

Primary care genetics education and clinic location: a cluster randomised trial

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
23/08/2005	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
16/09/2005	Completed	<input checked="" type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
08/01/2009	Other	

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

SUHT R&D ref: RHM CHI 0250 - UoS ref: 3348

Study information

Scientific Title

Study objectives

Primary care genetics education will increase the referral rate to clinical genetics and nurse counsellor genetics clinics in primary care will be acceptable to patients seen.

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)

Screening

Health condition(s) or problem(s) studied

Family history of a genetic condition

Interventions

Randomised GP practices to receive genetics education and patients referred with trial GP practices to receive a genetics clinic appointment in primary or secondary care

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Education: GP referral rate to clinical genetics

Location: clinic attendance rate

Key secondary outcome(s)

Education: referral type, referral practice and attitudes to clinical genetics

Location: Patient satisfaction, patient clinic travel costs, NHS clinic costs, clinic waiting times, patients remaining in primary care

Completion date

01/05/2005

Eligibility

Key inclusion criteria

All patients referred to the Wessex Clinical Genetics Service (WCGS) registered with a trial GP practice. All patients of all ages. If a child is referred one parent/guardian will complete the questionnaire, both sexes, all ethnic groups.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Other

Sex

All

Key exclusion criteria

1. Any patient who has previously been seen by the WCGS
2. Non GP referrals will be excluded from the education comparison
3. Urgent and ward referrals will be excluded from the location comparison
4. Patients receiving a home visit as a pre-clinic contact will be excluded from the location comparison

Date of first enrolment

01/07/2002

Date of final enrolment

01/05/2005

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Wessex Clinical Genetics Service

Southampton

United Kingdom

SO16 5YA

Sponsor information

Organisation

Southampton University Hospitals Trust and University of Southampton (UK)

ROR

<https://ror.org/0485axj58>

Funder(s)

Funder type

Government

Funder Name

Genetics Policy Unit, Department of Health (UK)

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?
Results article	Results	01/03/2006		Yes	No