Participant Study Number:	_		Participant initials
	Centre	Study ID	

ALternatives **T**o prophylactic **A**ntibiotics for the treatment of **R**ecurrent urinary tract infection in women (the ALTAR Study)



Complete	d by:			
Name:			Signature:	
Date				
	Day	Month	Year	
TRUCTION	FOR COMPL	ETION		

Please place a [X] or insert [requested information] in appropriate box

If you make any errors while completing this form, please strikethrough through the incorrect data with a horizontal line and initial and date any changes.

Please enter the unique MACRO identification number generated on starting this form at the top of Page 4. Study ID will only be available if the participant is subsequently randomised.

Please contact your local recruitment co-ordinator or Central Trial Office if you have any uncertainty regarding completion

Participant Study Number: -						Participant initials		
	C	Centr	е	Stud	y ID			

CONTACTS	
A. PARTICIPANT	
Name	

Address	1 st line:	
	2 nd line:	
	Town:	
	County:	
	Postcode:	

D.O.B								
	D	D	Μ	M	Y	Υ	Y	Y

Hospital Number		
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NHS or CHI Number					

Sex [X]	Female	Male

Preferred telephone number						
Alternative telephone number						

Email Address					
Preferred means of Contact:	Post	Email	Telephone (Landline)	Telephone (Mobile)	Text
1 [most] to 5 [least]					

Participant Study Number: -			Participant initials		
	Centre	Study ID			

B. GENERAL PRACTITIONER		
Surname and Initials		

Practice Name

	1 st line:	
	2nd line:	
Address	Town:	
	County:	
	Postcode:	
Telephone number		

C. HOSPITAL CONSULTANT or GENERAL PRACTITIONER RESPONSIBLE FOR CLINICAL CARE				
DURING WASHOUT PERIOD				
NHS Trust/GP Practice				
[Research Site]				

Consultant/GP Name [Initial]		

GMC Number [7 digit]	
Contact telephone number	
E mail Address	

Participant Study Number: -		_			Participant initials		
	Cent	re	Stud	y ID			

|--|

Sec	tion 1: CURRENT PROPHYLAXIS		
1.	Approximate number of episodes of symptomatic urinary tract infection experienced by participant in 12 months PRIOR to starting current period of prophylaxis	Enter number = 00 –	99
2.	Approximate number of episodes of symptomatic urinary tract infection experienced by participant SINCE starting current period of prophylaxis	Enter number = 00 –	99
3.	Approximate months of current period of continuous use of antibiotic prophylaxis for UTI	Enter number = 00 –	99
4.	Last agent used for current antibiotic prophylaxis for UTI [X]	Amoxycillin	Trimethoprim
		Nitrofurantoin	Cefalexin
		Co-amoxyclav	Ciprofloxacin
		Other (enter name)	

Section 2: OUTCOME OF WASHOUT PERIOD						
1.	Date of start of washout period					
2.	Date of end of washout period					

OUTCOME	
3. Randomised	4. Withdrew
	(check reason below)
	Continued on Returned to
	no prophylaxis brophylaxis brophylaxis
	Other 🗌
	(state)

Participant Study Number: -			Participant initials
	Centre	Study ID	

SY	MPTOMATIC URINARY TRACT INFE	CTIONS DUR	ING WASHOUT PERIO	D
5.	Approximate number of episodes	of	Enter number:	
	symptomatic urinary tract infection	on experience	ed 🛛	
	by participant during washout per	iod		
6.	Severity of symptomatic urinary	Lower UTI	Pyelonephritis	UTI requiring
	tract infections experienced		(fever +/- loin pain)	hospital
	during washout period.			admission
	[State number of UTIs 00 – 99]			