchy? By pinching their skin, squeezing a ball or using everyday items such as a ruler or pen to scratch themselves with?

What usually precedes a bout of itching (e.g. does the child become quiet and withdrawn or louder and more disruptive)? Do they suddenly go very still or do they continually fidget? Do they go red in the face, etc?

How sore are the child’s hands? Can they hold a pencil comfortably? Can they take part in ‘fine work’ such as sewing?
Activities

Eczema is a complex condition and one that affects a growing number of families in the UK. Highly visible, it attracts attention, speculation and comment, which is why it’s so important for a child’s peers to have a better understanding of its causes, symptoms and treatment as well as the everyday challenges that it presents.

If you had no experience of eczema prior to reading this guide, we appreciate that there’s a lot of information to take in and that you might not immediately feel confident giving a lesson on the condition. Equally, we recognise how time-consuming lesson planning can be.

That’s why we’ve worked with a team of highly experienced teachers to produce a series of lesson plans, including a variety of activities and resources.

We suggest the following activities for different age ranges, but please feel free to use your discretion as to what will work for your class.

3–5-year-olds | 5–7-year-olds | 7–11-year-olds
Activities 1, 2 and 3 | Activities 4, 5 and 6 | Activities 7, 8 and 9

Designed to encourage understanding and compassion, each lesson aims to be no more than 20–30 minutes long, with an emphasis on both the physical and the emotional impact of eczema on a child.

Finally, whether you use the lesson plans as they have been provided or decide to modify them to suit the specific needs of your pupils, we hope that they provide a useful framework for teaching the next generation about this incurable and increasingly common condition.

Thank you once again for your help and support.
Today we are going to be learning about this word (have the word ‘Eczema’ displayed on the board). It says ‘eczema’.

Has anybody heard this word before?

Explain to the class what eczema is – a dry skin condition where the person has red patches of skin that are very sore and itchy.

**Activity and Feedback**

- Give each child a piece of play dough and ask them to roll it flat.
  - *How does this feel?*
- Now ask them to scratch the surface.
  - *What happens?*
- Now ask them to scratch it again.
  - *What happens?*
- Repeat the scratching one more time.
  - *What happens?*

**Conclusion**

Explain to the class that this is what happens to a person’s skin when they scratch their eczema, which is why they have to try to stop themselves scratching as it will make their skin worse.

This is why a person with eczema needs to keep putting cream on their skin as it helps to stop it being so dry and itchy and to get better.
Can anybody remember what this word (have ‘Eczema’ written on the board) says? Remind the class what the word says and that it is a dry skin condition where the sufferer has red patches of skin that are very sore and itchy.

Activity
Take the class into the hall or another large space and invite the child with eczema to stand at the front and take on the role of PE teacher, asking the other children to do ten star jumps or hop on the spot – any sort of physical activity. It should be for about 2–3 minutes.

Feedback
With the class sitting down, ask them how they are feeling, how does their skin (particularly their face) feel and how do they feel in their clothes.

Conclusion
Explain to the class that how they are feeling would not be good for a person with eczema as it will make their skin even more sore and itchy and that’s why you asked the child with eczema to be the PE teacher instead.

Explain to the class that many things in everyday life can make a person’s eczema worse, even if they have put their cream on.

Show the class the trigger list, found on page 26 or from our website www.eczema.org/eczema-at-school-activity2
Triggers...

- Clothing and uniform
- Sweat
- Temperature
- Pollen
- Soap and water
- Animals
- Messy or wet play, art and cookery
- Chairs
- Dust
- Carpet
- Swimming
- Food
- Damp and mould
- Fragrance
Remind the class that eczema is a dry skin condition where the sufferer has red patches of skin that are very sore and itchy.

People with eczema need to put cream on to protect their skin and help it to get better.

There are also many everyday objects or items that can make it worse. Show the class the feeling cards (only use Happy, Sad, Angry and Excited) and explain to them what each feeling is. A template is available on page 28 and from our website www.eczema.org/eczema-at-school-activity3

Activity

Sit the class in a circle with each child having a set of feeling cards (Angry, Sad, Happy and Excited). Ask the children to listen to the following situations and, at the end of each, to hold up a card to say how they would feel in this situation.

> Everyone else in your class is going to Ben’s birthday but your Mummy says you can’t because you’re already doing something else.

> Your best friend has a cat but you can’t have one because cats make your Daddy sneeze.

> You’re at the beach and want to go and play but your Mummy is spending ages putting sun cream on you because she doesn’t want you to burn.

> You’ve been poorly so Daddy says you can’t go swimming this weekend in case it makes you ill again.

> Some children at school keep pointing and laughing at you because you have to wear glasses.

Feedback

Every time the children hold up a feeling card, ask one or two children to explain why they are feeling like this.

Conclusion

Tell the class that how they have felt in these situations can often be how people with eczema feel as the condition makes their life more difficult. We need to be kind and friendly to everyone we meet so that people don’t feel sad.
Feeling cards

Happy

Sad

Angry

Excited

Worried

Frustrated

Activity three  Recommended for 3–5-year-olds
Activity six  Recommended for 5–7-year-olds
Today we are going to be learning about this word (have the word ‘Eczema’ displayed on the board). Does anyone know what this word says? Tell the class if they don’t. Explain to the class what eczema is – a dry skin condition where the sufferer has red patches of skin that are very sore and itchy.

Activity

Explain to the class that people with eczema have very itchy skin and that scratching makes it worse so they have to try not to do it. Do you think this would be easy to do? Ask the class to sit on their hands and tell them that you are going to ask them a series of questions but they mustn’t put their hand up to answer (or shout out). In order to generate a reaction from the children, pick questions that they will really struggle not to answer such as ‘Who likes chocolate?’, ‘Who has a pet?’ and ‘Who likes football?’

Feedback

You should hopefully find that the children (well most of them!) were not able to sit on their hands but put their hands up to answer your questions! Ask the children to put their thumbs up or down depending on how easy they felt it was to not put their hand up to answer questions.

Conclusion

Explain to the class that this is how a person with eczema feels – they know they mustn’t scratch because it will make their eczema worse but it is very hard not to. Add that this is why a person with eczema needs to keep putting cream on their skin as it helps to stop it being so dry and itchy.
Spot the Triggers

Remind the class that eczema is a dry skin condition that makes the skin red, sore and very itchy.

People with eczema need to put cream on to soothe and moisturise the skin so that they don’t scratch and make it worse. Tell the class that many things in everyday life can make a person’s eczema worse, even if they have put their cream on.

Show children the list of triggers and explain how and why each is a trigger. It’s available on page 26 and from our website www.eczema.org/eczema-at-school-activity5

Activity
Children should be in groups of four or five and have the picture of a classroom (see page 31 of the pack and also available from www.eczema.org/eczema-at-school-activity5).

Keeping the trigger list on the board, ask the children to draw a circle around the items that they think are triggers.

Feedback
Ask each group to say what they have drawn their circles around. Refer to the trigger list and ask if that was on the list.

If you feel it is appropriate, you may want the children to explain why they have drawn circles around an object.

Conclusion
Introduce the feelings that a child with eczema may have because they cannot do what their friends can do so easily – Angry, Frustrated, Worried and Sad.
Activity five

Spot the triggers

Recommended for 5-7-year-olds

Produced by the National Eczema Society  www.eczema.org
Social and emotional issues

Remind the class that eczema is a dry skin condition that makes the skin red and very itchy.

People with eczema need to put cream on to protect their skin and help it to get better.

There are also many everyday objects and items that can make it worse. Show the class the feeling cards and explain to them what each feeling is. If you do not already have these, a template is available on page 28 or from our website www.eczema.org/eczema-at-school-activity6

Activity

Sit the class in a circle with each child having a set of feeling cards (Angry, Sad, Worried, Happy, Excited and Frustrated).

Ask the children to listen to the stories (see page 33 of the pack and also available from www.eczema.org/eczema-at-school-activity6) you are going to read and, at the end of each story, they are to hold up a card to say how they think the child in the story is feeling.

Feedback

At the end of each story as the children are holding up a feeling card, ask one or two children to explain why they have chosen that feeling.

Conclusion

Tell the class that eczema makes life more difficult for people and that they can feel angry, sad, worried and frustrated as a result.

We need to be kind and friendly to everyone we meet so that people don’t feel sad.

Ask the class to give you examples of how they could make another child happy.
Here are five mini stories about children with eczema.

**The party**
Abigail’s birthday party was in one week and she was VERY excited. But there was a problem. She’d seen the prettiest dress in the whole world in a shop window in town. But her Mum said that the stiff, scratchy material would only irritate her skin. It wasn’t fair. It just wasn’t fair. Just for one day she wanted to not have to worry about her eczema and surely that one day should be her birthday? Just for one day couldn’t she be like all her other friends and just wear the pretty dress? Even if it did make her itch!

**New school**
Tom is starting a new school. His Dad is driving him there. His skin is not feeling good today. Tom’s cheeks are red and there are sore patches on his neck. Why couldn’t his eczema have just stayed hidden, today of all days? Now everyone is going to know he is ‘different’ to them. What if it is like his old school? It had taken months for the other children to get used to how he looked and his creams. What if nobody wanted to talk to him or play with him? The car pulled into the car park and Tom wished his Dad would take him straight back home.

**Sweet dreams**
The big box of chocolates suddenly leapt off the shelf and started chasing Sally around the store. Faster and faster she ran but it wouldn’t leave her alone. Past the fruit and vegetables, down the cereal aisle, it was definitely catching her up! With a start, Sally woke up. Looking around her room she wondered what had jolted her from her sleep and then, there it was, the dreaded itch. Itch, itch, itch, itch. She made a fist with each hand and tried counting to ten. Don’t scratch, don’t scratch, don’t scratch. Lying in her bed Sally tried to think of nice things – seeing her friend Maisie at the weekend, her favourite TV programme, that new song she’d heard on the radio – but the itch just wouldn’t go away. Sally yawned. And yawned again. She was going to find it really difficult to sleep.

**Spike**
Sam waves to Amy across the street and shouts, ‘Come and see what I’ve got’, pointing excitedly at his back garden. As she walks down the path, Amy wonders what Sam’s new toy might be. A climbing frame? A bike? Maybe even the treehouse he asked for? Turning the corner she sees that Joe and Lisa are sitting on Sam’s lawn, with their eyes glued to a patch of grass at his feet. They see a flash of brown fur and then they hear a loud ‘woof’. Sam reaches down and picks up a very cute, very wriggly puppy and holds it up for Amy to see. ‘Come and meet Spike!’ Amy looks at Sam and shakes her head. She’d love to stroke the pup and let him cover her in wet kisses, like the others are doing now, but she knows that furry animals always make her skin itch if she gets too close.

**Splash**
Ryan watched the other children put on their swimming costumes and hurry out of the changing room into the pool. They couldn’t wait to play with all the inflatables and toys that were bobbing in the water. Ryan couldn’t wait either but his Mum was busy applying his cream and it was taking aaaaaaaaaages. The clock ticked. ‘Can I go now? Preeeeeeease?’ Mum shook her head. Tick, tick, tick. He could hear the other children shouting and laughing and tried to imagine what they were doing. ‘I have to do it for your eczema’ she added. How many times had Ryan heard that? Sighing, he wondered what it must be like not to have such troublesome skin.
Have the word ‘Eczema’ displayed on the board. Ask the children to write down what they think eczema is. Take feedback from the class and write down their ideas on the board.

**Activity**

- With the class in pairs ask them to build a wall using Lego/Duplo/Multilink.
- Now pour shredded paper or sand over it.  
  **What happens?**
- Repeat the exercise but this time ask them to leave lots of gaps in the wall starting from the top row down.  
  **What happens?**
- Repeat again (with the gaps) but place a solid barrier over the top of the wall.  
  **What happens?**

**Feedback and Conclusion**

Show the class the diagrams from page 8 of the pack – also available from [www.eczema.org/eczema-at-school-activity7](http://www.eczema.org/eczema-at-school-activity7) – and discuss how:

- ‘Healthy skin’ with its solid brick wall keeps moisture in and external irritants out.
- A child with eczema has skin like the second wall the class built, with lots of little gaps between the skin cells. These gaps allow moisture to escape from the body, causing the skin to become very dry, and allow external irritants in (as demonstrated by the shredded paper or sand) causing it to become red, inflamed and itchy.
- Children with eczema use creams that form a barrier to help prevent further moisture loss and penetration by irritants, which in turn means that the skin isn’t as red, sore and itchy.
Triggers

Take feedback from the class on what eczema is and how it can affect children. Tell the class that many things in everyday life (triggers) can cause eczema to worsen and these vary from person to person.

Show children the list of possible triggers (available on page 26 or from www.eczema.org/eczema-at-school-activity8) and explain how and why each is a trigger.

Activity

Put children in pairs or groups of four or five for this activity.

Give each pair or group a different scenario (see page 36 of the pack and also available from www.eczema.org/eczema-at-school-activity8) and ask them to say what possible triggers a child in this situation would face.

Feedback

Take feedback from each pair or group as to what possible triggers a child would face. Does the rest of the class agree? Are there any others anyone can think of?

Possible answers:

- **Sports day** – the heat of the day, getting hot from running, sweat, grass, pollen and shorts elastic or material rubbing on bare skin could all make eczema worse.
- **Sleepover** – make-up, hairspray, perfume, the heat from the radiator and dust in the room could all make eczema worse.
- **School play** – overheating in the costume, overheating from the spotlights, the costume material, dust, paint and glue could all make eczema worse.
- **Camping** – going from hot to cold, the label or seam chafing her neck, grass, trees, pollen and the dog could all make eczema worse.
- **Beach** – heat from the sun, sand, sea (salt can sting if you have sore skin already, and going from hot to cold can also cause an itching frenzy), face paint, sweat and a rough towel could all make eczema worse.

Conclusion

Show how a child with eczema often has to:

- take extra care in order to participate (e.g. Jessica comes to the sleepover with her own duvet and towels).
- experience discomfort in order to participate (e.g. Adam doesn’t like the beach as the sand makes him itch but he doesn’t want to miss out on a family trip).
- not participate (e.g. Amy couldn’t play with the dog she met at the top of the hill as animals make her eczema worse).

Introduce the emotions that a child with eczema may have because they can’t do as easily what their peers can – anger, frustration, feeling self-conscious and left out.
Scenarios

Here are descriptions of five different scenarios that a child with eczema finds themselves in.

Sports day

It’s the annual school sports day and Ben is competing in the 100m sprint and relay race. In between events he sits on the grass to cheer on the rest of his team. It’s warm and sunny, and lots of parents have come to watch. Ben can see his little brother waving at him. In the end it all comes down to the relay race. If Ben’s team wins it, they’ve won overall. Ben adjusts his new shorts, waits for the word ‘go’ and runs as fast as he possibly can, faster than he ever has before, and crosses the line… first!

Sleepover

It’s Alice’s birthday sleepover, and Jessica and the rest of the girls are just arriving. Jessica’s bag is nearly as big as she is as she’s brought her own duvet and towels with her. Alice knows it’s because the washing powder her Mum uses to do the laundry might irritate Jessica’s skin but the others don’t and are whispering behind her back. Chloe leans against the radiator and suggests giving each other makeovers using the make-up, nail varnish and hairspray she’s ‘borrowed’ from her older sister’s room. Alice gives everyone a go with her Mum’s fancy perfume. Soon the smell of pizza drifts up the stairs....

School play

Jamie has the lead role of King in the school play and is really struggling to learn his lines. Miss Langley is not helping. ‘No, no, no, NO!’ is all he’s heard for the last half an hour. Worse still, even when he did finally come out with the right line, it turned out to be someone else’s! Standing in an oversized – and very heavy! – robe and crown, Jamie can see the dust swirling in the spotlight every time the curtain is pulled back and another pupil is thrust onto the stage to join him. If only he’d volunteered to help with the scenery – it looked a lot more fun painting castle walls and gluing leaves on to fake trees. Maybe if he offered to help....

Camping

It’s warm in the car and Amy is determined to stay in it. But her Dad has other ideas. ‘It will be fun’, he keeps saying as if, the more often he says it, the more fun everything will suddenly be. Amy and her brother peer out of the windows at the grey sky. Still, Dad begins taking the tent out of the boot and putting it up. With a groan Amy pulls on her fleece, waterproof jacket and gloves and follows him. The cold breeze makes her ears burn and she pulls on the woolly hat that Granny knitted. Once the tent is finished it’s time for a walk and a picnic. As they march up the hill, one after the other, Amy can feel her nose getting colder and something rubbing her neck but, with so many layers on and her gloves, she can’t reach whatever it is. At the top, hiding in the long grass, are two furry ears and a wagging tail.

Day trip to the beach

It’s hot, very hot, and the Robertson family have gone to the beach for the day. Adam doesn’t like the beach as the sand makes his skin itch. Mum has already covered them all in sun cream, which means the sand now sticks to his skin even more! To avoid building sandcastles with his little sister he offers to collect shells for her instead. The sea looks lovely and cool to paddle in as he walks down towards the pier where two girls are doing face painting for charity. By the time he gets back and sits down in a deckchair, it’s even hotter and sweat is running down his back. Reaching for a beach towel he rubs his face.
Children should either be in pairs or groups of four or five for this activity. Ask the children to write down what they already know about eczema.

Take feedback from each group. Tell the children that they will be focusing on the feelings a child with eczema may have and what social difficulties they may face.

**Activity**
Give each pair or group a different ‘Ask Amy’ question (see pages 38 and 39 of the pack and also available from www.eczema.org/eczema-at-school-activity9). Ask them to discuss how the child in the question is feeling and then ask them to write a response.

**Feedback**
Take feedback from each pair or group and then read how Amy actually responded to the question (see pages 38 and 39 of the pack).

**Conclusion**
Remind the class that children with eczema face additional challenges in everyday life and that understanding is needed.
Ask Amy

Q My little sister is seven and has eczema. The other week I found out that people at school are teasing her about it and calling her names. She hasn’t mentioned it to Mum and Dad and the only reason I know is because a friend of mine caught a group of girls laughing at her and saying she had fleas because she can’t stop scratching. I’ve tried to talk to her about it but she keeps telling me that it’s ok. I don’t want to go behind her back and tell our parents or a teacher but I don’t know what else to do. What can I do to help her?

Steven

Amy says...

School can be tough for anyone who stands out from the crowd. Whether you have a funny surname, an unusual hobby or like music that nobody else in your class likes, at some point someone is likely to pick up on why you’re different and tease you about it.

The most important thing your sister can do is to show that she’s a normal girl just like them. She should therefore make a real effort to participate in things that the other girls in her class are doing both in and out of school. Yes, there may be things she can’t do due to her eczema or that require some changes in order not to inflame her skin but, for the most part, there’s no reason she shouldn’t do a lot of the things that her classmates do.

Ask your friends to keep an eye out for her so you know if things seem to be getting better or worse without constantly asking her. Make sure she knows that you are always there to talk to if she needs a friendly ear.

In terms of telling an adult, I would always advise that someone is kept informed. If you feel that your sister would react very badly to this, then perhaps see how the situation develops over the coming weeks and, if it’s getting worse, speak to a teacher.

Q I love football and am a member of the school team but this year the cold, wet weather is making my eczema worse and worse. A group of boys have also started giving me hassle in the changing room afterwards as I have to apply my creams, which they think makes me a girl. What can I do?

Andy

Amy says...

It’s clear that playing sport is making your eczema worse, so ask your teacher if you can be a spectator for the next 2–3 weeks while your skin repairs itself.

Once your skin is under control again, rejoin your team but be sure to take the following extra steps:

▶ Apply cream before a game as well as afterwards to ensure your skin is fully protected.

▶ Ask your teacher if you can wear trousers instead of shorts to create a further barrier between your skin and the harsh elements and tackling.

▶ Make sure all your sports kit is made of cotton, which will enable your skin to breathe, reduce the risk of chafing and prevent overheating.

Respond to any comments that are made by saying that your doctor has prescribed the creams to treat your sore skin and that it’s something you just have to do to get better in the same way an asthma sufferer needs their inhaler or someone with a broken leg needs to use crutches. If the problem persists, tell your teacher so that he can intervene if any comments are made.
Q I’ve started going out with a girl who I really like but I find it impossible to talk about my eczema with her. We’ve only known each other a few months during which my eczema has been really good but my skin is always worse in winter and I’m dreading how she’ll react when it does flare up. What should I say to her?

Joe

Amy says...

You’re not alone as many young people write to me about how to handle their eczema in a relationship.

The positive thing to take from all of this is that she likes you for who you are and clearly wants to be with you, so finding out you have eczema should not affect the way she feels.

Not everyone knows about eczema, so it’s worth starting with the basics so she fully understands why you have the condition and what it means for you. Explain what a flare-up is like and which aspects of the winter months are responsible for your skin getting worse – for example, the cold weather, central heating, heavy clothing, etc.

Finally, encourage her to ask you as many questions as she wants – the more she understands the condition, the easier it will be for both of you.

Q I’m due to start a new school in September and I’m really nervous as I won’t know anybody. At my old school I had a big group of friends and we hung around together all the time. Mum says I’ll make new friends but I’m scared that nobody will want to talk to me or sit next to me because of my eczema. What can I do?

Katie

Amy says...

Starting a new school is challenging for everyone so you’re not alone in your fears!

There are ways in which you can reduce your anxiety though. Here are my top tips:

> I’m sure you won’t be the only new pupil joining your year group in September, so why not ask your parents to speak to the Head about arranging a get-together? Not only will you be able to meet all the other new starters – all of whom will be as nervous as you! – and get to know them before the new term starts, but you might also find you make some good friends in the process.

> You are obviously a really good friend, as you had so many at your old school, so why not ask them what it is they like about you? This is a fantastic way to boost your confidence as it will remind you of all the great things you have to offer to potential new friends!

> Before term starts, think about all the questions people might ask you about your eczema. If you know in advance how you want to answer each of these, you’ll make sure you come across as friendly and confident rather than shy and embarrassed. Not everyone knows what eczema is, so by answering questions you can help to alleviate any concerns they may have, such as it being contagious.

> Finally, don’t be upset if someone refuses to be friends with you because of your eczema. After all, if they decide who they want to be friends with solely based on how they look, are they actually worth being friends with at all?