



Analysis of re-prescribing of antibiotics used for urinary tract infection in the community using national information

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Background

Lower urinary tract infection (UTI) is common, particularly in women, and is the second most common reason for antibiotic use in the community¹. Initial antibiotic treatment is usually empirical and in Scotland *E. Coli* is the most frequent cause of UTI. National guidance recommends a 3-day course of trimethoprim or nitrofurantoin for non-pregnant women of any age with acute lower UTI^{2,3}.

Antibiotic susceptibility has been stable over the last 5 years although non-susceptibility to trimethoprim remained high at 36.7%. The majority (97.3%) of isolates remain susceptible to nitrofurantoin⁴. However samples submitted to microbiology departments will be biased towards resistance as samples may only be submitted in complicated cases or where patients have failed on empirical treatment.

A meta-analysis published in 2015 reported 3-day courses of nitrofurantoin had lower clinical efficacy than longer courses⁵ and this observation had been reported anecdotally by some prescribers in Scotland.

Objectives

To identify the proportion of adult women who received a prescription for nitrofurantoin or trimethoprim returning for a further course of antibiotics within 7 days of completion of the initial course.

To determine if adult women who receive 3-day courses had more repeat courses than those who receive 5 or 7-day courses of nitrofurantoin or trimethoprim.

Methods

Records from the Prescribing Information System (PIS) E-messaging were linked to SMR-01 data to:

- identify females aged ≥ 16 years with a community prescription for nitrofurantoin 50mg tablets/capsules or 100mg modified release capsules or trimethoprim 200mg tablets from 1 January to 31 December 2016
- split the cohort into three groups: 3; 5; and 7-day courses on the basis of the quantity of tablets supplied
- exclude patients who had antibiotics prescribed or a hospital admission in the previous 90 days.

The outcome of interest was the proportion of individuals who received a further prescription for one of the following antibiotics within 7 days of the date of completion of the initial antibiotic course: cefaclor, cefalexin, ciprofloxacin, co-amoxiclav, fosfomycin, methenamine hippurate, nitrofurantoin, norfloxacin, ofloxacin, pivmecillinam, trimethoprim.

Logistic regression was used to test the effect of course length on treatment success. Age group; Charlson score (prior 5 years); number of medicines by BNF paragraphs prescribed in prior 12 months; care home status and exposure to antibiotics (DDDs) in the prior 12 months were used as additional predictors.

Results

Within the study period 144,004 (61.8%) of 233,248 patients met the inclusion criteria: 35,387 (24.6%) were prescribed nitrofurantoin and 108,617 (75.4%) were prescribed trimethoprim. The number of patients included was similar across all age groups. Only 1% of patients were care home residents. Around half of patients had an unknown Charlson score and 40% had a score of zero.

In females who received trimethoprim 92.8% did not have a further prescription for UTI antibiotics within 7 days of the end of the initial course. In those receiving nitrofurantoin 94.2% had no further UTI antibiotics prescribed within 7 days (Table 1).

Table 1 Number (%) of patient prescribed trimethoprim and nitrofurantoin that returned within 7 days for a further prescription.

Antibiotic	Total Number (%)	No repeat antibiotic within 7 days Number (%)	Repeated antibiotic within 7 days Number (%)
Trimethoprim	108617	100820 (92.8%)	7797 (7.2%)
Nitrofurantoin	35387	33276 (94.2%)	2111 (5.8%)

After adjusting for other factors 3-day courses of trimethoprim were not associated with an increased rate of re-prescribing compared to 5 or 7-days. Seven day courses were associated with a statistically significant increase in repeat antibiotics compared to 3-day course (Figure 1).

In those who received nitrofurantoin after adjusting for other factors the re-prescribing rate in those who received 3-day courses (6.5%) was higher than those who received 5-day courses (5.3%) and 7-day courses (5.2%; Figure 2).

Fig. 1 Trimethoprim: re-prescribing rate for 3, 5 and 7-day courses

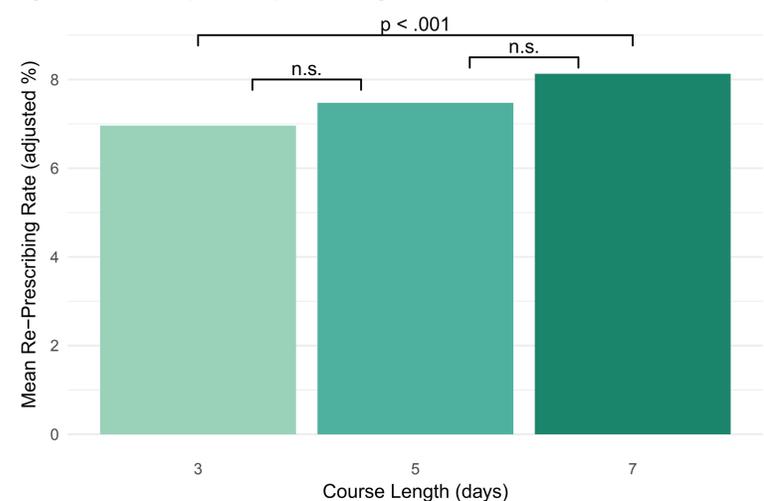
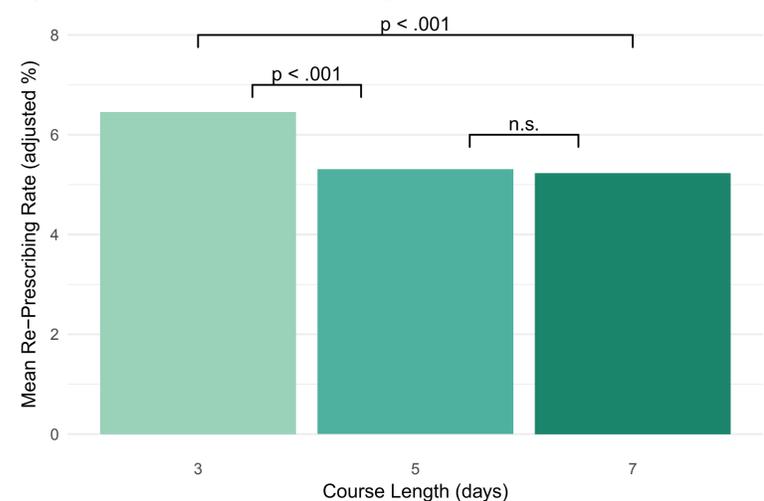


Fig. 2 Nitrofurantoin: re-prescribing rate for 3, 5 and 7-day courses



Discussion

This study using routine national data indicates that despite the reported high levels of trimethoprim non-susceptibility in *E. Coli* in urine isolates only 7.2% of adult females returned within 7 days of the date of completion of the course for a further prescription for a UTI antibiotic. Moreover only 5.8% of females who received nitrofurantoin had a further UTI antibiotic within 7 days.

Three day courses of nitrofurantoin but not trimethoprim were associated with higher rates of repeat antibiotic prescribing compared to 5 or 7-day courses. However the absolute difference was low.

Conclusion

These data should provide reassurance to clinicians that trimethoprim and nitrofurantoin remain appropriate first choice empirical antibiotics in females aged ≥ 16 years with symptoms of simple uncomplicated lower urinary tract infection.

References

1. Christiaan F, Dolk K, Pouwels KB et al. Antibiotics in primary care in England: which antibiotics are prescribed and for which conditions? J Antimicrob Chemother 2018; 73 Suppl 2:ii2-ii10
2. Scottish Intercollegiate Guidelines Network. SIGN 88 – Management of suspected bacterial urinary tract infection in adults.
3. Public Health England. Managing common infections in primary care guidance for consultation and adaptation.
4. Health Protection Scotland. Scottish One Health Antimicrobial Use and Antimicrobial Resistance Report 2016. Health Protection Scotland, 2017
5. Huttner A, Verhaegh EM, Harbarth s et al. Nitrofurantoin revisited: a systematic review and meta-analysis of controlled trials. J Antimicrob Chemother 2015; 70: 2456-2464