

Professor D. Sharp, Cotham House, Bristol BS6

participant unique ID [                    ]

participant DOB [                    ]



**PATIENT CONSENT FORM B**  
(ISRCTN 16479417)

**Antidepressant drug therapy vs a community based psychosocial intervention for the treatment of moderate postnatal depression: a pragmatic randomised controlled trial**

(Please circle one)

- |   |     |    |
|---|-----|----|
| 1. Have you read and understood the patient information sheet (Version 3, 27/07/07)   | YES | NO |
| 2. Have you received enough information about the study?  | YES | NO |
| 3. Have you had an opportunity to discuss this study and ask any questions?   | YES | NO |
| 4. Have you had satisfactory answers to all of your questions?  | YES | NO |
| 5. Have you had sufficient time to come to your decision?   | YES | NO |
| 6. Do you understand that if you consent to this part of the study, you are agreeing to participate in a study comparing two treatments for postnatal depression: antidepressants and counselling from a Health Visitor?  | YES | NO |
| 7. Do you understand that if you consent to this part of study, you are agreeing to let the Research Associate, or other responsible members of the research team, to look at the medical records for you and your baby?  | YES | NO |
| 8. Do you understand that you are free to withdraw from the study: <ul style="list-style-type: none"><li>• At any time?</li><li>• Without having to give a reason?</li><li>• Without affecting your current or future medical care?</li><li>• That details of your participation up to the time of withdrawal will be stored anonymously on file and may be used in the final analysis of data?</li></ul> | YES | NO |
| 9. Do you agree to possibly being contacted by a Research Associate to talk about your views on the treatment of postnatal depression?  | YES | NO |
| 10. Do you agree to participate in this study?  | YES | NO |

PATIENT'S Signature .....

Name (BLOCK LETTERS) ..... Date .....

Version 3 July 27 2007