



Developing a Tool for Prospective Assessment of Treatment Appropriateness in Urinary Tract Infections



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Background

- Antimicrobial resistance is an increasingly serious threat to global public health¹
- Audits and endorsement of appropriate antimicrobial use should be a priority for all antimicrobial stewardship programs²
- To assess treatment appropriateness, it is recommended that a standardized tool be developed³ for quality improvement
- Many UTI treatment guidelines exist, however there are limited published tools designed for prospectively assessing treatment appropriateness

Objective

- Develop an assessment tool for auditing UTI treatment that assesses appropriateness, based on guideline concordance:
 - Prospective
 - Standardized to drive the approach
 - High inter-rater agreement

Methods

- The project team drafted a survey tool for assessing appropriateness of antibiotic therapy in patients with UTI
- The tool was developed using an iterative approach
- Two auditors independently reviewed UTI antibiotic therapy in 50 cases in October 2016
 - On the basis of local UTI guidelines, the auditors noted whether the therapy was “appropriate” or “suboptimal” (i.e. guideline concordant, or not)
- Inter-rater agreement between the two auditors was estimated with Cohen’s kappa statistic⁴
- A minimum of 48 cases needed to be assessed to detect a statistically significant kappa of 0.80 or greater in a two-tailed test ($P \leq 0.05$), with 80% power⁵

Figure 1: UTI Treatment Appropriateness Assessment Tool

Assess Based on Day of Therapy: 1-2 (Refer to A + B) 3-4 (Refer to A + C + D) 5 & later (Refer to A + C + D + E)

A. Diagnosis		<input type="checkbox"/> Signs & symptoms of UTI documented
a) Signs & symptoms documented by physician:		<input type="checkbox"/> No symptoms documented or delirium only
<input type="checkbox"/> Dysuria or urgency or frequency or suprapubic pain <input type="checkbox"/> Flank pain or back pain or CVA tenderness <input type="checkbox"/> New fevers or rigors or neutrophilia without other source <input type="checkbox"/> Non-specific symptoms in a patient with spinal cord injury or paralysis <input type="checkbox"/> Sepsis (per qSOFA score) without other known etiology		
b) Investigations:		
<input type="checkbox"/> Urinalysis performed <input type="checkbox"/> Urine culture performed		
B. Empiric Therapy		<input type="checkbox"/> Appropriate
Suspected Infection of Urinary Source, based on symptoms:		<input type="checkbox"/> Suboptimal
<input type="checkbox"/> No Documented Symptoms <input type="checkbox"/> Uncomplicated Cystitis <input type="checkbox"/> Complicated Cystitis <input type="checkbox"/> Pyelonephritis <input type="checkbox"/> Urosepsis/Febrile UTI <input type="checkbox"/> CAUTI (<input type="checkbox"/> Catheter removed if feasible)		
Drug	Dose	
<input type="checkbox"/> Appropriate per local guideline on back page, patient history, or as recommended by ID physician <input type="checkbox"/> Therapy Too Broad Spectrum <input type="checkbox"/> Inappropriate (Consider allergy and recent antibiotic use)	<input type="checkbox"/> Appropriate <input type="checkbox"/> Suboptimal (Consider renal function)	
C. Culture-Directed Therapy		<input type="checkbox"/> Appropriate
<input type="checkbox"/> No Documented Symptoms <input type="checkbox"/> Uncomplicated Cystitis <input type="checkbox"/> Complicated Cystitis <input type="checkbox"/> Pyelonephritis <input type="checkbox"/> Urosepsis/Febrile UTI <input type="checkbox"/> CAUTI (<input type="checkbox"/> Catheter removed if feasible)		<input type="checkbox"/> Suboptimal
Drug (check all that apply)	Dose	
<input type="checkbox"/> Appropriate for confirmed or presumed pathogen, or as recommended by ID physician <input type="checkbox"/> Opportunity for de-escalation <input type="checkbox"/> Bug-drug mismatch <input type="checkbox"/> Therapeutic duplication <input type="checkbox"/> Suboptimal (Consider allergy)	<input type="checkbox"/> Appropriate <input type="checkbox"/> Suboptimal (Consider renal function)	
D. Route		<input type="checkbox"/> Appropriate
<input type="checkbox"/> Current route indicated <input type="checkbox"/> IV to PO step-down indicated (Clinically stable & symptoms improving, functional GI, tolerates oral) <input type="checkbox"/> Organism not susceptible to oral options		<input type="checkbox"/> Suboptimal
E. Duration (Total treatment)		<input type="checkbox"/> Appropriate
<input type="checkbox"/> Appropriate per local guideline on back page or as recommended by ID physician <input type="checkbox"/> Too long <input type="checkbox"/> Too short		<input type="checkbox"/> Suboptimal
Ultimately, was this UTI treated appropriately? (Must be symptomatic* and “Appropriate” in all applicable categories)		<input type="checkbox"/> Appropriate
*Unless pregnant, history of solid organ transplant, awaiting urologic procedure with expected bleeding		<input type="checkbox"/> Suboptimal

Figure 2: Inter-rater Agreement

		Auditor #1		
		Appropriate	Suboptimal	Total
Auditor #2	Appropriate	22	3	25
	Suboptimal	2	23	25
	Total	24	26	50

Kappa = 0.80 (95% CI: 0.63-0.97)
“Substantial Agreement”

Results

- A tool with five sections was developed (Figure 1):

Day of therapy	Sections assessed by auditors
1-2	A+B
3-4	A+C+D
5 & later	A+C+D+E

- 50 cases were assessed as “appropriate” or “suboptimal”, in different sections of the tool, depending on day of therapy
- If a case was deemed “appropriate” in all applicable sections, it was adjudicated as “ultimately appropriate” and inter-rater agreement was estimated on this final adjudication
- The auditors had the same adjudication in 45 of 50 cases (90% agreement) (Figure 2), kappa was 0.80 (95% CI: 0.63-0.97)

Discussion

- Our group has designed a tool that can be used for prospective auditing with a standardized approach
- The tool had substantial inter-rater agreement⁴ for assessing appropriateness of UTI therapy in field testing
- A unique design feature of this tool is that patients were assessed using different sections of the form, depending on where they were in the course of antimicrobial therapy
- This tool is useful for conducting point prevalence surveys and serves as a template for other antimicrobial stewardship teams to modify according to their needs
- With modification, the described assessment process is also applicable to other infectious disease syndromes
- The main limitation of this tool is that it has not been tested by other trained auditors for usability

Conclusions

- The tool and its development process provide a template that can be used by other antimicrobial stewardship teams to implement audits of treatment appropriateness and improve quality of care.

References available upon request

