female patients with UTI symptoms fulfilling inclusion criteria

informed consent

randomization to trial treatment, baseline data

1x1 placebo granule sachet
3x400 mg ibuprofen tablets (3 days)

1x1 fosfomycin-trometarol granule sachet
3x1 placebo tablets (3 days)

telephone-interviews: evaluation of symptoms, AIA, antibiotic intake, AE, SAE

day 0

day 1, 3, 5, 7

day 8-27

day 28

telephone-interview: evaluation of symptoms, activity impairment-assessment, antibiotic intake, AE, SAE

6month

telephone-interview: Evaluation of UTI, urosepsis, pyelonephritis

12 month

final telephone-interview: evaluation of UTI, urosepsis, pyelonephritis

study completion

Figure 1: study plan
Abbreviations: UTI= Urinary Tract Infection, AE=Adverse Event, SAE=Serious Adverse Event, AIA=Activity Impairment Assessment