Fever and Rash Pathway

Clinical Assessment/ Management tool for Children





First Draft Version: November 2017 Review Date: November 2019.

Management - Acute Setting

Could petechiae be explained by vomiting/coughing? **SVC** distribution Not progressive (all children with purpura need urgent paediatric review) Manage underlying cause. Provide patient advice sheet

Anaphylaxis / angioedema affecting airway YES - Haemodynamic compromise **ACUTELY ILL** Purpura/petechiae/bruising? **NOT ACUTELY ILL** Skin blisters / significant erosions Mucosal blistering Skin pain YES Fever ≥38° in child under 3 months of age YES 90% Body Surface Area = ERYTHRODERMA YES 5 days of fever >38°C Systemically unwell Atypical / severe rash: e.g. coalescing rash or intense erythema or pustules or large surface area involvement YES If no red or amber flags, consider: Infectious exanthems (Viral or bacterial) Primary bacterial infections of the skin Allergy / Urticaria

EMERGENCY ACTION: Contact Lead ED / Paediatric Doctor. Move to Resuscitation Area. Resus Call ("2222") for Paediatric Arrest - Anaphylaxis or sepsis

Immediate medical intervention as appropriate such as: Anaphylaxis: IM Adrenaline Sepsis: immediately initiate sepsis 6 (See sepsis pathway)

IMMEDIATE REFERRAL TO PAEDS

Purpura may be a feature of Vasculitis (purpura are palpable). Causes of fever and vasculitis: think infection. Other causes possible e.g. autoimmune, drug reactions etc. Associated joint or abdo pain: think Henoch Schonlein Purpura

If non-palpable purpura, may be early vasculitis, but other causes may not need urgent referral: platelet dysfunction (e.g. NSAIDs), bruising, related to vigorous coughing

Fever and blistering may be a feature of: Stevens Johnson Syndrome, Toxic epidermal necrolysis, Staphylococcal scalded skin syndrome, Eczema herpeticum, Toxic shock syndrome

Possible late onset sepsis. Refer to "fever in children less than 5 years" clinical pathway.

IMMEDIATE REFERRAL TO PAEDS

Fever and widespread rash may be a feature of:

Infected eczema, drug reaction

Systemic inflammatory process

Kawasaki disease

CLOSE MONITORING IN 1° CARE / ED OR SEEK ADVICE FROM PAEDS. Provide patient advice sheet.

Fever and severe/atypical rashes may be a feature of:

Drug Hypersensitivity

Sweets disease

Atypical infections

Erythema Multiforme

Infected eczema

Acute psoriasis

This guidance was written in collaboration with the SE Coast SCN and involved extensive consultation with healthcare professionals in Wessex

Provide appropriate and clear

guidance to the parent / carer

and treatment if required.

Provide patient advice sheet.

Confirm they are comfortable

with the decisions / advice

given and then think

"Safeguarding" before sending home.

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Differential Diagnosis	Distinguishing features
Viral infections	
Measles	Erythematous maculopapular rash over hairline/forehead and behind the ears, spreading caudally. Koplik spots (gray papules on buccal mucosa). Overwhelming misery. Obtain vaccine history.
Rubella	Eythematous maculopapular rash on face, spreads to extremities. Tender lymphadenopathy (occipital, postauricular, cervical)
Chicken pox	Consider bacterial secondary infection if fever 3 or more days after onset of chickenpox or increasing erythema around lesions. Consider toxic shock syndrome if haemodynamically unstable, generalised rash (erythroderma) and mucosal erythema (red eyes, red lips).
Eczema herpeticum	Disseminated viral infection (usually HSV1/HSV2) characterised by fever and clusters of itchy blisters or punched-out erosions. Severe eczema herpeticum may affect multiple organs. Risk of bacterial secondary infection (staph aureus or Gp A strep).
Other viral exanthems (enterovirus, adenovirus, HHV6, parvovirus, Coxsackie virus etc)	'Slapped cheek' (parvovirus B19 –also causes hydrops fetalis) - macular erythema over cheeks, lacy eruption on extremities; Papular purpuric gloves and socks syndrome (parvovirus B19) – erythema, oedema, petechiae/purpura on palms and soles with burning/pruritis; Roseola (HHV-6) – circular/elliptical macules/papules on trunk +/- surrounding white halo – rash as fever subsides; Herpangina (various enteroviruses) – exanthem (often absent), painful grey oral vesicles; Hand-Foot-Mouth Disease (Coxsackie A16>enterovirus 71) – grey vesicles, pustules and erosions on hands, feet and buttocks with oral vesicles/erosions on an erythematous base; Infectious mononucleosis (EBV) – morbilliform rash, pharyngitis, fatigue, myalgia, hepatosplenomegally, lymphadenopathy.
Bacterial infections	
Meningococcal disease	Short prodromal phase (fever, lethargy, malaise, nausea/vomiting), followed by the more specific and severe symptoms of meningitis and sepsis (limb pain, cold hands/ feet, pale or mottled skin). Associated with non-blanching rash - a rapidly evolving petechial or purpuric rash is a sign of very severe disease.
Scarlet fever	Erythema of axilla, neck, chest, evolution to pink papules on erythematous background. Pastia's lines (linear petechial streaks in body folds). Red strawberry tongue. 7-10 days later hand and foot desquamation.
Cellulitis	Erythema, oedema and pain.
Impetigo	Bullous (only staph aureus) – flaccid see through bullae which rupture leaving a shiny dry erosion with an erythematous base +/- fever/diarrhoea/lethargy; Non bullous (S aureus >> Gp A strep) – eythematous macule – pustule/vesicle – erosion with golden crust – fever should be minimal.
Staph scalded skin syndrome	Fever, conjunctivitis, skin pain and flexural erythema with subsequent desquamation. NB culture of bullae negative.
Staph/strep toxic shock syndrome	Macular exanthem (on trunk spreading outwards), palmoplantar erythema and oedema with subsequent desquamation, conjunctival hyperaemia + hypotension + involvement of three or more organs. Risk factors include recent chickenpox and minor burns.
Secondarily infected eczema	Weeping, crusting or pain occurring on the background of eczema should prompt consideration of secondary bacterial infection.

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Differential Diagnosis	Distinguishing features
Drug Reactions	
Severe - Steven Johnson syndrome (SJS) / toxic epidermal necrolysis (TEN)	Occurring after 7-21 after exposure to a new drug. Rash often preceded by prodromal illness (fever, sore throat, myalgia, conjunctivitis). Rash starts on trunk, extends to limbs (spares soles and palms). SJS < 10% BSA, widespread purpuric macules or atypical targets. TEN > 30% BSA +/- widespread purpuric macules. Mucosal involvement including eyes, mouth/lips, pharynx/oesophagus, genitalia, upper respiratory tract and GI tract. (SJS often occurs in response to infection in the absence of a precipitating drug. Mycoplasma pneumoniae is commonly implicated, often producing a more marked mucosal pattern of disease).
Drug Hypersensitivity (DRESS – Drug Rash with Eosinophilia and Systemic Symptoms)	Occurring 7-40 days after exposure to new drug. Often morbilliform in appearance, worse initially over the face and upper body. Facial oedema is frequent. Atypical targetoid lesions, pustules, vesicles and purpura may occur. Fever, eosinophilia, lymphadenopathy, internal involvement organ (most frequently hepatitis – risk of liver failure).
Acute Generalised Exanthemous Pustulosis (AGEP)	Erythema and swelling with large numbers of overlying superficial pustules. Most commonly affecting the face and flexures initially. Occurs shortly after or within 4 days of exposure of a culprit drug (commonly antibiotics, but numerous drugs have been reported).
Frequently non-infective causes	
Sweet's Syndrome	Erythematous tender nodules and plaques, often with associated pustules. Neutrophilia and fever. Occurring in response to underlying malignancy, upper respiratory tract infection, inflammatory bowel disease, rarely drug-induced.
Erythema multiforme	Erythematous targertoid-lesion (bulls eye appearance). Minimal associated itch. Common over acral sites, but any part of the body may be affected. In some cases it can be associated with blistering and/or mucosal lesions. May be idiopathic, but Herpes Simplex Virus and mycoplasma pneumonia infection are commonly implicated.
Henoch Schonlein purpura (HSP)	Classically presents with symmetrical palpable purpura on legs and buttocks in an otherwise well child. May involve joint pain/swelling, abdominal pain and haematuria. Most commonly occurs in children aged 2-11 years. Monitor BP and urine (for blood and protein)
Kawasaki disease	Fever >5 days, rash, bilateral non-exudative conjunctivitis, oral signs (red, cracked lips), oedema of hands/feet and cervical LAN (>1.5cm). Overwhelming misery extremely common. Signs may appear and disappear before others arise. 80% of cases occur in children <5 years of age (peak incidence 1 to 2 years).