## Supplementary document 11: Participant Information Sheet and Consent Form

# Women's interviews following completion of Slimming World® groups

Rec Reference Number: 16/LO/1422

## Supporting women with postnatal weight management (SWAN study)

We would like to invite you to take part in an interview about your experience of taking part in the above study. Please take your time to consider this information and discuss it with others if you wish. Please ask us if you need further information.

Part 1 tells you the purpose of this study and what will happen to you if you take part. Part 2 gives you more detailed information about the conduct of the study.

#### Part 1 of the information sheet

## What is the purpose of the study?

As you kindly agreed to take part in the SWAN study we would now like to ask you about your experiences of being offered the opportunity of joining weight management groups offered by Slimming World® (Alfreton, UK). We would like to find out:

- How easy it was to access a group in your area and how often you were able to attend.
- Whether or not you liked being part of a group and whether attending has helped you make changes in your diet and lifestyle.
- Your experience of and contact with other sources of support which help you to manage your weight.
- Whether or not you think other women will be willing to take part in a future study about postnatal weight management using Slimming World.

#### Why have I been invited?

You have been invited as you took part in the SWAN study and joined a Slimming World® (Alfreton, UK) group.

### Do I have to take part?

No, it is up to you to decide. You can change your mind at any time and withdraw from the study without giving a reason

# What will happen to me if I take part in the interview?

If you are happy to take part:

- You will be asked to sign a consent form to take part when your baby is around 4-6 months old
- A researcher will contact you by telephone to ask a few questions about your experience of taking part in the SWAN study which will take about 30-40 minutes.
- The researcher will ask your permission to audio-record the interview. The interview would be transcribed prior to analysis and we will remove your name and any other identifying information from the typed transcript. Should you wish to review and, if necessary, correct the transcript of your interview, you can request a copy from the Investigator, Professor Debra Bick.

### Will my information be confidential?

We will follow ethical and legal practice and all information about you will be handled in confidence.

All information we collect will be stored securely and kept in locked cupboards in locked offices, only accessible to authorised members of the study team. All study information will be kept in a confidential form, and codes rather than names will be used to identify the interview recording.

We won't use your name or any information that can be used to identify you in any publications. Quotations from the interviews may be used in reports, presentations and papers but will not be traceable to you. All published and unpublished reports will disguise the identity of the individuals.

Your interview transcript will kept for 5 years in a secure location, to allow for any audit of the research that may be required and your interview audio-recording will be destroyed when the study results are published. Only members of the research team will have access to the information you give. The only exception to this will be if you tell us information which suggests a risk of serious danger to yourself or others. In this case, we would inform your health care provider.

### What are the possible benefits of taking part?

The information we obtain will help to show how we can best support women with their weight management after having a baby in the future.

### What are the possible disadvantages and risks of taking part?

We do not anticipate that there will be any disadvantages or risks in taking part. If answering any questions causes any anxiety about your health, information on who to contact to discuss this is included at the end of this leaflet. If you have any problems or concerns about your health, the researcher will be able to suggest you contact your GP or health visitor.

### What will happen to the results of the study?

We will publish the study results in reports required by funders and also in articles published in academic journals. We will also present the results of the study at conferences which may be attended by other parents, those working within maternity services and policy makers. No names of any of the study participants will ever be used in any publication.

#### Who is organising and funding the study?

The research is being funded by the National Institute for Health Research Public Health Research programme. The sponsor of the study is King's College London.

#### Who has reviewed the study?

All research in the NHS is looked at by independent group of people, called a Research Ethics Committee, to protect your interests. This study has been reviewed and given favourable opinion by London Camberwell St Giles Research Ethics Committee.

You have been given a copy of this information sheet to keep and when you complete the consent form you will also be given a signed consent form to keep.

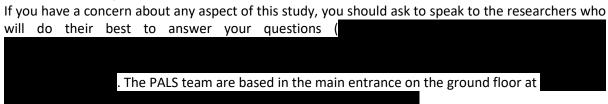
## **Further information and contact details**

You can find out more about this study from the funder's website (http://www.nets.nihr.ac.uk/projects)

## Is there a contact point where I can seek independent advice about participating in the study?

If you would like more information about the study itself you can ask to speak to the Research Midwife. Their contact details are on the back page of this leaflet. The hospital trust's Patient Advice and Liaison Service (PALS) can also be contacted. They will give you advice about how to contact someone for independent advice. Their phone number is on the back page of this leaflet.

## What if there is a problem?



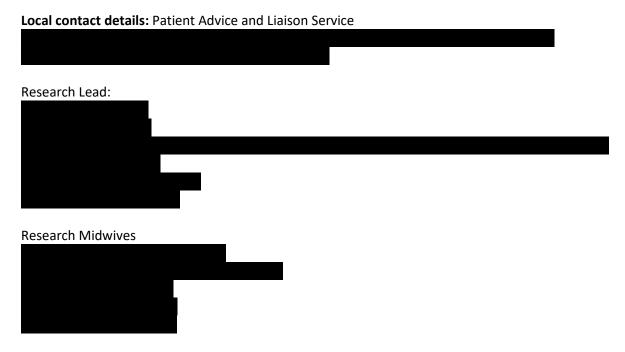
In the event that something does go wrong and you are harmed during the research you may have grounds for legal action for compensation

Trust and/or King's College London but you may have to pay your legal costs. The normal National Health Service complaints mechanisms will still be available to you (if appropriate).

# Where can I find the results of the study?

The results of the study will be published in a medical journal and on the King's College London and the National Institute for Health Research websites when the study has finished. We will send a summary of the study findings to all women who took part in the study when it has finished.

Thank you for taking the time to read this information.



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#### **CONSENT FORM**

Name of Researcher:		Please i	initial box
1. I confirm that I have read an (version 4) for the above study information, ask questions and	. I have had the opportu	•	
2. I understand that my participatime without giving any reason	•	that I am free to withdraw at any gal rights being affected.	
3. I agree to the use of audio-tomy interview	aping if I am interviewed	d, with possible use of verbatim quoto	es from
4. I agree to take part in the a	bove study.		
Name of Patient	Date	Signature	
Name of person		Signature	
taking consent			

An original copy of the participant information sheet and completed informed consent form is to be given to the participant, in addition to the original copy that is filed in the investigator file