Comparing a patient held reminder card to a doctor held reminder card to improve epilepsy care in the community.

Submission date	Recruitment status	Prospectively registered
23/01/2004	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
23/01/2004	Completed	[X] Results
Last Edited	Condition category	Individual participant data
20/11/2009	Nervous System Diseases	

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Study objectives

To evaluate the effectiveness and usefulness of an epilepsy prompt and record card for the management of people with epilepsy in primary care. In addition the effectiveness of a patient held card will be compared to that of a card kept in the medical records. Evaluation was via a practice based randomised controlled trial over a period of 24 months. The card includes reminders to record key information about the clinical condition as well as prompts to carry out key tasks that have been identified as important in the research literature.

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)

GP practice

Study type(s)

Other

Participant information sheet

Health condition(s) or problem(s) studied

Epilepsy

Interventions

- 1. Doctor held card (card inserted into the patients records)
- 2. Patient held card

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Relate to usage of the card, recording information, the completion of key clinical tasks, seizure severity as well as measures of patient and doctor satisfaction.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/04/1997

Completion date

01/08/1999

Eligibility

Key inclusion criteria

Patients with active epilepsy (either a seizure in the last 2 years or on anti epilepsy medication with a history of epilepsy)

Participant type(s)

Patient

Age group

Not Specified

Sex

Both

Target number of participants

1275 patients, 82 practices (added 20/11/09)

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/04/1997

Date of final enrolment

01/08/1999

Locations

Countries of recruitment

United Kingdom

Wales

Study participating centre

Department of General Practice (CeReS Office)

Cardiff United Kingdom CF4 4XN

Sponsor information

Organisation

Record Provided by the NHS R&D 'Time-Limited' National Programme Register - Department of Health (UK)

Sponsor details

The Department of Health Richmond House 79 Whitehall London United Kingdom SW1A 2NL

Sponsor type

Government

Website

http://www.doh.gov.uk

Funder(s)

Funder type

Government

Funder Name

NHS Evaluation of Methods to Promote the Implementation of Research Findings (National Programme) (UK)

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 summaryNot provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/02/2002		Yes	No