

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
Registration date 21/10/2010	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 30/09/2016	Condition category Urological and Genital Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

Study website
<http://www.endocul.co.uk>

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

Secondary study design

Non randomised study

Study setting(s)

Other

Study type(s)

Other

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet

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 measure

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

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/04/2008

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

Age group

Adult

Lower age limit

18 Years

Sex

Female

Target number of participants

Planned sample size: 130; UK sample size: 130

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**

England

United Kingdom

Study participating centre

Birmingham City University

Birmingham

United Kingdom

B42 2SU

Sponsor information**Organisation**

Birmingham Women's NHS Foundation Trust (UK)

Sponsor details

Mindelsohn Way

Edgbaston

Birmingham

England

United Kingdom

B15 2TG

Sponsor type

Hospital/treatment centre

Website

<http://www.bwhct.nhs.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

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