Revised AAP Guideline on UTI in Febrile Infants and Young Children

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In 2011, the American Academy of Pediatrics released a revision of its 1999 clinical practice guideline on urinary tract infections in febrile infants and young children two to 24 months of age. The new clinical practice guideline has several important updates based on evidence generated over the past decade. The updated guideline includes clinical criteria for collecting urine specimens. Diagnosis now requires evidence of infection from both abnormal urinalysis results and positive urine culture results (the criterion for a positive culture has been reduced from at least 100,000 colony-forming units per mL to at least 50,000 colony-forming units per mL). Oral treatment now is considered to be as effective as parenteral treatment. Renal and bladder ultrasonography is still recommended, but the biggest change in the current guideline is that routine voiding cystourethrography is no longer recommended after the first urinary tract infection. Follow-up is based on evaluating children for urinary tract infection during subsequent febrile episodes, rather than routinely performing repeat urine cultures. (Am Fam Physician. 2012;86(10):940-946. Copyright © 2012 American Academy of Family Physicians.)
importance in this age group. Moreover, the evidence from which the action statements are based comes from studies of children two to 24 months of age, and it is not clear whether the data can be applied to other children. An algorithmic summary of the guideline is presented in Figure 1. Readers are encouraged to consult the full guideline and technical report for additional information and rationale.

Presumptive Therapy for Ill-Appearing Children

Action Statement 1: If a clinician decides that a febrile infant or young child with no apparent source for the fever requires antimicrobial therapy because of ill appearance or another pressing reason, the clinician should ensure that a urine specimen is obtained for both culture and urinalysis before an antimicrobial is administered. The specimen should be obtained by catheterization, because the diagnosis of UTI cannot be established reliably by a culture of urine collected in a bag. (Evidence Quality A; Strong Recommendation.)

The only change from the 1999 guideline is that both urinalysis and culture should be performed to assure a diagnosis of true UTI rather than asymptomatic bacteriuria in a child whose fever is unrelated to the urinary tract (see action statement 3). A dependable specimen needs to be obtained before the administration of an antimicrobial because most common agents will sterilize the urine rapidly, obscuring a UTI diagnosis.

Determining the Likelihood of UTI

The second set of action statements addresses patients whom clinicians feel do not need immediate antimicrobial therapy.

Action Statement 2: If a clinician determines that an infant or young child with unexplained fever is not ill enough to require immediate antimicrobial therapy, the clinician should assess the likelihood of UTI.

Action Statement 2a: If the clinician determines this patient has a low likelihood of UTI, clinical follow-up without testing is sufficient, recognizing that the likelihood may change during the course of the illness (e.g., persistence of fever, increased height of fever). (Evidence Quality A; Strong Recommendation.)

Action Statement 2b: If the clinician determines that this patient is not in a low-risk group, there are two choices: obtain a urine specimen by catheterization for culture and urinalysis, or obtain a urine specimen by the most convenient means and perform a urinalysis. If the urinalysis suggests a UTI (i.e., dipstick testing positive for leukocyte esterase or nitrites, or microscopy positive for white blood cells or bacteria), obtain and culture a urine specimen by catheterization or suprapubic aspiration. If urinalysis of fresh urine (less than one hour since void) is negative for leukocyte esterase and nitrites, it is reasonable to follow the clinical course without initiating antimicrobial therapy, recognizing that a negative urinalysis result does not rule out a UTI with certainty. (Evidence Quality A; Strong Recommendation.)

The purpose of these action statements is to acknowledge that although the overall rate of UTI in infants and young children with fever and no apparent source is about 5 percent, some patients have a much higher rate, and others a much lower rate. The guideline committee did not operationalize “low likelihood,” but instead chose to leave it up to the clinician whether to set the

<table>
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<tr>
<th>Area of management</th>
<th>Updates from the 1999 guideline</th>
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<td>Diagnosis</td>
<td>Both an abnormal urinalysis result and a positive urine culture result are needed to confirm inflammation</td>
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<td>A positive culture result is defined as at least 50,000 colony-forming units per mL, rather than the previous criterion of at least 100,000 colony-forming units per mL</td>
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<td>Guidance is added for using clinical criteria to establish a threshold to decide whether to obtain a urine specimen</td>
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<td>Treatment</td>
<td>Oral treatment is as effective as parenteral treatment</td>
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<td>Imaging</td>
<td>Voiding cystourethrography is not recommended routinely after the first febrile UTI; ultrasonography should include the bladder and the kidneys</td>
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<td>Follow-up</td>
<td>Emphasis is on urine testing with subsequent febrile illnesses, rather than on regularly repeated urine cultures after treatment</td>
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NOTE: The guideline applies to infants and children two to 24 months of age with unexplained fever.

AAP = American Academy of Pediatrics; UTI = urinary tract infection.

Information from reference 4.
Diagnosis and Management of UTI in Febrile Infants and Young Children

1. Infant 2-24 mo with fever >38°C

2. Is patient judged to require immediate antimicrobial therapy?

   No

   4. Is likelihood of UTI <1%?

   Yes

   5. Perform urinalysis.

   Option

   6. Obtain urine for urinalysis only by catheter or SPA or bag.

   Option

   7. Conduct enhanced urinalysis with microscope and counting chamber.

   8. Conduct dipstick urinalysis; considered positive if LE and/or nitrite is positive.

   9. Urinalysis positive?

      No

      10. Culture urine obtained by catheterization or SPA.

      11. Treat with antimicrobials effective against common uropathogens according to local sensitivity patterns; oral or parenteral.

      12. Urinalysis and culture positive?

         No

         13. Adjust antimicrobial therapy according to sensitivities. Treat 7-14 d.

         Stop

         Yes

         14. Obtain RBUS any time after UTI is confirmed.

         15. Follow clinical course, reevaluate if fever persists.


         Stop

         17. Second or higher proven UTI or VCUG indicated by RBUS?

            No

            18. Obtain VCUG to evaluate for grade IV-V VUR.

            19. Instruct family to seek medical care for future fevers to ensure timely treatment of UTI.

            Yes

            20. Urologic management as indicated by imaging.

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AAP UTI Guidelines

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1. Risk of UTI is ~5%.
2. A clinician may decide that a febrile infant requires antimicrobial therapy to be administered because of ill appearance or other pressing reason.
3. A urine sample suitable for culture should be obtained before initiating antimicrobials.
4. See Figure 2 for girls and Figure 3 for boys.
5. A urinalysis helps interpret the results of the urine culture, distinguishing UTI from asymptomatic bacteruria.
6. SPA is not recommended unless necessary, because it produces more distress than catheterization.
7. UA that includes microscopy with a hemocytometer has higher sensitivity and specificity, but may not be available.
8. Urine dipstick is slightly less sensitive, but satisfactory if microscopy not available. Positive LE or nitrites or microscopy positive for WBCs or bacteria is a positive urinalysis.
9. If urinalysis is negative, UTI is unlikely (<0.3%).
10. Satisfactory culture is necessary to document a true UTI and to guide antimicrobial management. Only urine obtained by catheterization (or SPA) is suitable for culture.
11. Sensitivities vary by region and time. Base route on practical consideration, eg, unable to retain oral fluids.
12. Pure growth of ≥50,000 CFUs/mL of a uropathogen and urinalysis demonstrating bacteruria or pyuria.
13. Antimicrobial sensitivities of isolated bacteria should be used to adjust antimicrobial choice.
14. Look for anatomic abnormalities that require further evaluation.
15. Follow-up in 1-2 d is important to ensure risk factors have not emerged that would increase UTI risk.
16. Discontinuation of antimicrobials assumes that urine culture was obtained before any antimicrobials were started. Unnecessary antimicrobials can contribute to antimicrobial resistance and may increase risk of UTI.
17. “Proven UTI” means a positive urine culture obtained by suprapubic tap or catheterization. RBUS indications for VCUG should be judged by the clinician.
18. After a second UTI, the risk of grade IV-V VUR, ie, hydronephrosis, is estimated to be 18%.
19. Evaluation ideally within 48 h. Early detection and treatment of febrile UTI may reduce the risk of renal scarring.

Guideline for Obtaining Urine Specimens in Febrile Girls

Is the child ill enough to warrant immediate antimicrobial therapy based on initial clinical assessment?

Yes

Obtain urine specimen and treat (see Figure 1)

No

What is the clinician’s threshold for evaluation?

Lower threshold: Risk of UTI needs to be greater than 1 percent for evaluation (leads to more urine specimens)

Higher threshold: Risk of UTI needs to be greater than 2 percent for evaluation (leads to fewer urine specimens)

How many UTI risk factors* are present?

None or one

Obtain specimen

No specimen

Two or more

Obtain specimen

None to two

Obtain specimen

Three or more

Obtain specimen

*—Risk factors: white race, age younger than 12 months, temperature of at least 102.2°F (39°C), fever lasting at least two days, absence of another source of infection.

Figure 2. Algorithm for deciding when to obtain a urine specimen in girls two to 24 months of age with unexplained fever; based on the clinician’s threshold for risk. (UTI = urinary tract infection.)

Information from reference 4.

Criteria for Diagnosis

The third action statement addresses the importance of requiring both urinalysis and culture.

Action Statement 3: To establish the diagnosis of UTI, clinicians should require both urinalysis results that suggest infection (pyuria or bacteriuria) and the presence of at least 50,000 colony-forming units per mL of a uropathogen cultured from a urine specimen obtained by catheterization or suprapubic aspiration. (Evidence Quality C; Recommendation.)

In the past, a positive urinalysis result was not considered sensitive or specific enough to be included as a diagnostic criterion for
UTI. It is now recognized that pyuria is a hallmark of true UTI and helps distinguish UTI from asymptomatic bacteriuria. It is the host’s inflammatory response that results in scarring; therefore, the presence of white blood cells is an important feature of true UTI. Another change in this action statement is that the threshold criterion for a positive urinalysis result was reduced from 100,000 to 50,000 colony-forming units per mL. This change is based on a reexamination of data from which the previous recommendation was derived, as well as consideration of newer evidence.

**Antimicrobial Therapy**

**Action Statement 4a:** When initiating treatment, the route of antimicrobial administration should be based on practical considerations. Initiating treatment orally or parenterally is equally effective. The clinician should base the choice of agent on local antimicrobial sensitivity patterns (if available) and should adjust the choice according to sensitivity testing of the isolated uropathogen. (Evidence Quality A; Strong Recommendation.)

**Action Statement 4b:** The duration of antimicrobial therapy should be seven to 14 days. (Evidence Quality B; Recommendation.)

The committee would have preferred to recommend a specific number of days of antimicrobial therapy but was unable to find evidence comparing seven, 10, and 14 days head-to-head. Thus, duration of therapy was identified as an area for further research.

**Urologic Imaging**

The biggest change from the 1999 guideline is in the section on imaging. The recommendation to perform renal and bladder ultrasonography is the same, but performing voiding cystourethrography (VCUG) routinely after a first UTI is no longer recommended.

**Action Statement 5:** Febrile infants and young children with UTI should undergo renal and bladder ultrasonography. (Evidence Quality C; Recommendation.)

**Action Statement 6a:** Although VCUG should not be performed routinely after the first febrile UTI, it is indicated if renal and bladder ultrasonography reveals hydronephrosis, scarring, or other findings that suggest high-grade VUR or obstructive uropathy, and in other atypical or complex clinical circumstances. (Evidence Quality B; Recommendation.)

**Action Statement 6b:** Further evaluation should be conducted if there is a recurrence of febrile UTI. (Evidence Quality X; Recommendation.)

Evaluation with VCUG identifies and helps determine the severity of VUR (grades I through V). The test involves radiation, expense, and considerable discomfort but has been recommended because results affect treatment decisions; infants and young children with VUR receive a long course of daily therapy.
antimicrobial prophylaxis, and those with high-grade VUR are candidates for surgical intervention.

In the past decade, the effectiveness of antimicrobial prophylaxis has been questioned. Since 2006, six RCTs comparing prophylaxis with no prophylaxis have been published. To determine the effectiveness in children two to 24 months of age, the authors of the six trials were asked to provide data for children in this age group who were randomized to receive prophylaxis or no prophylaxis after a febrile UTI. The data were provided by grade of VUR, with febrile UTI recurrence as the outcome measure. Data from all six trials were provided, resulting in a data set of 1,091 children (Figure 4). No statistically significant benefit could be demonstrated for prophylaxis in those without VUR or those with grades I to IV VUR (only five children with grade V were included in the trials).

Because recurrence is more common in infants and young children with high-grade VUR, whether they have received prophylaxis or not, postponing VCUG until after a recurrence accomplishes two things: it avoids the radiation, cost, and discomfort of VCUG in 90 percent of these patients with a first febrile UTI, and it identifies those with high-grade VUR, albeit after a recurrence.

In a hypothetical cohort of infants with a first UTI, 65 percent did not have VUR, 29 percent had grade I to III VUR, 5 percent had grade IV, and 1 percent had grade V. Of those with a recurrent UTI, 26 percent did not have VUR, 56 percent had grade I to III VUR, 12 percent had grade IV, and 6 percent had grade V. Waiting for recurrence is considered an acceptable trade-off because the increased percentage of patients who will develop renal scarring is small.

Pediatric urologists have expressed concern that not performing VCUG routinely will lead to undetected VUR, and these concerns have been addressed by the AAP committee. Population studies from two countries on two continents also find that the case for identifying and treating VUR is unconvincing. Guidelines from Britain and Italy now recommend against routine VCUG.

**Parental Education and Follow-up**

**Action Statement 7:** After confirmation of UTI, the clinician should instruct the child’s caregivers to seek prompt (ideally within 48 hours) medical evaluation for future febrile illnesses to ensure that recurrent infection can be quickly detected and treated. (Evidence Quality C; Recommendation.)

In the past, frequent follow-up cultures were recommended to identify asymptomatic recurrences. However, it is now recognized that this likely misidentified girls with asymptomatic bacteriuria as having recurrent or chronic UTI. Because it is the host response that causes scarring, and the host response includes both fever and inflammation (white blood cells), it is more appropriate to detect and treat febrile recurrences than to perform periodic cultures.

**Areas for Further Research**

The extensive literature review and analysis form the basis for the evidence-based guideline, but they also identify gaps in the evidence. Therefore, a section of the guideline is devoted to areas for research with the hope that the next revision of the guideline, like the current one, will be based on new evidence that permits refinement of recommendations for diagnosis and management. The

![Figure 4](https://example.com/figure4.png)

**Figure 4.** Recurrence rate of urinary tract infection or pyelonephritis in febrile infants and young children two to 24 months of age with and without antimicrobial prophylaxis.

Information from reference 4.
Table 2. UTI in Febrile Infants and Young Children: Areas for Further Research

The relationship between UTI and reduced renal function/hypertension: It has long been presumed that UTIs, particularly when coupled with VUR, lead to reduced renal function/hypertension. This presumption has been challenged, however.9

Alternatives to invasive urine collection and culture: Diagnostic methods that do not require invasive urine collection would be desirable, but are not currently available.

The role of VUR (and, thus, voiding cystourethrography): Additional studies are underway (see below).

The role of prophylaxis: The RIVUR (Randomized Intervention for Children with Vesicoureteral Reflux) study has completed enrollment, and results are expected in the next few years.

Genetics: VUR is known to be familial, and asymptomatic bacteriuria also appears to have genetic determinants. Further elucidation of these determinants may prove useful.

UTIs in Hispanics: Although data about UTI incidence and recurrence rates in blacks and whites are available, less is published about rates in Hispanics.

Further treatment: Delaying voiding cystourethrography until after a recurrence defers the decision about what to do when VUR is identified. Evidence is needed to guide further treatment (i.e., what kind of treatment is best for which children?).

Duration of treatment: Head-to-head trials comparing various treatment durations (e.g., seven, 10, or 14 days) would be useful.

UTI = urinary tract infection; VUR = vesicoureteral reflux.

Information from references 4 and 9.

guideline identifies eight areas for research, which are outlined in Table 2.4,9

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REFERENCES


