

Assessment of self administration of medicines (SAM) for hospital in-patients on a haematology ward

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 30/10/2019	Condition category Other	<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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Assessment of self administration of medicines (SAM) for hospital in-patients on a haematology ward

Study objectives

To assess whether self-administration of medicines (SAM) improves patients compliance and knowledge of their medicines and disease

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)

Hospital

Study type(s)

Other

Participant information sheet

Health condition(s) or problem(s) studied

Not Applicable

Interventions

A control group will be interviewed to assess baseline knowledge.

Randomised Controlled Trial randomised to:

1. SAM
2. Non SAM Group

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

- 1, Patient compliance
2. Patient knowledge of their medicines and disease

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/09/2002

Completion date

30/09/2004

Eligibility**Key inclusion criteria**

At least 20 patients between 18 and 70 suitable for self administration of medicines (SAM)

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Upper age limit

70 Years

Sex

Both

Target number of participants

20

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/09/2002

Date of final enrolment

30/09/2004

Locations**Countries of recruitment**

England

United Kingdom

Study participating centre
Level 1, The Old Building
Bristol
United Kingdom
BS2 8HW

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
United Bristol Healthcare NHS Trust (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 summary

Not provided at time of registration