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# Support and Reassurance in Antenatal Care

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|---|---|---|
| **Submission date**<br>09/09/2005 | **Recruitment status**<br>No longer recruiting | [_] Prospectively registered<br>[_] Protocol<br>[_] Statistical analysis plan<br>[X] Results<br>[_] Individual participant data |
| **Registration date**<br>23/11/2005 | **Overall study status**<br>Completed | |
| **Last Edited**<br>05/11/2014 | **Condition category**<br>Pregnancy and Childbirth | |

**Plain English summary of protocol**
Not provided at time of registration

## Contact information

**Type(s)**
Scientific

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## Additional identifiers

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**
N&AHP/04/001

## Study information

**Scientific Title**

## Acronym
SRAC

## Study objectives
The project aims to evaluate the effects of two packages of care that may provide women with greater support and reassurance during the third trimester of pregnancy without increasing the number of antenatal contacts that they receive.

Hypothesis: A structured telephone support intervention (with or without supplemental uterine artery Doppler screening) encourages women to adhere to the antenatal visit schedule proposed by the National Institute of Clinical Excellence (2003).

## Ethics approval required
Old ethics approval format

## Ethics approval(s)
Not provided at time of registration

## Study design
Randomised controlled trial

## Primary study design
Interventional

## Secondary study design
Randomised controlled trial

## Study setting(s)
Other

## Study type(s)
Other

## Participant information sheet

## Health condition(s) or problem(s) studied
Antenatal care

## Interventions
Randomised controlled trial of a structured telephone support intervention provided by a midwife +/- uterine artery Doppler screening.

## Intervention Type
Other

## Phase
Not Applicable

### Primary outcome measure
Number of antenatal contacts with health professionals.

### Secondary outcome measures
1. Number of antenatal admissions to hospital
2. Anxiety
3. Levels of perceived social support
4. Satisfaction with care
5. Economic evaluation
6. Major clinical outcomes

### Overall study start date
01/02/2004

### Completion date
30/09/2006

## Eligibility

### Key inclusion criteria
Low risk primiparous women booked for delivery at the Royal Victoria Infirmary, Newcastle upon Tyne

### Participant type(s)
Patient

### Age group
Adult

### Sex
Female

### Target number of participants
764

### Key exclusion criteria
Women who are unable to understand English

### Date of first enrolment
01/02/2004

### Date of final enrolment
30/09/2006

## Locations

### Countries of recruitment
England

United Kingdom

**Study participating centre**
**Research Midwives Office**
Newcastle upon Tyne
United Kingdom
NE1 4LP

# Sponsor information

**Organisation**
Newcastle upon Tyne Hospitals NHS Trust (UK)

**Sponsor details**
Research and Development
Royal Victoria Infirmary
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**Sponsor type**
Hospital/treatment centre

**ROR**
https://ror.org/05p40t847

# Funder(s)

**Funder type**
Government

**Funder Name**
National Co-ordinating Centre for Research Capacity Building (UK) - Nursing and Allied Health Professions Award (ref: N&AHP/04/001)

# Results and Publications

**Publication and dissemination plan**

Not provided at time of registration

## Intention to publish date

## Individual participant data (IPD) sharing plan

## IPD sharing plan summary

Not provided at time of registration

## Study outputs

| Output type | Details | Date created | Date added | Peer reviewed? | Patient-facing? |
|---|---|---|---|---|---|
| Results article | results | 31/03/2014 | | Yes | No |