

# Support and Reassurance in Antenatal Care

<b>Submission date</b> 09/09/2005	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 23/11/2005	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 05/11/2014	<b>Condition category</b> Pregnancy and Childbirth	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

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

## Contact information

**Type(s)**  
Scientific

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## Additional identifiers

**Protocol serial number**  
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

**Study type(s)**

Other

**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(s)**

Number of antenatal contacts with health professionals.

**Key secondary outcome(s)**

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

**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

## Healthy volunteers allowed

No

## Age group

Adult

## Sex

Female

## 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

United Kingdom

England

## Study participating centre

Research Midwives Office

Newcastle upon Tyne

United Kingdom

NE1 4LP

# Sponsor information

## Organisation

Newcastle upon Tyne Hospitals NHS Trust (UK)

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

### 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?
<a href="#">Results article</a>	results	31/03/2014		Yes	No