



30000			
Ethics numb	er: 14/SC/1143		
Centre ID:			
Participant's	Initials: (If no middle initial insert '-')		
Participant T	rial ID: (Centre ID + GP's Initials)		
	ATAFUTI		
	native Treatments of Adult Female Urinary Tract Infection: a double blind, ontrolled, factorial randomised trial of Uva ursi and open pragmatic trial of ibuprofen		
	GP CONSENT FORM – GP INTERVIEW		
Name of Rese	earcher: Professor Michael Moore		
The informed	consent consists of two parts – the information sheet and this consent form.		
	Please Initial Box		
;	I confirm that I have read and understand the participant information sheet (version 3 dated 17/03/15) for the above study. I have had the opportunity to consider the information, ask questions and have had these answered satisfactorily.		
\	understand that my participation is voluntary and that I am free to withdraw at any time without giving any reason, and without my legal ights being affected.		
	consent to the collection and use of information about me from the nterview in accordance with the participant information sheet.		
r	give permission for my interview with the researcher to be audio recorded. I understand that my name will not be used in this process and that any quotes used will remain anonymous.		





5.		data collected from me up e to be used in the above st		
6.	I agree to take part in the	e interviews for the above stu	dy.	
and any inqui	iries relating to your perso	on Act 1998 the data contro onal information may be add eral Hospital, Southampton	dressed to the Southamp	
Name of Participant		Signature	Date	
Name of Pers	son taking consent	Signature	Date	
When comple		land in Madical Nata		

1 (original) signed consent form to be kept in Medical Notes1 copy for the participant1 copy for the researcher site file

