

Endometriosis and cultural diversity: improving services for minority ethnic women

Submission date 21/10/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 21/10/2010	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 30/09/2016	Condition category Urological and Genital Diseases	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

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
4371

Study information

Scientific Title
Endometriosis and cultural diversity: improving services for minority ethnic women

Acronym

Endocul

Study objectives

To develop, in collaboration with community groups and patients, evidence based and culturally sensitive resources for women with endometriosis and to develop good practice guidance for the health professionals providing services to women of minority ethnic origin.

Ethics approval required

Old ethics approval format

Ethics approval(s)

MREC, ref: 08/H1204/8

Study design

Multicentre non-randomised interventional process of care pilot/feasibility study

Primary study design

Interventional

Study type(s)

Other

Health condition(s) or problem(s) studied

Topic: Reproductive Health and Childb; Subtopic: Oral and Gastrointestinal (all Subtopics), Reproductive Health & Childbirth (all Subtopics); Disease: Reproductive Health & Childbirth

Interventions

1. Develop and pilot a culturally appropriate resource for women with endometriosis in English, Chinese, Greek, Gujarati and Urdu
2. Explore the impact of the socio-cultural context on the meanings of endometriosis among minority ethnic women
3. Develop and pilot good practice in the provision of culturally competent care for providers of endometriosis services in primary and secondary care

Follow up length: 0 months

Study entry: Other

Details: purposive recruitment

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Develop good practice guidance for the health professionals providing services to women of minority

Key secondary outcome(s))

Not provided at time of registration

Completion date

31/10/2009

Eligibility

Key inclusion criteria

Focus groups:

1. Women of African Caribbean, Chinese, Greek/Greek Cypriots, Pakistani, or Indian origin
2. Aged between 18 and 50 years old

Interviewees:

3. Women of African Caribbean, Chinese, Greek/Greek Cypriots, Pakistani, or Indian origin
4. Aged between 18 and 50 years old
5. Current clinical diagnosis of endometriosis
6. Duration of illness greater than 1 year prior to the study
7. Receiving care from mainstream health care services (NHS)

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

Female

Key exclusion criteria

1. Outside of ethnic communities or age range in inclusion criteria
2. No diagnosis of endometriosis

Date of first enrolment

01/04/2008

Date of final enrolment

31/10/2009

Locations

Countries of recruitment

United Kingdom

England

Study participating centre
Birmingham City University
Birmingham
United Kingdom
B42 2SU

Sponsor information

Organisation
Birmingham Women's NHS Foundation Trust (UK)

ROR
<https://ror.org/056ajev02>

Funder(s)

Funder type
Government

Funder Name
National Institute for Health Research (NIHR) (UK) Research for Patient Benefit (RfPB) programme

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?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes
Study website	Study website	11/11/2025	11/11/2025	No	Yes