Children and Infants with Fever - Acute Management

Document Number: PD2010_063
Publication date: 11-Oct-2010
Functional Sub group: Clinical/ Patient Services - Baby and child
Clinical/ Patient Services - Medical Treatment
Summary: Basic Clinical Practice Guidelines for the acute treatment of infants and children with fever.
Replaces Doc. No.: Children and Infants with Fever - Acute Management [PD2005_388]
Author Branch: NSW Kids and Families
Branch contact: NSW Kids & Families 9391 9503
Audience: Emergency Depts, Paediatric Units
Distributed to: Public Health System, Divisions of General Practice, Government Medical Officers, NSW Ambulance Service, Private Hospitals and Day Procedure Centres, Tertiary Education Institutes
Review date: 11-Oct-2015
Policy Manual: Patient Matters
File No.
Status: Active

Director-General

This Policy Directive may be varied, withdrawn or replaced at any time. Compliance with this directive is mandatory for NSW Health and is a condition of subsidy for public health organisations.
INFANTS AND CHILDREN: ACUTE MANAGEMENT OF FEVER

PURPOSE

The infants and children: acute management of fever clinical practice guideline (attached) has been developed to provide direction to clinicians and is aimed at achieving the best possible paediatric care in all parts of the state.

The clinical practice guideline was prepared for the NSW Department of Health by an expert clinical reference group under the auspice of the state wide Paediatric Clinical Practice Guideline Steering Group.

MANDATORY REQUIREMENTS

This policy applies to all facilities where paediatric patients are managed. It requires all Health Services to have local guidelines/protocols based on the attached clinical practice guideline in place in all hospitals and facilities likely to be required to assess or manage children with fever.

The clinical practice guideline reflects what is currently regarded as a safe and appropriate approach to the acute management of fever in infants and children. However, as in any clinical situation there may be factors which cannot be covered by a single set of guidelines. This document should be used as a guide, rather than as a complete authoritative statement of procedures to be followed in respect of each individual presentation. It does not replace the need for the application of clinical judgement to each individual presentation.

IMPLEMENTATION

Chief Executives must ensure:

- Local protocols are developed based on the infants and children: acute management of fever clinical practice guideline.
- Local protocols are in place in all hospitals and facilities likely to be required to assess or manage paediatric patients with fever.
- Ensure that all staff treating paediatric patients are educated in the use of the locally developed paediatric protocols.

Directors of Clinical Governance are required to inform relevant clinical staff treating paediatric patients of the revised protocols.

REVISION HISTORY

<table>
<thead>
<tr>
<th>Version</th>
<th>Approved by</th>
<th>Amendment notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>December 2004</td>
<td>Director-General</td>
<td>New policy</td>
</tr>
<tr>
<td>(PD2005_388)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>October 2010</td>
<td>Deputy Director-General</td>
<td>Second edition</td>
</tr>
<tr>
<td>(PD2010_063)</td>
<td>Strategic Development</td>
<td></td>
</tr>
</tbody>
</table>

ATTACHMENT

1. Infants and Children: Acute Management of Fever – Clinical Practice Guideline.
Contents

Introduction .................................................................................................................................. 3

Changes to previous clinical practice guidelines ......................................................... 4

Overview ........................................................................................................................................ 5

Rationale for clinical approach ................................................................................................. 5

Assessment and initial management ......................................................................................... 6

Flowchart for child < 5 years old with fever (>38°C axillary) .............................................. 6

Toxicity: ABCD ................................................................................................................................. 7

Focus of infection ............................................................................................................................ 7

Subjective features ......................................................................................................................... 7

Rash and fever ................................................................................................................................. 7

Investigations ................................................................................................................................. 8

Lumbar puncture ............................................................................................................................... 8

Urine culture ................................................................................................................................... 9

Antipyretics ...................................................................................................................................... 9

Tepid sponging ................................................................................................................................. 9

Follow-up ....................................................................................................................................... 9

Evidence base for the acute management of fever ................................................................. 11

NHMRC designation of levels of evidence ............................................................................... 11

What is fever? ................................................................................................................................. 11

How should temperatures be measured? .................................................................................. 11

Clinical recommendations .......................................................................................................... 13
Appendix 1 – References .................................................................................................................. 16
  How should temperatures be measured? ..................................................................................... 16
  Clinical assessment .................................................................................................................. 17
  Antipyretics ........................................................................................................................... 18

Appendix 2 – Resources ............................................................................................................. 19

Appendix 3 – Parent information .............................................................................................. 20

Appendix 4 – Working party members ..................................................................................... 21
Introduction

These Guidelines are aimed at achieving the best possible paediatric care in all parts of the State. The document should not be seen as a stringent set of rules to be applied without the clinical input and discretion of the managing professionals. Each patient should be individually evaluated and a decision made as to appropriate management in order to achieve the best clinical outcome.

The formal definition of clinical practice guidelines comes from the National Health and Medical Research Council:

‘systematically developed statements to assist practitioner and patient decisions about appropriate health care for specific clinical circumstances.’


It should be noted that this document reflects what is currently regarded as a safe and appropriate approach to care. However, as in any clinical situation there may be factors, which cannot be covered by a single set of guidelines, this document should be used as a guide, rather than as a complete authoritative statement of procedures to be followed in respect of each individual presentation. It does not replace the need for the application of clinical judgment to each individual presentation.

This document represents basic clinical practice guidelines for the acute management of fever in children and infants. Further information may be required in practice; suitable widely available resources are included as appendix two.

Each Area Health Service is responsible for ensuring that local protocols based on these guidelines are developed. Area Health Services are also responsible for ensuring that all staff treating paediatric patients are educated in the use of the locally developed paediatric guidelines and protocols.

In the interests of patient care it is critical that contemporaneous, accurate and complete documentation is maintained during the course of patient management from arrival to discharge.

Parental anxiety should not be discounted: it is often of significance even if the child does not appear especially unwell.
Changes to previous clinical practice guidelines

The following outlines significant changes to the document:

- Now includes children up to 5 years of age (previously up to 3 years of age).

- Following the introduction of pneumococcal vaccination, the rates of occult bacteraemia have markedly decreased. Accordingly non-toxic febrile children older than 3 months of age who have no obvious source of infection are no longer screened for occult bacteraemia.

- Urinalysis has been introduced as a screening investigation for non-toxic febrile children older than 3 months of age who have no obvious source of infection.

- Increased emphasis has been placed upon the timely diagnosis of urinary tract infection, Meningococcal disease and Kawasaki disease.
Overview

Fever is one of the most common acute presentations in childhood. Many children will be only mildly unwell and will have a focus of infection identified on clinical examination.

Our aim is to detect those children with serious causes of fever such as meningitis, pneumonia and pyelonephritis without subjecting too many children to too many procedures or tests. This requires a combination of clinical judgement, specific investigations and serial observation.

Fever changes rapidly over time and a parent’s perception of the presence of fever in a child prior to presentation should not be discounted.

Key factors are:
- the child’s age
- presence of signs of toxicity
- presence of a focus of infection

When dealing with children suspected of having an infectious disease, it is essential that infection control measures be implemented to prevent cross contamination and spread.

Rationale for clinical approach

(1) Age

Neonates and young infants:
- May not have the characteristic signs of serious infection (temperature can be high or low).
- Localising features may be absent.
- Can deteriorate rapidly.
- May be infected with organisms from the birth canal.

Young infants with fever, especially those under three months of age, need rapid assessment and investigation, and admission to hospital. Consult a senior colleague about the extent of investigations (full blood count, cultures of blood, urine and CSF, chest x-ray) and the administration of antibiotics.

Older infants/toddlers:
- Localise infection better than neonates, but may still be pre verbal.
- Frequently exposed to infectious diseases in group childcare.
- Get viral infections as well as the ‘typical’ bacterial infections of pneumococcus, meningococcus and Hib (incidence of these infections significantly lessened by immunisation).
Assessment and initial management

Flowchart for child < 5 years old with fever (>38°C axillary)

- **Needs resuscitation?**
  - Yes: Get help, resuscitate. Investigate (FBC, B/C, Urine, CSF, CXR) Admit / transfer, IV antibiotics
  - No: Age < 3 months? (corrected for gestation)
    - Yes: Investigate (FBC, B/C, Urine, CSF, CXR) Admit / transfer, senior review / consult early. Investigate (FBC, B/C, Urine, CSF, CXR) IV antibiotics
    - No: Any sign of toxicity?
      - Yes: Toxicity
        - One or more of:
          - Alertness arousal or activity decreased
          - Breathing difficulties (tachypnoea, increased work of breathing)
          - Colour (pale or mottled), circulation (cool peripheries), or cry (weak, high pitched)
          - Decreased fluid intake or decreased urine output
      - No: Focus of infection?
        - Yes: Urinalysis. If positive, culture urine and commence antibiotics. If negative, review / consult next day if still febrile.
        - No: Investigate focus as appropriate and treat

- **Toxicity**
  - Alertness arousal or activity decreased
  - Breathing difficulties (tachypnoea, increased work of breathing)
  - Colour (pale or mottled), circulation (cool peripheries), or cry (weak, high pitched)
  - Decreased fluid intake or decreased urine output

Unimmunised children are at increased risk of serious bacterial infection

- **Axillary measurement of temperature is preferred in the 0-5 years age group.**
- **Oral and rectal measurements are not recommended because of safety concerns.**
- **Tympanic measurements may be inaccurate.**
- **When in doubt, ask for advice. No febrile child should be discharged from an Emergency Department without senior advice, particularly a child referred by a general practitioner, or a child representing with a febrile illness.**
- **At discharge the parent(s) should be educated on the detection and significance of toxicity, arrangements made for review, and a Fever Fact sheet and discharge summary provided.**
- **Err on the side of caution. If you are worried, admit / transfer the child.**
- **Only do a procedure or a test if it is going to contribute to a clinical decision. Use the flowchart to work out what tests you need. If in doubt about a child’s clinical condition consult with someone more experienced such as a paediatric specialist.**
  - If a specialist is not available, call NETS (the Newborn and paediatric Emergency Transport Service) on 1300 36 2500.
Older children:
- Usually verbalise and localise symptoms well.
- More tolerant to fluid loss – less likely to need IV rehydration.
- Can get ‘typical’ childhood organisms plus others such as mycoplasma and infectious mononucleosis.

(2) Toxicity: ABCD
Use this simple system to work out how sick a child appears to be:

‘A’ is for arousal, alertness or activity decreased

‘B’ is for breathing difficulties (tachypnoea, increased work of breathing)

‘C’ is for poor colour (pale or mottled), poor circulation (cold peripheries, increased capillary refill time) or cry (weak, high pitched)

‘D’ is for decreased fluid intake (less than half normal) and/or decreased urine output (fewer than four wet nappies a day)

The presence of any of these signs places the child at high risk of serious illness.

The presence of more than one sign increases the risk.

A ‘toxic’ child appears drowsy, lethargic or irritable, pale, mottled or tachycardic. Children with any of these signs must be seen urgently, investigated and treated as a priority.

The majority of children with signs of toxicity will receive antibiotic therapy.

The decision to administer antibiotics will be based upon age, degree of toxicity, height of fever, and height of white cell count and acute phase reactants.

(3) Focus of infection
Children with a definite focus of infection should only have investigations specific to that focus unless they are very young or toxic. For example, a mildly unwell child with definite acute otitis media does not need a urine culture, but a very unwell child who has acute otitis media needs a more thorough work-up as the child may have secondary bacteraemia, meningitis or an abscess.

Subjective features
Subjective features such as mild reddening of the throat or tympanic membranes should be interpreted with great caution especially in young children. Ask a more senior doctor to review the patient if the signs are mild or subjective.

Rash and fever
Not all rashes associated with fever are viral or ‘non-specific’. Meningococcal disease and Kawasaki disease are two important causes of rash which require timely diagnosis and therapy (see Box 1 and Box 2). If in any doubt ask a senior colleague for advice.
(4) Investigations

Perform an investigation only if the result is likely to alter management.

In urgent cases, such as a toxic child, do not wait for local anaesthetic to work. Get senior help immediately and get on with it.

Blood for culture should be taken whenever a blood count is performed on children with toxicity and/or a focus of infection.

White cell count and acute phase reactants can serve as a guide to the introduction of antibiotic therapy in children with toxicity and/or a focus of infection.

Chest x-ray is most useful if the child has signs of respiratory illness such as cough, tachypnoea, dullness or crackles. If there are no respiratory signs perform other investigations before the CXR.

Lumbar puncture should be considered in a young infant, toxic child, irritable child or a child with complex febrile convulsions, especially if the child is...
already on antibiotics. However, if the child is drowsy or requires resuscitation, resuscitation and antibiotics take precedence – do not delay. Please refer to Bacterial meningitis Clinical Practice Guideline.

**Urine culture** should be performed in all febrile children <3 months of age and all children who are toxic. A clean catch urine is appropriate however, timely collection is often difficult. Bag urine samples are inappropriate because of high contamination ratios. When it is urgent to get a urine specimen, a catheter urine sample is the recommended invasive technique.

Urine culture is essential prior to the commencement of antibiotics for suspected urinary tract infection.

For non-toxic, febrile children >3 months, dipstick urinalysis is an appropriate screening investigation, with urine culture being performed if the urinalysis is positive for leucocyte esterase or nitrites.

However, because of the difficulties in collecting satisfactory urine specimens in young children yet to be toilet trained, and because of concerns about possible renal damage associated with urinary tract infections during infancy, many practitioners will elect to simultaneously send urine for culture in these children, regardless of the urinalysis result.

**Antipyretics**

Antipyretics may provide comfort to a distressed child with fever. The presence of fever does not demand the use of antipyretics. There may be advantages to the child in not treating the fever.

<table>
<thead>
<tr>
<th><strong>Recommended doses</strong></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Paracetamol</strong></td>
<td>15mg/kg per dose given up to four-hourly up to a maximum of four doses each 24 hours. [ref Paracetamol Use PD2009_009, 26 Feb 2009]</td>
</tr>
<tr>
<td><strong>Ibuprofen</strong></td>
<td>(not recommended for children less than 6 months old). 10mg/kg per dose, given up to 6 hourly up to a maximum of four doses each 24 hours.</td>
</tr>
</tbody>
</table>

Alternating paracetamol and ibuprofen is theoretically unwise and not recommended. The response of fever to antipyretics is not of use in assessing the significance of an infection.

**Tepid sponging**

Tepid sponging and other physical methods of reducing temperature are not recommended and may be counterproductive. Unwrapping an overdressed child is appropriate.

**Follow-up**

- Children who are discharged home from an Emergency Department with fever should generally be followed up the following day, to assess progression of infection, response to treatment and results of investigations.

- Each facility will have in place its own system to facilitate this review and
these arrangements should be relayed to the parent(s) in writing.

- Although a child may be non-toxic when seen, no test can exclude the child becoming toxic and unwell later.

- Parents should be encouraged to look for toxicity every four to six hours, and to seek clinical review if the child becomes toxic or unwell.

- Clear communication from a doctor with empathy for the parents may enhance safety and improve the functioning of stressed families. A Fever fact sheet should be provided.

- The discharging Emergency Department doctor should write a note to the family doctor with the clinical diagnosis and a list of investigations performed.
What is fever?

- A fever is a rise in temperature, above normal, allowing for diurnal variation. Normal body temperature varies with time of day, but is generally less than 37.5°C centrally [Level III-2].

- Fever is commonly defined as a rectal temperature >38°C, which is approximately two standard deviations above the mean for infants under 3 months old (Herzog and Coyne 1993). In some circumstances, however, a lower temperature will be abnormal [Level IV].

- Children can have severe sepsis with no or minimal fever [Level IV].

How should temperatures be measured?

- Rectal temperatures are the ‘gold standard’ for measuring central body temperature [Level IV]. There are safety concerns (particularly in the very young) with its routine use as well as issues concerning lack of acceptability.

- Other methods of measurement such as axillary and tympanic membrane temperatures are less accurate and
less reliable than rectal temperatures [Level 1: Duce 1996, Craig et al 2000].

- In a study comparing axillary and rectal temperatures, a neonate's rectal temperature was on average 0.2°C higher than axillary, whereas the mean difference was 0.9°C for older children [Craig et al 2000, Level 111 - 2]. However, the confidence intervals were wide.

- Oral temperature measurement has not been systematically compared to rectal temperature. There are also concerns surrounding safety and acceptability.
Recommendation 1:
Axillary measurement of temperature is recommended for routine clinical use, but staff should be aware that axillary temperatures are up to 1°C lower than rectal temperatures. Rectal and oral temperatures are not recommended because of safety concerns and problems with acceptability. There is also a lack of data for oral temperatures. Tympanic temperatures are not recommended as they are unreliable.

Age of child
The younger the febrile infant, the greater is the incidence of a serious bacterial infection. For febrile neonates (0–4 weeks) the incidence is 12–32% [Level 111 - 2] (Neto 2004, Baker 1999, Kadish 2000).

For febrile infants aged 1–3 months, the risk of serious bacterial infection is somewhat lower but still significant (15–21% pre-pneumococcal immunisation) (Neto 2004, Baker 1999, Kadish et al 2000, Roberts et al 1977, Caspe et al 1983).

A variety of criteria eg Rochester, have been devised to attempt to identify a population of low risk infants aged 1–3 months who can be managed as outpatients (Dagan et al 1985, Dagan et al 1988). Estimations of the risk of an undetected serious bacterial infection in these low risk populations vary from 0.2% to 2% (Neto 2004, Baraff et al 1992, Klassen et al 1992).

Recommendation 2:
All febrile neonates should have a full septic workup and be admitted for parenteral antibiotics.

Infants aged 1–3 months will generally be managed in a similar fashion but there may be a place for outpatient management in carefully selected infants who are non-toxic, clinically stable over a period of observation, have reassuring pathology investigations and in whom close follow-up is assured.

Clinical assessment
The sensitivity of a ‘toxic appearance’ in detecting serious bacterial infection varied from 11% to 100% in different studies (Neto 2000) [Level I].

The most reliable infant observation scales were the Yale Observation Scale (McCarthy et al 1992) which examined quality of cry, reaction to parents, arousal, colour, hydration and social response, and the
Melbourne Study (Hewson et al 1990) which found the best predictors of serious bacterial infection to be feeding, breathing, hydration, activity, drowsiness and a history of being both pale and hot [Level II].

In older children with ‘occult bacteraemia’, the clinical appearance is very poorly predictive of the presence of bacteraemia, hence the term occult. Now much less common since the introduction of pneumococcal immunisation (Kuppermann et al 1998, Bulloch 2000) [Level I].

**Recommendation 3:**
Any child assessed as being ‘toxic’ should be admitted to hospital for investigation and, under most circumstances, administration of parenteral antibiotics.

**Meningococcal disease**
Invasive meningococcal group B disease continues to cause substantial morbidity and mortality. There is very commonly an early non-specific stage indistinguishable from a self-limiting viral illness.

Observational studies have associated leg pain, cold extremities and abnormal skin colour with developing invasive meningococcal disease (Thompson et al 2006).

Serial observation of febrile infants and children by experienced observers is an important strategy in the early detection of meningococcal disease (Thelen et al 2008).

**Kawasaki disease**
Kawasaki disease is an acute self-limited systemic vasculitis of unknown etiology which mainly affects infants and young children. Up to 25% of affected children, if untreated, develop coronary artery aneurysms.

The diagnosis is a clinical one, although there are often significant abnormalities of laboratory investigations. Timely diagnosis is important as the introduction of treatment, particularly with intravenous immunoglobulin, is efficacious in reducing symptoms and decreasing the formation of new coronary artery aneurysms (Newburger et al, 2004, Royle et al, 2005, Brogan et al, 2002, Oates-Whitehead, 2005).

**Urinary tract infection**
In febrile infants, history of a previous urinary tract infection, temperature higher than 40ºC and suprapubic tenderness are the findings most useful for identifying those with a urinary tract infection. Lack of circumcision among males, abdominal pain, back pain and lower urinary tract symptomatology also increase the likelihood of urinary infection (Nader et al 2007).

Urinary tract infection is the commonest serious bacterial infection in a febrile child with no clinically apparent focus of infection (Moyer 2004).

Recommendation 4:
The possibility of urinary tract infection needs to be considered in all febrile children who do not have an obvious source of infection. Although urinalysis is a useful screening investigation in these children, urine culture is essential prior to the commencement of antibiotics for suspected urinary tract infection.

Antipyretics

Ibuprofen is comparable as an antipyretic to paracetamol (Wilson et al 1991) [Level III–3].

Both the risks and benefits of paracetamol may have been exaggerated as parents are unable to guess from their child’s behaviour whether they received paracetamol or placebo (Kramer et al 1991) [Level II].


Recommendation 5:
The response to antipyretics should not be used as a diagnostic tool to try to differentiate bacterial from viral infection.
Appendix One – References

What is fever?

How should temperatures be measured?

Age of child

Clinical assessment


Meningococcal disease


Kawasaki disease


Urinary tract infection


Whiting P, Westwood M, Watt I, Cooper J, Kleijnen J. Rapid tests and urine sampling techniques for the diagnosis


Deville W LJM, Yzermans J C, Van Duijn NP, Bezemer PD, van der Windt D, Bouter L. The urine dipstick test useful to rule out infections. A meta-analysis of the accuracy. BMC Urology 2004: 4:1–14

Antipyretics


New References


Appendix Two – Resources

Fuller details may be necessary in practice, especially for the management of children with fever. Possible sources include:

NSW Health Department CIAP web site, Managing young children and infants with fever in Hospitals at: www.ciap.health.nsw.gov.au


Appendix Three – Parent information

A Fever fact sheet jointly developed by the John Hunter Children’s Hospital, Sydney Children’s Hospital and Children’s Hospital at Westmead is available at:


www.sch.edu.au/health/factsheets

www.chw.edu.au/parents/factsheets

 Disclaimer:
The fact sheet is for educational purposes only. Please consult with your doctor or other health professional to ensure this information is right for your child.
## Appendix Four – Working party members

<table>
<thead>
<tr>
<th>Name</th>
<th>Position</th>
<th>Affiliation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dr Des Mulcahy (Chair)</td>
<td>Paediatrician</td>
<td>Orange Base Hospital</td>
</tr>
<tr>
<td>Dr Shanika Attale</td>
<td>Paediatric Registrar</td>
<td>John Hunter Children’s Hospital</td>
</tr>
<tr>
<td>Ms Lucy Bates</td>
<td>Policy Officer</td>
<td>Northern Child Health Network</td>
</tr>
<tr>
<td>Ms Leanne Crittenden</td>
<td>Co-ordinator</td>
<td>Northern Child Health Network</td>
</tr>
<tr>
<td>Dr Michael Fasher</td>
<td>General Practitioner</td>
<td>Adjunct Associate Professor Western Clinical School</td>
</tr>
<tr>
<td>Dr Mark Birch</td>
<td>Infectious Diseases</td>
<td>John Hunter Children’s Hospital</td>
</tr>
<tr>
<td>Mr Audas Grant</td>
<td>CNC Emergency / Critical Care</td>
<td>Greater Southern (Albury)</td>
</tr>
<tr>
<td>Dr Brett Ireland</td>
<td>RMO</td>
<td>John Hunter Children’s Hospital</td>
</tr>
<tr>
<td>Prof David Isaacs</td>
<td>Immunologist</td>
<td>Children’s Hospital at Westmead</td>
</tr>
<tr>
<td>Mr Paul Kaye</td>
<td>After Hours Bed Manager / After Hours Nurse Manager</td>
<td>Greater Western Area Health Service, (Broken Hill)</td>
</tr>
<tr>
<td>Mr Martin Madejski</td>
<td>Transitional Nurse Practitioner</td>
<td>Emergency Department, Children’s Hospital at Westmead</td>
</tr>
<tr>
<td>Dr Matthew O’Meara</td>
<td>Paediatric Emergency Physician</td>
<td>Sydney Children’s Hospital</td>
</tr>
<tr>
<td>Dr Susan Piper</td>
<td>Medical Director</td>
<td>Paediatric Ambulatory Care Unit, Wyong</td>
</tr>
<tr>
<td>Dr Jo Rainbow</td>
<td>Paediatrician</td>
<td>Orange Base Hospital</td>
</tr>
<tr>
<td>Mr Thomas Ratoni</td>
<td>CNC Paediatrics</td>
<td>North Coast Area Health Service (Lismore)</td>
</tr>
<tr>
<td>Ms Rhonda Winskill</td>
<td>CNC Paediatrics</td>
<td>Northern Child Health Network (Maitland)</td>
</tr>
</tbody>
</table>