

# NATIONAL INSTITUTE FOR HEALTH AND CLINICAL EXCELLENCE

## Health and social care directorate

### Quality standards and indicators

#### Briefing paper

**Quality standard topic:** Urinary tract infection in children

**Output:** Prioritised quality improvement areas for development.

**Date of Quality Standards Advisory Committee meeting:** 22 November 2012

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# 1 Introduction

This briefing paper presents a structured overview of potential quality improvement areas for a NICE quality standard on urinary tract infection (UTI) in children.

It provides the Committee with a basis for discussion when prioritising quality improvement areas for developing the draft quality standard statements and measures. The draft quality standard will be the subject of public consultation.

## Key development source(s)

Unless otherwise stated, the key development source referenced in this briefing paper as follows:

[Urinary tract infection in children: diagnosis, treatment and long-term management](#). NICE clinical guideline 54 (2007). (Review decision date: May 2011; next review date: August 2013)

Where relevant, guideline recommendations from the key development source are presented alongside each of the suggested areas for quality improvement within the main body of the report.

# 2 Overview

## 2.1 *Focus of quality standard*

The focus of this quality standard is the care of infants and children from birth up to the age of 16 years with first or recurrent upper or lower tract UTI who are not already known to have underlying renal tract disease (specifically uropathy).

## 2.2 *Definition*

Urinary tract infection is a common bacterial infection causing illness in infants and children. A UTI is defined in NICE clinical guideline 54 by a combination of clinical features and the presence of bacteria in the urine (significant bacteriuria).

## 2.3 *Incidence and prevalence*

Urinary tract infection is a common bacterial infection. Around 1 in 10 girls and 1 in 30 boys will have had a UTI by the age of 16 years.

First occurrence of UTI is most common in infancy and affects boys most often in the first 3 months of life while in girls the peak incidence is after 6 months. Infants are

often systemically unwell and have acute pyelonephritis/upper urinary tract infection while older children more often have lower UTI and typical symptoms of cystitis.

The incidence of first UTI falls with age in both sexes but UTI is less common in boys than in girls after the first 6 months. Recurrent infections in boys are relatively uncommon whereas they are very common in girls.

It is estimated that a general practice of 10 000 patients, six GPs and 100 births per year, each GP will expect to have:

- two consultations a year with a child younger than 5 years about UTI (girls : boys = 3 : 1)
- one consultation a year with a boy (aged 0–14 years) about UTI
- four consultations a year with a girl (aged 0–14 years) about UTI.

## **2.4 Management**

Between 1991 and 2007 guidance issued by Royal College of Physicians and adopted in UK practice<sup>1</sup> recommended that UTI should be considered in every child with a fever or urinary symptoms, that the diagnosis should be confirmed by culture of a urine sample and that, following treatment of the acute illness, all infants and children younger than 7 years should have specific renal imaging and receive prophylactic antibiotics until these investigations have been completed.

This model of management, which includes imaging, antibiotic prophylaxis (long term treatment with antibiotics) and prolonged follow-up, places a heavy burden on NHS primary and secondary care resources including local radiology departments. This approach is costly, is based on limited evidence of clinical effectiveness, it is unpleasant for children and distressing for their parents or carers.

The publication of NICE clinical guideline 54 in 2007 led to a significant change in practice. The aim of NICE clinical guideline 54 is to achieve a more consistent and rational approach to the management of a UTI in children, with less diagnostic imaging investigations and antibiotic prophylaxis. This involves GPs making a diagnosis and deciding on appropriate referrals on who to refer to secondary care for further investigations.

A severe infection can make a child extremely unwell and may sometimes have serious consequences, and even minor infections can be distressing. Prompt and accurate diagnosis of UTI is essential if this condition is to be managed correctly. The initial assessment will usually involve a combination of clinical assessment and diagnostic testing. Most children with a first-time UTI in the UK present to primary care or to an emergency department.

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<sup>1</sup> Guidelines for the management of acute urinary tract infection in childhood. Report of a Working Group of the Research Unit, Royal College of Physicians. Published in 1991.

It should also be noted that NICE also issued more general guidance on the management of the febrile child in 2007 that offers a “traffic light” system for assessing possible illness severity and highlights the need for a UTI to be considered as the possible cause of a febrile illness. See NICE clinical guideline 47 on feverish illness in children for recommendations on urinary tract infection in children with fever. *Note: a NICE quality standard on feverish illness in children will be developed as part of the quality standards topic library.*

See appendices 1, 2 and 3 for patient pathway, algorithms and key priority for implementation recommendations from NICE clinical guideline 54.

### 3 Summary of suggestions

#### 3.1 Responses

In total six stakeholders submitted suggestions for quality improvement as part of the 2-week engagement exercise (19/09/12 – 03/10/12). Suggestions were also provided by specialist committee members.

**Table 1 Summary of suggested quality improvement areas**

Stakeholders were asked to suggest up to 5 areas for quality improvement. These have been merged and summarised in the table below for further consideration by the Committee (incorporating stakeholder and specialist committee member suggestions).

The full detail of the suggestions is provided in appendix 4 for information.

| <b>Suggested area for improvement</b>   | <b>Stakeholder (see table 2 for abbreviations)</b> |
|---|--|
| <u>Education and awareness</u> <ul style="list-style-type: none"> <li>Improved awareness amongst the public and first-line health care information providers and educators about UTI in children.</li> </ul>  | NUTH, NHS NorthEast, RCN                           |
| <u>Diagnosis</u> <ul style="list-style-type: none"> <li>Accurate diagnosis of UTI including accurate clinical differentiation of upper/lower UTI.</li> </ul>  | HPA, NHS NorthEast, NUTH, RCN, RCPH, SCM           |
| <u>Diagnosis</u> <ul style="list-style-type: none"> <li>Identification of risk factors for renal disease.</li> </ul>  | RCPH   |
| <u>Diagnosis</u> <ul style="list-style-type: none"> <li>Early treatment where high likelihood of UTI with adjustment depending on culture result.</li> </ul>  | NUTH, NHS NorthEast, RCN                           |
| <u>Acute management – immediate referral to paediatric specialist</u> <ul style="list-style-type: none"> <li>Referral of young babies with possible UTI to a secondary centre for confirmation of diagnosis and rapid treatment.</li> </ul>   | NUTH, NHS NorthEast, RCN,                          |
| <u>Acute Management – individualised care plans</u> <ul style="list-style-type: none"> <li>Individualised care plans for rapid access to diagnosis, treatment and management of further UTIs for children identified with a high risk of identifying abnormality i.e. children with scarring, family history of scarring or children with recurrent UTI's or other existing renal or functional anomalies.</li> </ul> | NHS NorthEast<br>NUTH<br>RCN                       |
| <u>Standards for imaging</u> <ul style="list-style-type: none"> <li>Standards for imaging in UTI.</li> </ul>  | BSPR/RCR   |

| <b>Suggested area for improvement</b>   | <b>Stakeholder (see table 2 for abbreviations)</b> |
|---|--|
| <u>Laboratory reporting standards</u> <ul style="list-style-type: none"> <li>Standards for definition of UTI in the laboratory</li> </ul> | SMC  |

### Additional areas

Additional suggestions provided in appendix 4 but not included in the above table are noted below. General statements cannot be developed so these need further discussion by the Committee to determine focus and suitability.

- Monitoring of vital signs. (RCN)
- Prompt investigations as per NICE guideline 54. (RCPH)
- Improvement of the use of antibiotic prophylaxis in children who have had a UTI. (SCM)
- Standards for definition of UTI in the laboratory with variable reporting between 105 colony forming units (CFU)/ml or 104 CFU/ml or 103 CFU/ml if pure culture. (HPA)

### **Table 2 Stakeholder details (abbreviations)**

The details of stakeholder organisations who submitted suggestions are provided in the table below.

| Abbreviation   | Full name  |
|----------------|--|
| BSPR/RCR       | British Society of Paediatric Radiology supported by The Royal College of Radiologists   |
| HPA            | Health Protection Agency Primary Care Unit - Microbiology Department   |
| NHS North East | Child Health and Maternity - UTI children sub-group of the Northern Child Health Clinical Network                                |
| NUTH           | The Newcastle Upon Tyne Hospitals NHS Foundation Trust, The Great North Children's Hospital, Department of Paediatric Nephrology |
| RCN            | Royal College of Nursing   |
| RCPH           | Royal College of Paediatrics and Child Health  |
| SCM            | Specialist Committee Member(s)   |

## **4 Suggested improvement area: education and awareness**

### **4.1 Summary of suggestions**

Stakeholders highlighted the need for improved awareness of signs and symptoms and signs of UTI, especially in very young children, amongst parents, the public and first-line health care information providers.

The importance of awareness was highlighted as crucial to reducing the risk associated with childhood urinary tract infections and ensuring that parents and carers act quickly and appropriately when their child is unwell by seeking help.

### **4.2 Selected recommendations from development source**

#### **General awareness of UTI**

NICE clinical guideline 54 (the key development source) does not provide specific recommendations to cover general awareness in the population.

*\*\*NICE clinical guideline 47 on feverish illness in children provides recommendations on urinary tract infection in children with fever. Note: a NICE quality standard on feverish illness in children will be developed.*

#### NICE CG47 Recommendation 1.2.2.7

Urinary tract infection should be considered in any child younger than 3 months with fever.

#### NICE CG47 Recommendation 1.2.2.8

Urinary tract infection should be considered in a child aged 3 months and older with fever and one or more of the following:

- vomiting
- poor feeding
- lethargy
- irritability
- abdominal pain or tenderness
- urinary frequency or dysuria
- offensive urine or haematuria.

#### **Symptoms and signs**

NICE clinical guideline 54 provides recommendations on recognition of diverse presenting symptoms and signs in infants and children with UTI, especially in the

very youngest. These are covered in the suggested improvement area for 'diagnosis' (section 5).

### **Information and advice for children, young people and parents or carers**

NICE clinical guideline 54 provides recommendations on advice that should be given to children and their parents/carers when they have had a UTI.

These are presented below to inform the Commission discussion.

#### NICE CG 54 - Recommendation 1.6.1.1

Healthcare professionals should ensure that when a child or young person has been identified as having a suspected UTI, they and their parents or carers as appropriate are given information about the need for treatment, the importance of completing any course of treatment and advice about prevention and possible long-term management.

#### NICE CG 54 - Recommendation 1.6.1.2

Healthcare professionals should ensure that children and young people, and their parents or carers as appropriate, are aware of the possibility of a UTI recurring and understand the need for vigilance and to seek prompt treatment from a healthcare professional for any suspected re-infection.

#### NICE CG 54 - Recommendation 1.6.1.3

Healthcare professionals should offer children and young people and/or their parents or carers appropriate advice and information on:

- prompt recognition of symptoms
- urine collection, storage and testing
- appropriate treatment options
- prevention
- the nature of and reason for any urinary tract investigation
- prognosis
- reasons and arrangements for long-term management if required.

### **4.3 Current UK practice**

The suggested improvement area for improvements in education and awareness is based on stakeholders' knowledge and experience. Stakeholders drew attention to existence of under diagnosis of UTI in some areas, although no published studies were provided.



## **5 Suggested improvement area: diagnosis**

### **5.1 Summary of suggestions**

Most of the stakeholders highlighted the importance of 'accurate and prompt diagnosis of UTI' to underpin appropriate further management.

It was noted that many children are often provided with 'possible' UTI without a conclusive diagnosis, resulting in some children being referred to a secondary care specialist centre and unnecessary tests.

The following key areas for quality improvement relating to diagnosis were highlighted:

- Symptoms and signs. In particular, stakeholders highlighted the importance of including children with unexplained fever who may not be adequately assessed for the presence of UTI and urine testing within 24 hours.
- Urine collection. Stakeholders highlighted the importance of collection of an uncontaminated urine sample.
- Speed of transportation of samples to laboratory and importance of urine preservation (if urine cannot be cultured within 4 hours of collection).
- Urine testing. In particular, as part of the urine testing strategies, stakeholders highlighted importance of prompt antibiotic treatment where there is high likelihood of UTI with subsequent adjustments of treatment according to culture result.
- History and examination on confirmed UTI, specifically the identification and recording of risk factors for renal disease.
- Clinical differentiation between upper and lower UTI.

### **5.2 Selected recommendations from development source**

Recommendations from the development source relating to the suggested improvement areas have been provisionally selected and are presented below to inform the Committee in their discussions.

#### **Symptoms and signs**

##### NICE CG 54 Recommendation 1.1.1.1 (KPI)

Infants and children presenting with unexplained fever of 38°C or higher should have a urine sample tested after 24 hours at the latest.

### NICE CG 54 Recommendation 1.1.1.3 (KPI)

Infants and children with symptoms and signs suggestive of urinary tract infection (UTI) should have a urine sample tested for infection. Table 1 is a guide to the symptoms and signs that infants and children present with.

## **Urine collection**

### NICE CG 54 Recommendation 1.1.3.1 (KPI)

A clean catch urine sample is the recommended method for urine collection. If a clean catch urine sample is unobtainable:

- Other non-invasive methods such as urine collection pads should be used. It is important to follow the manufacturer's instructions when using urine collection pads. Cotton wool balls, gauze and sanitary towels should not be used to collect urine in infants and children.
- When it is not possible or practical to collect urine by noninvasive methods, catheter samples or suprapubic aspiration (SPA) should be used.
- Before SPA is attempted, ultrasound guidance should be used to demonstrate the presence of urine in the bladder.

## **Urine preservation**

### NICE CG 54 Recommendation 1.1.4.1

If urine is to be cultured but cannot be cultured within 4 hours of collection, the sample should be refrigerated or preserved with boric acid immediately.

## **Urine testing**

### NICE CG 54 Recommendation 1.1.5.1 (KPI)

The urine-testing strategies shown in tables 2–5 are recommended.<sup>2</sup>

## **History and examination on confirmed UTI**

### NICE CG 54 Recommendation 1.1.7.1 (KPI)

The following risk factors for UTI and serious underlying pathology should be recorded:

- poor urine flow
- history suggesting previous UTI or confirmed previous UTI
- recurrent fever of uncertain origin

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<sup>2</sup> Assess the risk of serious illness in line with [Feverish illness in children](#) (NICE clinical guideline 47) to ensure appropriate urine tests and interpretation, both of which depend on the child's age and risk of serious illness.

- antenatally-diagnosed renal abnormality
- family history of vesicoureteric reflux (VUR) or renal disease
- constipation
- dysfunctional voiding
- enlarged bladder
- abdominal mass
- evidence of spinal lesion
- poor growth
- high blood pressure.

### **Clinical differentiation between acute pyelonephritis/upper urinary tract infection and cystitis/lower urinary tract infection**

#### NICE CG 54 Recommendation 1.1.8.1

Infants and children who have bacteriuria and fever of 38°C or higher should be considered to have acute pyelonephritis/upper urinary tract infection. Infants and children presenting with fever lower than 38°C with loin pain/tenderness and bacteriuria should also be considered to have acute pyelonephritis/upper urinary tract infection. All other infants and children who have bacteriuria but no systemic symptoms or signs should be considered to have cystitis/lower urinary tract infection.

### **5.3 Current UK practice**

There is currently a limited amount of published current practice information to fully assess baseline practice relating to diagnosis of UTI in children.

No published reports were highlighted by respondents, so these suggested areas for quality improvement are based on stakeholders' knowledge and experience. However, stakeholders drew attention to variations in respect to accurate and prompt diagnosis.

The NICE Guideline Development Group, based on their expert opinion, noted wide variation in practice in diagnosis of UTI in children, both in the choice and combination of tests, in the type of dipstick test used and in how laboratories perform microscopy and culture.

An online survey<sup>3</sup> of 186 paediatric trainees in a region assessed awareness of NICE clinical guideline 54. In respect to diagnosis, 99% reported that they diagnosed UTI in children based on positive dipstick testing and 80% reported they used clean catch as the primary collection method. It is noted that this is a limited study of small sample size.

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<sup>3</sup> Bhojani S et al. (2010) Compliance with NICE guidelines for urinary tract infections: A survey among the paediatric trainees of the London Deanery. Archives of Disease in Childhood. 95:12

## **6 Suggested improvement area: acute management**

### **6.1 Summary of suggestions**

The suggested key area for quality improvement is the 'referral of young babies with possible UTI to a secondary centre for confirmation of diagnosis and rapid treatment'.

Stakeholders highlighted this is an important quality marker, emphasising that youngest children are at increased likelihood of underlying renal pathology and may rapidly deteriorate with sepsis.

### **6.2 Selected recommendations from development source**

Recommendations from the development source relating to the suggested improvement area have been provisionally selected and are presented below to inform the Committee in their discussions.

#### NICE CG 54 Recommendation 1.2.1.2 (KPI)

Infants younger than 3 months with a possible UTI should be referred immediately to the care of a paediatric specialist. Treatment should be with parenteral antibiotics in line with [Feverish illness in children](#) (NICE clinical guideline 47).

### **6.3 Current UK practice**

Based on their knowledge and experience, stakeholders drew attention to variation in local arrangements for referral of infants to paediatric specialists.

## **7 Suggested improvement area: individualised care plans**

### **7.1 *Summary of suggestions***

The suggested key area for quality improvement is 'individualised care plans to support rapid access to diagnosis, treatment and management of further UTIs for children identified with a high risk of identifying abnormality i.e. children with scarring, family history of scarring or children with recurrent UTIs or other existing renal anomalies'.

This was suggested as an important area because children with underlying renal anatomical or functional abnormality identified after presenting with a UTI are more likely to have recurrence of UTI and have poorer outcomes.

### **7.2 *Selected recommendations from development source***

NICE clinical guideline 54 does not provide specific recommendations relating to the use of care plans.

No recommendations are presented.

Further discussion is required to determine focus and suitability for this suggested area within the draft quality standard.

### **7.3 *Current UK practice***

The suggested improvement area is based on stakeholders' knowledge and experience.

## **8 Suggested improvement area: standards for imaging**

### **8.1 *Summary of suggestions***

One stakeholder highlighted the importance of standards for imaging in UTI in children as a quality improvement area for consideration by the Committee.

It was highlighted that quality of imaging, if performed without appropriate level of expertise, affects diagnosis and management and can result in poor outcomes including adverse psychological outcomes.

To support the rationale for improvement, it is suggested that the level of expertise may have fallen as a result of reduced investigations following a change in current practice and implementation of the NICE guidance.

### **8.2 *Selected recommendations from development source***

Imaging is a key feature of NICE clinical guideline 54. The NICE Guideline Development Group reviewed and made recommendations on which tests are to be performed but not how they are to be carried out.

No recommendations are presented.

Further discussion is required to determine focus and suitability for this suggested area within the draft quality standard.

### **8.3 *Current UK practice***

The suggested improvement areas are based on stakeholders' knowledge and experience. No published reports were identified.

## 9 Suggested improvement area: laboratory reporting standards

### 9.1 Summary of suggestion

The suggested area for quality improvement is 'reporting of *E. Coli* coliforms which should be differentiated from non-*E. Coli* coliforms'.

This is highlighted as important for overall quality because of the need to ensure reporting of organisms in a way that will identify atypical UTIs. The identification of atypical UTI (non *E Coli*) is important for ensuring infants and children with atypical UTI have ultrasound of the urinary tract during the acute infection to identify structural abnormalities of the urinary tract such as obstruction.

### 9.2 Selected recommendations from development source

The identification of non-*E. Coli* organisms is part of the definition of atypical UTI as set out in NICE clinical guideline 54 and which underpin a number of recommendations relating to acute management.

NICE clinical guideline 54 provides the following definitions, including infection with non-*E. coli* organisms.

#### Box 1 Definitions of atypical and recurrent UTI

##### Atypical UTI includes:

- seriously ill (for more information refer to [Feverish illness in children](#) [NICE clinical guideline 47])
- poor urine flow
- abdominal or bladder mass
- raised creatinine
- septicaemia
- failure to respond to treatment with suitable antibiotics within 48 hours
- **infection with non-*E. coli* organisms.**

##### Recurrent UTI:

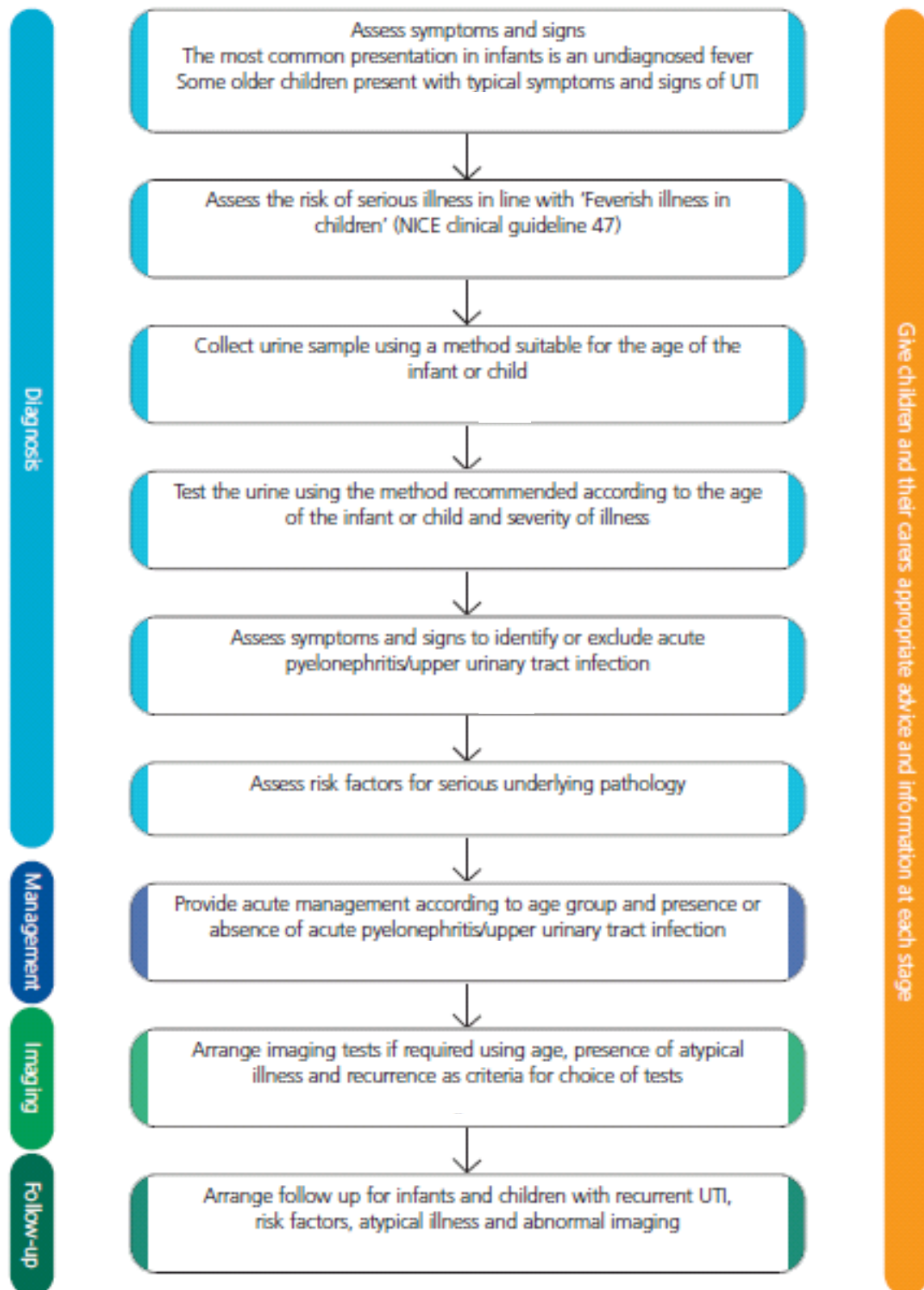
- two or more episodes of UTI with acute pyelonephritis/upper urinary tract infection, or
- one episode of UTI with acute pyelonephritis/upper urinary tract infection plus one or more episode of UTI with cystitis/lower urinary tract infection, or
- three or more episodes of UTI with cystitis/lower urinary tract infection.

### **9.3**      ***Current UK practice***

These suggested improvement areas are based on the respondents' knowledge and experience, citing that historically there is variable implementation of this criterion (E Coli vs non E Coli differentiation).



## Appendix 1 Pathway diagram (CG54)

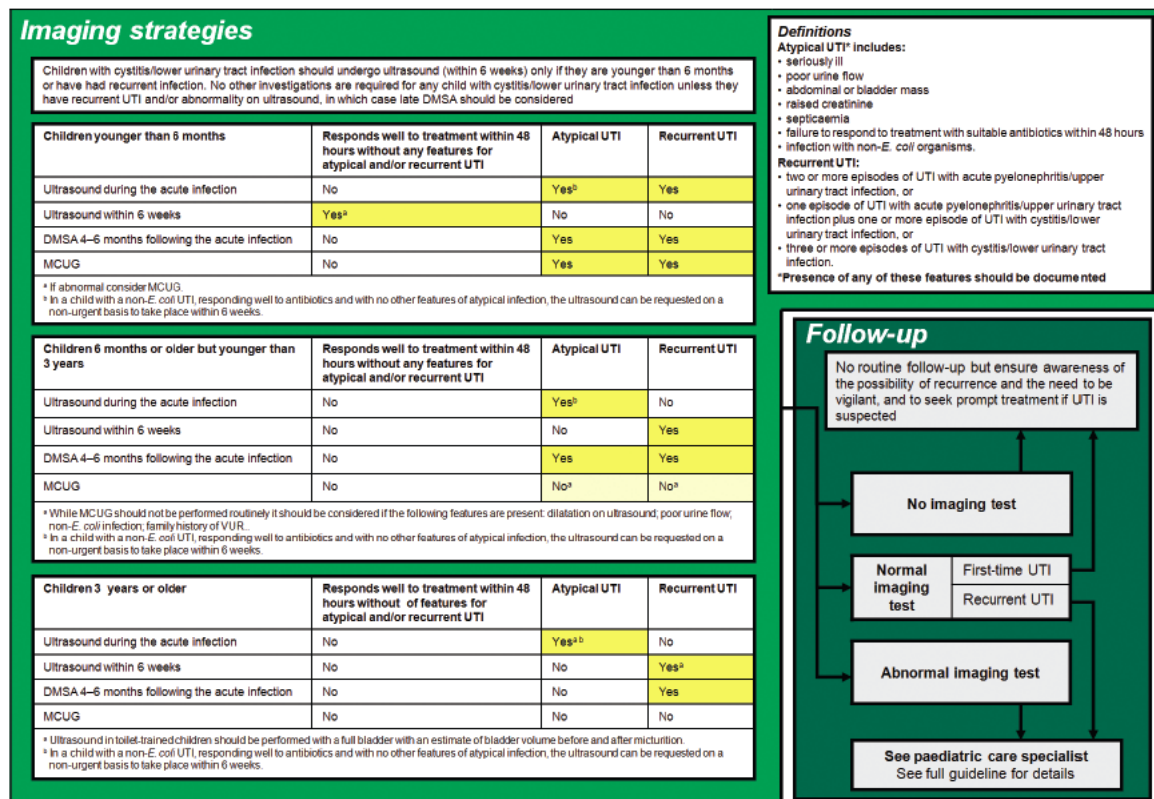
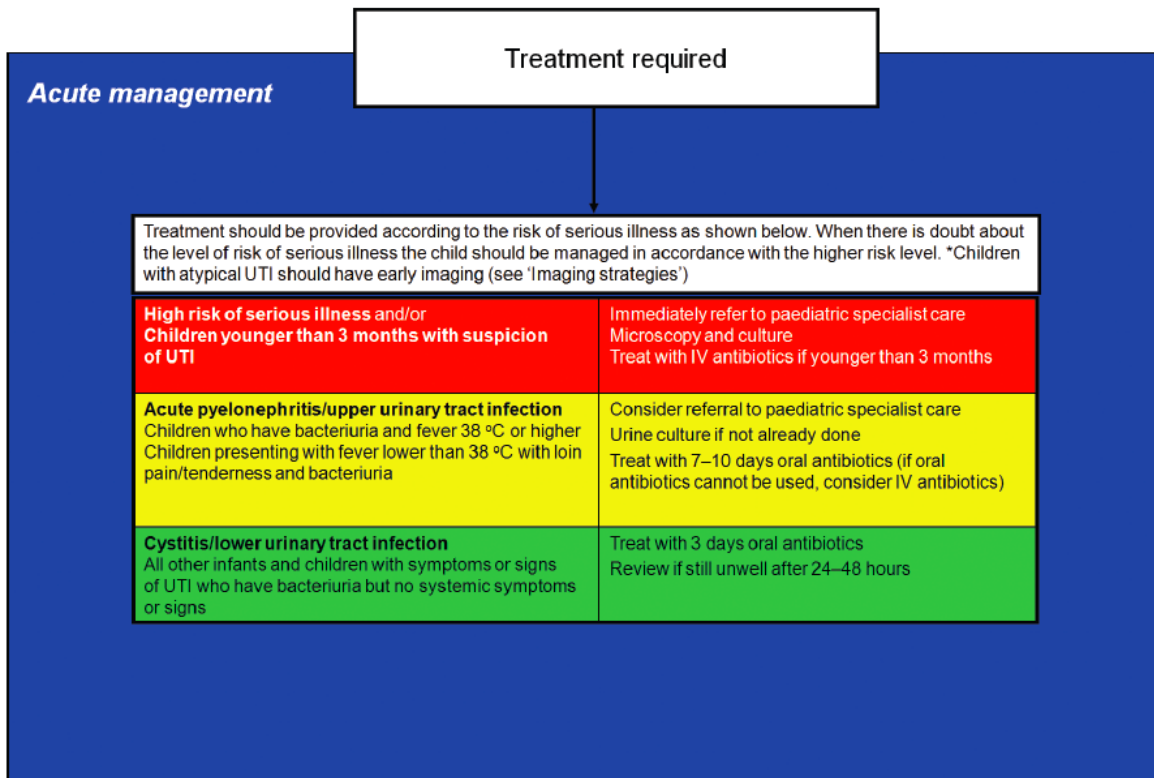


## Appendix 2 Algorithm from full clinical guideline (CG54)

The following algorithm is taken from the full clinical guideline 'Diagnosis, treatment and long-term management of urinary tract infection in infants and children'.

| Appropriate information and advice must be provided at each stage  |   |   |  |                |                |  |   |                                   |   |  |  |  |               |  |                |  |           |       |   |  |  |  |        |                      |   |   |  |  |
|--|---|---|--|----------------|----------------|--|---|-----------------------------------|---|--|--|--|---------------|--|----------------|--|-----------|-------|---|--|--|--|--------|----------------------|---|---|--|--|
| Infants younger than 3 months  | Infants and children 3 months or older but younger than 3 years | Children 3 years or older   |  |                |                |  |   |                                   |   |  |  |  |               |  |                |  |           |       |   |  |  |  |        |                      |   |   |  |  |
| <p><b>Suspecting UTI</b></p> <p style="text-align: center;"><b>Symptoms and signs suggesting UTI</b><br/>The presence of risk factors for UTI with serious underlying pathology should be recorded</p> <table border="1" style="width: 100%;"> <tr> <td colspan="2" style="text-align: center;">Most common ←</td> <td colspan="2" style="text-align: center;">→ Least common</td> </tr> <tr> <td style="width: 25%;">Fever<br/>Vomiting<br/>Lethargy<br/>Irritability</td> <td style="width: 25%;">Poor feeding<br/>Failure to thrive</td> <td style="width: 25%;">Abdominal pain<br/>Jaundice<br/>Haematuria<br/>Offensive urine</td> <td style="width: 25%;"></td> </tr> </table> <table border="1" style="width: 100%; margin-top: 10px;"> <tr> <td colspan="2"></td> <td colspan="2" style="text-align: center;">Most common ←</td> <td colspan="2" style="text-align: center;">→ Least common</td> </tr> <tr> <td style="width: 16.6%;">Preverbal</td> <td style="width: 16.6%;">Fever</td> <td style="width: 16.6%;">Abdominal pain<br/>Loin tenderness<br/>Vomiting<br/>Poor feeding</td> <td style="width: 16.6%;">Lethargy<br/>Irritability<br/>Haematuria<br/>Offensive urine<br/>Failure to thrive</td> <td style="width: 16.6%;"></td> <td style="width: 16.6%;"></td> </tr> <tr> <td>Verbal</td> <td>Frequency<br/>Dysuria</td> <td>Dysfunctional voiding<br/>Changes to continence<br/>Abdominal pain<br/>Loin tenderness</td> <td>Fever<br/>Malaise<br/>Vomiting<br/>Haematuria<br/>Offensive urine<br/>Cloudy urine</td> <td></td> <td></td> </tr> </table> <p style="text-align: center;">Assess the risk of serious illness using the traffic light system described in the <i>Feverish Illness in Children</i> guideline</p> |   |   | Most common ←  |                | → Least common |  | Fever<br>Vomiting<br>Lethargy<br>Irritability | Poor feeding<br>Failure to thrive | Abdominal pain<br>Jaundice<br>Haematuria<br>Offensive urine |  |  |  | Most common ← |  | → Least common |  | Preverbal | Fever | Abdominal pain<br>Loin tenderness<br>Vomiting<br>Poor feeding | Lethargy<br>Irritability<br>Haematuria<br>Offensive urine<br>Failure to thrive |  |  | Verbal | Frequency<br>Dysuria | Dysfunctional voiding<br>Changes to continence<br>Abdominal pain<br>Loin tenderness | Fever<br>Malaise<br>Vomiting<br>Haematuria<br>Offensive urine<br>Cloudy urine |  |  |
| Most common ←  |   | → Least common  |  |                |                |  |   |                                   |   |  |  |  |               |  |                |  |           |       |   |  |  |  |        |                      |   |   |  |  |
| Fever<br>Vomiting<br>Lethargy<br>Irritability  | Poor feeding<br>Failure to thrive                               | Abdominal pain<br>Jaundice<br>Haematuria<br>Offensive urine                         |  |                |                |  |   |                                   |   |  |  |  |               |  |                |  |           |       |   |  |  |  |        |                      |   |   |  |  |
|  |   | Most common ←   |  | → Least common |                |  |   |                                   |   |  |  |  |               |  |                |  |           |       |   |  |  |  |        |                      |   |   |  |  |
| Preverbal  | Fever   | Abdominal pain<br>Loin tenderness<br>Vomiting<br>Poor feeding                       | Lethargy<br>Irritability<br>Haematuria<br>Offensive urine<br>Failure to thrive |                |                |  |   |                                   |   |  |  |  |               |  |                |  |           |       |   |  |  |  |        |                      |   |   |  |  |
| Verbal   | Frequency<br>Dysuria  | Dysfunctional voiding<br>Changes to continence<br>Abdominal pain<br>Loin tenderness | Fever<br>Malaise<br>Vomiting<br>Haematuria<br>Offensive urine<br>Cloudy urine  |                |                |  |   |                                   |   |  |  |  |               |  |                |  |           |       |   |  |  |  |        |                      |   |   |  |  |
| <p><b>Collecting urine</b></p> <p style="text-align: center;">Urine should be tested in infants and children who have symptoms suggesting UTI described above. Those with unexplained fever of 38 °C or higher should have a urine sample tested after 24 hours at the latest</p> <p style="text-align: center;">Collect urine using a clean catch sample<br/>if not possible see guideline for details</p>  |   |   |  |                |                |  |   |                                   |   |  |  |  |               |  |                |  |           |       |   |  |  |  |        |                      |   |   |  |  |

| Testing urine   | Infants and children 3 months or older but younger than 3 years   | Children 3 years or older  |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |
|---|---|--|-----------------|--|-------------------------------|------------------------------|---|--------------------------------------|--|-----------------------------|---|---|--|---|--|--|--|---|--|---|
| <p>Infants younger than 3 months</p> <ul style="list-style-type: none"> <li>• Refer to paediatric specialist care</li> <li>• Urine sample for urgent microscopy and culture</li> <li>• Manage in line with 'Feverish illness in children' (NICE clinical guideline 47)</li> </ul>   | <p style="text-align: center;"><b>Use urgent microscopy and culture to diagnose UTI</b></p> <table border="1" style="width: 100%;"> <tr> <td style="width: 15%;">Specific urinary symptoms</td> <td style="width: 10%;"></td> <td style="width: 75%;">Urine sample for urgent microscopy and culture<br/>Start antibiotic treatment<br/>If urgent microscopy is not available, send a urine sample for microscopy and culture, and start antibiotic treatment.</td> </tr> <tr> <td rowspan="3">Non-specific urinary symptoms</td> <td style="background-color: #ff0000; color: white; text-align: center;">High risk of serious illness</td> <td>Urgent referral to paediatric specialist care<br/>Urine sample for urgent microscopy and culture<br/>Manage in line with 'Feverish illness in children' (NICE clinical guideline 47).</td> </tr> <tr> <td style="background-color: #ffff00; text-align: center;">Intermediate risk of serious illness</td> <td>Consider urgent referral to a paediatric specialist as described in 'Feverish illness in children' (NICE clinical guideline 47)<br/>When specialist paediatric referral is not required:                             <ul style="list-style-type: none"> <li>• Urgent microscopy and culture should be arranged</li> <li>• Antibiotic treatment should be started if microscopy is positive</li> <li>• When urgent microscopy is not available, dipstick testing may be used as a substitute</li> <li>• The presence of nitrites suggests the possibility of infection and antibiotic treatment should be started</li> </ul>                             In all cases, a urine sample should be sent for microscopy and culture.                         </td> </tr> <tr> <td style="background-color: #008000; color: white; text-align: center;">Low risk of serious illness</td> <td>Urine sample for microscopy and culture<br/>Start antibiotic treatment if microscopy or culture is positive.</td> </tr> </table> | Specific urinary symptoms  |                 | Urine sample for urgent microscopy and culture<br>Start antibiotic treatment<br>If urgent microscopy is not available, send a urine sample for microscopy and culture, and start antibiotic treatment. | Non-specific urinary symptoms | High risk of serious illness | Urgent referral to paediatric specialist care<br>Urine sample for urgent microscopy and culture<br>Manage in line with 'Feverish illness in children' (NICE clinical guideline 47). | Intermediate risk of serious illness | Consider urgent referral to a paediatric specialist as described in 'Feverish illness in children' (NICE clinical guideline 47)<br>When specialist paediatric referral is not required: <ul style="list-style-type: none"> <li>• Urgent microscopy and culture should be arranged</li> <li>• Antibiotic treatment should be started if microscopy is positive</li> <li>• When urgent microscopy is not available, dipstick testing may be used as a substitute</li> <li>• The presence of nitrites suggests the possibility of infection and antibiotic treatment should be started</li> </ul> In all cases, a urine sample should be sent for microscopy and culture. | Low risk of serious illness | Urine sample for microscopy and culture<br>Start antibiotic treatment if microscopy or culture is positive.   | <p style="text-align: center;"><b>Use dipstick test to diagnose UTI</b></p> <table border="1" style="width: 100%;"> <tr> <td style="width: 15%;">Both leucocyte esterase and nitrite positive</td> <td style="width: 85%;">Start antibiotic treatment for UTI<br/>If high or intermediate risk of serious illness or past history of UTI, send urine sample for culture</td> </tr> <tr> <td>Leucocyte esterase negative and nitrite positive</td> <td>Start antibiotic treatment if fresh sample was tested. Send urine sample for culture</td> </tr> <tr> <td>Leucocyte esterase positive and nitrite negative</td> <td>Send urine sample for microscopy and culture<br/>Only start antibiotic treatment for UTI if there is good clinical evidence of UTI<br/>Result may indicate infection elsewhere<br/>Treat depending on results of culture</td> </tr> <tr> <td>Both leucocyte esterase and nitrite negative</td> <td>Do not start treatment for UTI<br/>Explore other causes of illness<br/>Do not send urine sample for culture unless recommended in 'Indications for sending for culture'</td> </tr> </table> | Both leucocyte esterase and nitrite positive | Start antibiotic treatment for UTI<br>If high or intermediate risk of serious illness or past history of UTI, send urine sample for culture | Leucocyte esterase negative and nitrite positive | Start antibiotic treatment if fresh sample was tested. Send urine sample for culture | Leucocyte esterase positive and nitrite negative | Send urine sample for microscopy and culture<br>Only start antibiotic treatment for UTI if there is good clinical evidence of UTI<br>Result may indicate infection elsewhere<br>Treat depending on results of culture | Both leucocyte esterase and nitrite negative | Do not start treatment for UTI<br>Explore other causes of illness<br>Do not send urine sample for culture unless recommended in 'Indications for sending for culture' |
| Specific urinary symptoms   |   | Urine sample for urgent microscopy and culture<br>Start antibiotic treatment<br>If urgent microscopy is not available, send a urine sample for microscopy and culture, and start antibiotic treatment.   |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |
| Non-specific urinary symptoms   | High risk of serious illness  | Urgent referral to paediatric specialist care<br>Urine sample for urgent microscopy and culture<br>Manage in line with 'Feverish illness in children' (NICE clinical guideline 47).  |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |
|   | Intermediate risk of serious illness  | Consider urgent referral to a paediatric specialist as described in 'Feverish illness in children' (NICE clinical guideline 47)<br>When specialist paediatric referral is not required: <ul style="list-style-type: none"> <li>• Urgent microscopy and culture should be arranged</li> <li>• Antibiotic treatment should be started if microscopy is positive</li> <li>• When urgent microscopy is not available, dipstick testing may be used as a substitute</li> <li>• The presence of nitrites suggests the possibility of infection and antibiotic treatment should be started</li> </ul> In all cases, a urine sample should be sent for microscopy and culture. |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |
|   | Low risk of serious illness   | Urine sample for microscopy and culture<br>Start antibiotic treatment if microscopy or culture is positive.  |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |
| Both leucocyte esterase and nitrite positive  | Start antibiotic treatment for UTI<br>If high or intermediate risk of serious illness or past history of UTI, send urine sample for culture   |  |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |
| Leucocyte esterase negative and nitrite positive  | Start antibiotic treatment if fresh sample was tested. Send urine sample for culture  |  |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |
| Leucocyte esterase positive and nitrite negative  | Send urine sample for microscopy and culture<br>Only start antibiotic treatment for UTI if there is good clinical evidence of UTI<br>Result may indicate infection elsewhere<br>Treat depending on results of culture   |  |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |
| Both leucocyte esterase and nitrite negative  | Do not start treatment for UTI<br>Explore other causes of illness<br>Do not send urine sample for culture unless recommended in 'Indications for sending for culture'   |  |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |
| <p><b>Guidance on microscopy results</b></p> <table border="1" style="width: 100%;"> <tr> <td></td> <td style="width: 25%;">Pyuria positive</td> <td style="width: 25%;">Pyuria negative</td> </tr> <tr> <td>Bacteriuria positive</td> <td>Having UTI</td> <td>Having UTI</td> </tr> <tr> <td>Bacteriuria negative</td> <td>Start antibiotics if clinically UTI</td> <td>Not having UTI</td> </tr> </table> |   |  | Pyuria positive | Pyuria negative  | Bacteriuria positive          | Having UTI                   | Having UTI  | Bacteriuria negative                 | Start antibiotics if clinically UTI  | Not having UTI              | <p><b>Indications for sending for culture</b></p> <ul style="list-style-type: none"> <li>• acute pyelonephritis/upper urinary tract infection</li> <li>• a high to intermediate risk of serious illness</li> <li>• younger than 3 years</li> <li>• a single positive result for leucocyte esterase or nitrite</li> <li>• recurrent UTI</li> <li>• an infection that does not respond to treatment within 24–48 hours</li> <li>• when clinical symptoms and dipstick tests do not correlate.</li> </ul> <p style="text-align: center;"><b>Symptomatic infants or children with a positive urine culture should be treated.</b></p> |   |  |   |  |  |  |   |  |   |
|   | Pyuria positive   | Pyuria negative  |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |
| Bacteriuria positive  | Having UTI  | Having UTI   |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |
| Bacteriuria negative  | Start antibiotics if clinically UTI   | Not having UTI   |                 |  |                               |                              |   |                                      |  |                             |   |   |  |   |  |  |  |   |  |   |



## **Appendix 3 Key priorities for implementation recommendations (CG54)**

### **Symptoms and signs**

- Infants and children presenting with unexplained fever of 38°C or higher should have a urine sample tested after 24 hours at the latest.
- Infants and children with symptoms and signs suggestive of urinary tract infection (UTI) should have a urine sample tested for infection. [Table 1](#) is a guide to the symptoms and signs that infants and children present with.

### **Urine collection**

- A clean catch urine sample is the recommended method for urine collection. If a clean catch urine sample is unobtainable:
  - Other non-invasive methods such as urine collection pads should be used. It is important to follow the manufacturers' instructions when using urine collection pads. Cotton wool balls, gauze and sanitary towels should not be used to collect urine in infants and children.
  - When it is not possible or practical to collect urine by noninvasive methods, catheter samples or suprapubic aspiration (SPA) should be used.
  - Before SPA is attempted, ultrasound guidance should be used to demonstrate the presence of urine in the bladder.

### **Urine testing**

- The urine-testing strategies shown in [tables 2–5](#) are recommended.<sup>[1]</sup>

### **History and examination on confirmed UTI**

- The following risk factors for UTI and serious underlying pathology should be recorded:
  - poor urine flow
  - history suggesting previous UTI or confirmed previous UTI
  - recurrent fever of uncertain origin
  - antenatally-diagnosed renal abnormality

- family history of vesicoureteric reflux (VUR) or renal disease
- constipation
- dysfunctional voiding
- enlarged bladder
- abdominal mass
- evidence of spinal lesion
- poor growth
- high blood pressure.

### **Acute management**

- Infants younger than 3 months with a possible UTI should be referred immediately to the care of a paediatric specialist. Treatment should be with parenteral antibiotics in line with [Feverish illness in children](#) (NICE clinical guideline 47).
- For infants and children 3 months or older with acute pyelonephritis/upper urinary tract infection:
  - consider referral to a paediatric specialist
  - treat with oral antibiotics for 7–10 days. The use of an oral antibiotic with low resistance patterns is recommended, for example cephalosporin or co-amoxiclav.
  - if oral antibiotics cannot be used, treat with an intravenous (IV) antibiotic agent such as cefotaxime or ceftriaxone for 2–4 days followed by oral antibiotics for a total duration of 10 days.
- For infants and children 3 months or older with cystitis/lower urinary tract infection:
  - treat with oral antibiotics for 3 days. The choice of antibiotics should be directed by locally developed multidisciplinary guidance. Trimethoprim, nitrofurantoin, cephalosporin or amoxicillin may be suitable
  - the parents or carers should be advised to bring the infant or child for reassessment if the infant or child is still unwell after 24–48 hours. If an alternative diagnosis is not made, a urine sample should be sent for culture to identify the presence of bacteria and determine antibiotic sensitivity if urine culture has not already been carried out.

### **Antibiotic prophylaxis**

- Antibiotic prophylaxis should not be routinely recommended in infants and children following first-time UTI.

### **Imaging tests**

- Infants and children who have had a UTI should be imaged as outlined in [tables 6, 7 and 8](#).

**NICE clinical guideline 54 tables**

**Table 1 Presenting symptoms and signs in infants and children with UTI**

| <b>Age group</b>                     |  | <b>Symptoms and signs</b><br>Most common -----> Least common |   |   |
|--------------------------------------|--|--|---|---|
| <b>Infants younger than 3 months</b> |  | Fever<br>Vomiting<br>Lethargy<br>Irritability                | Poor feeding<br>Failure to thrive   | Abdominal pain<br>Jaundice<br>Haematuria<br>Offensive urine                   |
|                                      | <b>Infants and children, 3 months or older</b> | <b>Preverbal</b>   | Fever   | Abdominal pain<br>Loin tenderness<br>Vomiting<br>Poor feeding                 |
| <b>Verbal</b>                        |  | Frequency<br>Dysuria   | Dysfunctional voiding<br>Changes to continence<br>Abdominal pain<br>Loin tenderness | Fever<br>Malaise<br>Vomiting<br>Haematuria<br>Offensive urine<br>Cloudy urine |

**Table 2 Urine-testing strategy for infants younger than 3 months**

All infants younger than 3 months with suspected UTI (see table 1) should be referred to paediatric specialist care and a urine sample should be sent for urgent microscopy and culture. These infants should be managed in accordance with the recommendations for this age group in [Feverish illness in children](#) (NICE clinical guideline 47).

**Table 3 Urine-testing strategies for infants and children 3 months or older but younger than 3 years**

|  |  |
|--|--|
| <p>Urgent microscopy and culture is the preferred method for diagnosing UTI in this age group; this should be used where possible.</p> |  |
| <p><b>If the infant or child has specific urinary symptoms</b></p>   | <p>Urgent microscopy and culture should be arranged and antibiotic treatment should be started.</p> <p>When urgent microscopy is not available, a urine sample should be sent for microscopy and culture, and antibiotic treatment should be started.</p>  |
| <p><b>If the symptoms are non-specific to UTI</b></p>  | <ul style="list-style-type: none"> <li>• For an infant or child with a high risk of serious illness: the infant or child should be urgently referred to a paediatric specialist where a urine sample should be sent for urgent microscopy and culture. Such infants and children should be managed in line with <a href="#">Feverish illness in children</a> (NICE clinical guideline 47).</li> <li>• For an infant or child with an intermediate risk of serious illness: if the situation demands, the infant or child may be referred urgently to a paediatric specialist. For infants and children who do not require paediatric specialist referral, urgent microscopy and culture should be arranged. Antibiotic treatment should be started if microscopy is positive (see table 5). When urgent microscopy is not</li> </ul> |



|  |  |
|--|--|
|  | <p>available, dipstick testing may act as a substitute. The presence of nitrites suggests the possibility of infection and antibiotic treatment should be started (see table 4). In all cases, a urine sample should be sent for microscopy and culture.</p> <ul style="list-style-type: none"> <li>• For an infant or child with a low risk of serious illness: microscopy and culture should be arranged. Antibiotic treatment should only be started if microscopy or culture is positive.</li> </ul> |
|--|--|

**Table 4 Urine-testing strategies for children 3 years or older**

|   |   |
|---|---|
| <p>Dipstick testing for leukocyte esterase and nitrite is diagnostically as useful as microscopy and culture, and can safely be used.</p> |   |
| <p><b>If both leukocyte esterase and nitrite are positive</b></p>   | <p>The child should be regarded as having UTI and antibiotic treatment should be started. If a child has a high or intermediate risk of serious illness and/or a past history of previous UTI, a urine sample should be sent for culture.</p>   |
| <p><b>If leukocyte esterase is negative and nitrite is positive</b></p>   | <p>Antibiotic treatment should be started if the urine test was carried out on a fresh sample of urine. A urine sample should be sent for culture. Subsequent management will depend upon the result of urine culture.</p>  |
| <p><b>If leukocyte esterase is positive and nitrite is negative</b></p>   | <p>A urine sample should be sent for microscopy and culture. Antibiotic treatment for UTI should not be started unless there is good clinical evidence of UTI (for example, obvious urinary symptoms). Leukocyte esterase may be indicative of an infection outside the urinary tract which may need to be managed differently.</p> |
| <p><b>If both leukocyte esterase and nitrite are negative</b></p>   | <p>The child should not be regarded as having UTI. Antibiotic treatment for UTI should not be started, and a urine sample should not be sent for culture. Other causes of illness should be explored.</p>   |

**Table 5 Guidance on the interpretation of microscopy results**

| <b>Microscopy results</b>   | <b>Pyuria positive</b>                                   | <b>Pyuria negative</b>                                   |
|-----------------------------|--|--|
| <b>Bacteriuria positive</b> | The infant or child should be regarded as having UTI     | The infant or child should be regarded as having UTI     |
| <b>Bacteriuria negative</b> | Antibiotic treatment should be started if clinically UTI | The infant or child should be regarded as not having UTI |

## Appendix 4 Suggestions from stakeholder engagement exercise

| ID  | Stakeholder                   | Suggested key area for quality improvement  | Why is this important?  | Why is this a key area for quality improvement?   | Supporting information  |
|-----|-------------------------------|---|---|---|---|
| 001 | BSPR/RCR                      | <ul style="list-style-type: none"> <li>Standards for imaging in UTI</li> </ul>  | <p>Imaging is a key feature of the current NICE guidance, when they were introduced imaging pathways were standardised for the first time and practice was affected in many centres. The guidance specifies which tests are to be performed but not how they are to be carried out.</p> <p>The frequency of some investigations (MCUG, DMSA) has been drastically reduced which means that in many centres the level of expertise has fallen.</p> | <p>The quality of the imaging affects the diagnosis, management and outcome.</p> <p>Imaging tests are potentially traumatic for the child and their family and may have adverse psychological outcomes if performed inexpertly. There are new techniques which can vastly improve patient experience (e.g. entenox administration for catheterisation) which are not used in many centres.</p> <p>Some of the investigations involve ionising radiation with potentially harmful effects. The radiation exposure can be limited if guidance is issued and adhered to.</p> | None provided   |
| 002 | Newcastle Hospitals NHS Trust | <ul style="list-style-type: none"> <li>Improved awareness of symptoms &amp; signs of UTI especially in the very young amongst public and first-line health care info providers educators eg HVs, practice nurses, NHS direct type services</li> </ul> | <p>There is evidence that increased risk of scarring is associated with delay in treatment. Delays in diagnosis &amp; starting treatment are likely to be multi-factorial &amp; will require a multi faceted improvement.</p> <p>Some evidence of under diagnosis of UTI in some areas compared to expected incidence.</p>  | Improved parent education & information is associated with empowerment, better & efficient use of health care resources & higher satisfaction with health services  | <p>NICE CG 54 emphasises diverse symptoms &amp; signs of UTI especially in the very youngest; emphasises importance of prompt treatment &amp; ensuring appropriate advice &amp; info given to families.</p> <p>NICE CG 37 postnatal</p> |

| ID  | Stakeholder                   | Suggested key area for quality improvement   | Why is this important?   | Why is this a key area for quality improvement?   | Supporting information   |
|-----|-------------------------------|--|--|---|--|
|     |                               |  |  |   | <p>care<br/>Parents should be offered information &amp; advice to enable them to identify symptoms &amp; signs of common health problems seen in babies.</p> <p>Compare public awareness of meningitis after recent health education campaigns compared to awareness re UTI in babies.</p> |
| 003 | Newcastle Hospitals NHS Trust | <ul style="list-style-type: none"> <li>Accurate diagnosis of UTI &amp; accurate diagnosis of upper verses lower UTI</li> </ul> | <p>Accurate diagnosis underpins appropriate further management.</p> <p>Differentiation between upper &amp; lower UTI underpins CG54 management advice (eg. length of time of treatment with antibiotics, number of UTIs needed to have investigations performed)</p> | <p>Despite emphasis on diagnosis in NICE CG54</p> <p>Many children still being labelled as 'possible' UTI without a conclusive diagnosis, resulting in some children being referred to secondary services with dysuria from other causes etc This may waste resources &amp; also may submit some children to unnecessary tests.</p> <p>Differentiation between upper &amp; lower UTI as advised in NICE CG 54 is difficult to do clinically &amp; outcomes need auditing.</p> | NICE CG54  |
| 004 | Newcastle                     | <ul style="list-style-type: none"> <li>Starting treatment early</li> </ul>   | Evidence of increased risk of  | Evidence of variation in practice which may   | CG54   |

| ID  | Stakeholder                      | Suggested key area for quality improvement   | Why is this important?   | Why is this a key area for quality improvement?  | Supporting information   |
|-----|----------------------------------|--|--|--|--|
|     | Hospitals<br>NHS Trust           | if high likelihood of UTI, sending urine for culture & adjusting treatment depending on culture result.  | scarring with delay in treatment   | contribute to delay in treatment.  |  |
| 005 | Newcastle Hospitals<br>NHS Trust | <ul style="list-style-type: none"> <li>Investigation plans should focus on those with highest likelihood of identifying abnormality eg those with recurrent UTI, the youngest. Those who are identified after 1st UTI or recurrence to have scarring or VUR or other anatomical or functional urinary tract problem, should have individualised care plan for rapid access to diagnosis &amp; treatment &amp; management of any further UTIs.</li> </ul> | <p>Evidence that those with underlying renal anatomical or functional abnormality identified after presenting with a UTI are more likely to have recurrence of UTI &amp; poorer outcome.</p> <p>Access to prompt diagnosis of UTI &amp; rapid treatment important &amp; needs to be individualised to circumstances of the child, family &amp; local services.</p> | <p>Is there any point of detecting significant underlying renal abnormality if no active management started?</p> <p>Management may include treatments like antibiotics but also education &amp; advice about regular voiding, avoiding vulval irritants, drinking plenty &amp; managing constipation etc.</p> <p>Prevention of worsening of chronic kidney disease.</p> <p>Prevention of prolonged &amp; recurrent illness</p> | <p>NICE CG54</p> <p>NICE Quality standards for management of Chronic Kidney Disease.</p> |
| 006 | Newcastle Hospitals<br>NHS Trust | <ul style="list-style-type: none"> <li>Young babies with possible UTI refer to secondary centre for confirmation of diagnosis and rapid treatment.</li> </ul>  | Evidence youngest children most likely to have underlying renal pathology & get most unwell with sepsis.   | Local arrangements for referral vary from place to place. Delay in accessing secondary assessment has been implicated in some case reviews with adverse outcomes.  | CG54<br>CG47   |

| ID  | Stakeholder   | Suggested key area for quality improvement   | Why is this important?   | Why is this a key area for quality improvement?  | Supporting information   |
|-----|---------------|--|--|--|--|
|     |               |  |  | Age range needs careful consideration; CG54 advises <3m but older infants & children who are unwell should also be included –as advised in CG 47 Feverish illness in children.   |  |
| 007 | NHS NorthEast | <ul style="list-style-type: none"> <li>Public and first-line health care education &amp; awareness of symptoms &amp; signs of UTI especially in the very young.</li> </ul> | <p>Delays in diagnosis &amp; treatment are multi-factorial.</p> <p>Some evidence of under diagnosis of UTI in some areas compared to expected incidence.</p> <p>Evidence that increased risk of scarring associated with delay in treatment.</p> | Improved parent education & information is associated with empowerment, better use of health care resources & higher satisfaction with health services   | <p>NICE CG 37 postnatal care</p> <p>Parents should be offered information &amp; advice to enable them to identify symptoms &amp; signs of common health problems seen in babies.</p> <p>Compare public awareness of meningitis after recent health education campaigns compared to awareness re UTI in babies.</p> |
| 008 | NHS NorthEast | <ul style="list-style-type: none"> <li>Accurate diagnosis of UTI</li> </ul>  | Accurate diagnosis underpins appropriate further management.   | <p>Despite emphasis on diagnosis in NICE CG54</p> <p>Many children still being labelled as 'possible' UTI without a conclusive diagnosis, resulting in some children being referred to secondary services with dysuria from other causes etc</p> | CG54   |
| 009 | NHS           | <ul style="list-style-type: none"> <li>Start treatment early if</li> </ul>   | Evidence of increased risk of  | Evidence of variation in practice  | CG54   |

| ID  | Stakeholder   | Suggested key area for quality improvement  | Why is this important?   | Why is this a key area for quality improvement?  | Supporting information   |
|-----|---------------|---|--|--|--|
|     | NorthEast     | high likelihood of UTI & adjust depending on culture result.  | scarring with delay in treatment   |  |  |
| 010 | NHS NorthEast | <ul style="list-style-type: none"> <li>Investigation plans should focus on those with high likelihood of identifying abnormality eg. recurrent UTI, youngest. Those identified after 1<sup>st</sup> UTI or recurrence with scarring or VUR or other anatomical or functional urinary tract problem, should have individualised care plan for rapid access to diagnosis &amp; treatment of further UTIs</li> </ul> | Evidence that those with underlying renal anatomical or functional abnormality identified more likely to have recurrence of UTI & poorer outcome. Access to prompt diagnosis of UTI & rapid treatment important & needs to be individualised to circumstances of the child, family & local services. | Prevention of worsening of chronic kidney disease.<br>Prevention of prolonged & recurrent illness  | NICE Quality standards for management of Chronic Kidney Disease. |
| 011 | NHS NorthEast | <ul style="list-style-type: none"> <li>Young babies with possible UTI refer to secondary centre for confirmation of diagnosis and rapid treatment.</li> </ul>   | Evidence youngest children most likely to have underlying renal pathology & get most unwell with sepsis. Age range needs careful consideration; CG54 says <3m but older infants & children who are unwell should also be included - CG 47 Feverish illness in children.                              | Local arrangements for referral vary from place to place. Delay in secondary assessment implicated in some case reviews with adverse outcomes. | CG54<br>CG47   |

| ID  | Stakeholder | Suggested key area for quality improvement   | Why is this important?  | Why is this a key area for quality improvement?  | Supporting information   |
|-----|-------------|--|---|--|--|
| 012 | RCN         | <ul style="list-style-type: none"> <li>• Monitoring of vital signs</li> <li>• First-line health care and public awareness of the importance of the signs and symptoms of childhood UTI specifically in very young babies.</li> </ul> | <p>1. Important information gained by assessing and measuring vital signs can be indicators of health and ill health. However, we believe they should not be performed in isolation to the broader observation and assessment of the infant, child or young person.</p> <p>In many instances vital signs will be assessed, measured and monitored by health care assistants and nursing students, under the direction and supervision of a registered nurse.</p> <p>There is variation in practice and quality standard in this area will be beneficial and will improve standard of practice.</p> <p>2. Delay in diagnosis and treatment are multi factorial, some evidence of under diagnosis in some areas compared to expected incidence. Evidence that there is an increased risk of scarring is associated with delay</p> | <p>1.The assessment, measurement and monitoring of vital signs are important basic skills for all health care practitioners working with infants, children and young people.</p> <p>2.Improvements in both parental and professional education is associated with parental empowerment, improved appropriate use of health care resources and improved satisfaction in health care</p> | <p>1.Standards for assessing, measuring and monitoring vital signs in infants, children and young people:<br/>RCN guidance for children’s nurses and nurses working with children and young people</p> <p><a href="http://www.rcn.org.uk/_data/assets/pdf_file/0004/114484/003196.pdf">http://www.rcn.org.uk/_data/assets/pdf_file/0004/114484/003196.pdf</a></p> <p>2. NICE CG 37 postnatal care stresses the need for parents to be offered appropriate information to enable them to identify signs and symptoms of common health conditions in babies.</p> |



| ID  | Stakeholder | Suggested key area for quality improvement  | Why is this important?   | Why is this a key area for quality improvement?   | Supporting information  |
|-----|-------------|---|--|---|---|
|     |             |   | in recognition and treatment.  |   |   |
| 013 | RCN         | <ul style="list-style-type: none"> <li>Accuracy in the diagnosis of Childhood UTI</li> </ul>  | Accurate prompt diagnosis underpins prompt treatment and appropriate further management  | Inspite of the emphasis placed on accurate diagnosis by NICE in CG54 many children are still being inadequately diagnosed with inconclusive diagnoses being made  | NICE CG54   |
| 014 | RCN         | <ul style="list-style-type: none"> <li>Prompt treatment of high risk of UTI with subsequent adjustments of treatment according to culture result</li> </ul>   | Increased risk of renal scarring if treatment delayed  | Evidence of practice variation  | NICE CG 54  |
| 015 | RCN         | <ul style="list-style-type: none"> <li>Children identified with a high risk of identifying abnormality i.e. children with scarring, family history of scarring or children with recurrent UTI's or other existing renal anomalies should have a individualised plan for rapid access for prompt accurate diagnosis and treatment</li> </ul> | Evidence for children with underlying renal or functional abnormalities are at increased risk of poor outcome and should have a individualised plan for rapid access for prompt accurate diagnosis and treatment | Prevention of worsening of chronic kidney disease<br><br>Prevention of prolonged recurrent disease with associated costs to the individual and society            | NICE Quality standards for management of Chronic kidney disease |
| 016 | RCN         | <ul style="list-style-type: none"> <li>Young babies with potential UTI should be referred to a secondary centre for accurate confirmation of a diagnosis of UTI and</li> </ul>  | 1.Evidence that the youngest children are at increased likelihood of underlying renal pathology and may rapidly deteriorate with sepsis. Age range needs to be carefully considered:                             | Arrangements for referral vary greatly from place to place. Delays in secondary assessment have been implicated in some case reviews with marked adverse outcome. | CG54<br>CG47  |

| ID  | Stakeholder | Suggested key area for quality improvement  | Why is this important?  | Why is this a key area for quality improvement?   | Supporting information  |
|-----|-------------|---|---|---|---|
|     |             | rapid appropriate treatment   | CG54 states that <3 months but older children who are unwell should also be included. Also CG 47 (as in 1 <sup>st</sup> key area)   |   |   |
| 017 | RCPH        | <ul style="list-style-type: none"> <li>Urine infections in children should be diagnosed and treated promptly</li> </ul> | Urine infections in children (especially young children) can be difficult to diagnose as symptoms can be very non-specific. There is good evidence that untreated urine infections can, in some children, lead to renal scarring with consequent risk of hypertension and chronic kidney disease. | The collection of an uncontaminated urine sample is essential for the diagnosis of UTI but can be difficult in some children. There is anecdotal evidence of great variability in practice. In particular, children with unexplained fever may not be adequately assessed for the presence of UTI. There is also variability in the speed with which samples are transported to the laboratory. | <p>Two ongoing studies are likely to provide very useful information regarding the diagnosis and management of children with UTI:</p> <p>The HQIP funded multisite audit on Childhood UTI which is looking comprehensively at the standards of care delivery in children with UTI; auditing across primary, secondary and tertiary care as well as Microbiology standards derived from the NICE guideline.</p> <p>the DUTY study, looking at symptoms of UTI and urine results in Primary Care</p> <p>Both of these studies are expected to report within</p> |

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|     |             |   |   |   | 12 months.             |
| 018 | RCPH        | <ul style="list-style-type: none"> <li>Children diagnosed to have a urine infection should be investigated promptly as per NICE guideline (54)</li> </ul> | Some children with UTI have underlying abnormalities of the urinary tract and bladder that predispose them to UTI. These abnormalities must be recognised and managed appropriately to avoid the risk of recurrent UTI.                   | UTI can lead to renal scarring with risks of hypertension and chronic kidney disease. Prevention of recurrent UTI reduces the risk of renal damage and the long term health implications accruing from such damage. |                        |
| 019 | RCPH        | Identification of risk factors for renal disease  | Identification of risk factors for renal disease is key to determining further investigations   | Anecdotal evidence suggests that recording of risk factors for renal disease in children with suspected UTI is poor   |                        |
|     | HPA         | Diagnosis of UTI in under 3 year olds   | this is still very valuable in primary and secondary care and therefore rates of diagnosis, or rate of submission of urines in this age group may give you some indication of numbers being diagnoses and how much improvement is needed. |   |                        |
|     | HPA         | How to improve identification of those children with UTI which require further investigation due to increased risk of underlying pathology.               | NICE needs to be very definite about who should be referred and then audit the parameters chosen.   |   |                        |
|     | HPA         | Standards for definition of   | Standards for definition of UTI in  | Currently there is variable reporting by labs.  |                        |

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|    |                               | UTI in the laboratory                      | the laboratory – 105 CFU/ml or 104 CFU/ml or 103 CFU/ml if pure culture?  |  |   |
|    | Specialist committee member 1 | Diagnosis (clinical) -1                    | Under-diagnosis, especially in young children may lead to adverse health outcomes and an inability to stratify future management according to risk  | Accurate diagnosis prevents future morbidity including end-stage renal failure in a small minority. This is a key issue for both the North-East guideline group and NICE but they have come to moderately different conclusions, most particularly on localisation of UTI              | CG54  |
|    | Specialist committee member 1 | Diagnosis (clinical)-2                     | Inappropriate diagnosis: as diagnostic tests have an element of inaccuracy, and a diagnosis of UTI leads to investigations that may be unpleasant and harmful, it is important to ensure that tests are undertaken appropriately. | Urine samples should not be tested where there is an alternative source of fever (and where an alternative diagnosis would adequately explain the symptoms and level of fever)   | CG54 (bracketed info is non-evidence-based)                           |
|    | Specialist committee member 1 | Investigations 1                           | When UTI is suspected, urine testing should be undertaken within 24 hrs   | Ensures accurate diagnosis and risk stratification. There are practical difficulties, especially in primary care that may prevent this from occurring.   | CG54  |
|    | Specialist committee member 1 | Investigations 2: urine collection         | Accurate sampling is essential to ensure neither under- nor over-diagnosis  | Clean-catch is recommended and pads if this cannot be achieved. Prior to CG 54 it was common to use U-bags. Clean-catch is seen as difficult but gives good results and may be quicker and simpler. Potties, suitable washed, may provide an alternative, but this method has not been | CG54 (+ some personal anecdote). Potties: Vernon S, Coulthard M et al |

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|    |                               |   |  | extensively trialled.   |   |
|    | Specialist committee member 1 | Investigations 3: urine containers  | Refrigeration or boric acid containers should be used if urine cannot reliably be cultured within 4hrs. (Smaller volume urine collection tubes should be used for children)  | In primary care, especially at more remote locations, refrigeration and/or transport to a laboratory within 4 hrs are impractical, so boric acid tubes are used. However, adult tubes require larger volumes and risk under-filling and therefore, false negative results | CG54 (bracketed info from North-East guidelines)  |
|    | Specialist committee member 1 | Investigations 4: laboratory reports  | Urine reports should report the organisms grown in a way that will identify atypical UTIs  | There is historically variable implementation of this criteria – presumably for reasons of cost – but E Coli should be differentiated from non E Coli coliforms   | CG54/ personal experience/ HQIP audit Implementation of NICE guidelines in primary and secondary care (NB pre-publication so confidential information)  |
|    | Specialist committee member 2 | Accurate diagnosis of UTI and in particular of true bacteriuria in infants and small children | Collection of clean urine is very difficult in infants and small children. There is a high chance both for over and under diagnosis of true UTIs.<br>Infants and small children have a high chance to either be over treated for and wrongly diagnosed UTI or not be treated for a missed UTI. | Over treatment of a true UTI exposes the child to unnecessary antibiotic treatment and unnecessary follow-up.<br>Not treating a missed febrile UTI gives that child a chance to develop kidney scarring.  | Recent recommendations for GPs published in the BMJ last year do not suggest how these problems can be solved.<br>Hay AD et al, How to best diagnose urinary tract infection in preschool children on primary care? BMJ 2011 Oct 25;343:d6316 |
|    | Specialist committee member 2 | Upper UTI (acute pyelonephritis) and lower UTI (acute cystitis) are not                       | Acute pyelonephritis and acute cystitis needs very different treatment and follow-up.  | Children with too little treatment for an upper UTI risk kidney scarring from that under treatment. Children with too much antibiotic   | Recent recommendations as published in the BMJ, see above, do not   |

| ID | Stakeholder                   | Suggested key area for quality improvement                                      | Why is this important?   | Why is this a key area for quality improvement?   | Supporting information   |
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|    |                               | dealt with a separate entities.   |  | treatment of a cystitis risk unnecessary side effects from that treatment which also causes unnecessary development of antibiotic resistance. | separate the management of upper and lower UTI.  |
|    | Specialist committee member 2 | Improvement of the use of antibiotic prophylaxis in children who have had a UTI | Appropriate use of prophylactic antibiotics can protect children from developing kidney scarring. Over use of antibiotic prophylaxis will cause unnecessary side effects and bacterial resistance. | Protecting kidney function and reducing development of antimicrobial resistance are too important goals in medicine.                          | At the time the recent UTI guidelines were written by NICE no scientific knowledge existed in this field. Since then six good studies have been published. |