

Randomised controlled trial of Dihydrocodeine (DHC) and methadone in the treatment of opiate dependent patients

Submission date 08/03/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 20/04/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 18/09/2007	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English Summary

Not provided at time of registration

Study website

<http://www.show.scot.nhs.uk/CSO/Publications/ExecSumms/JanFeb05/9.8%20Robertson.pdf>

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

K/OPR/2/2/D399

Study information

Scientific Title

Study hypothesis

To determine the efficacy of dihydrocodeine compared to methadone in the treatment of opiate dependence.

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)

Not specified

Study type(s)

Treatment

Participant information sheet

Condition

Opiate dependence

Interventions

Substitute treatment with methadone or dihydrocodeine.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Dihydrocodeine (DHC), methadone

Primary outcome measure

1. Survival
2. Retention in treatment
3. Continued drug use (particularly injecting)
4. Criminal behaviour

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/09/2000

Overall study end date

01/03/2005

Eligibility

Participant inclusion criteria

Opiate dependent patients

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

250

Participant exclusion criteria

Not provided at time of registration

Recruitment start date

01/09/2000

Recruitment end date

01/03/2005

Locations

Countries of recruitment

Scotland

United Kingdom

Study participating centre
Muirhouse Medical Group
Edinburgh
United Kingdom
EH4 4PL

Sponsor information

Organisation

Chief Scientist Office (UK) - Scottish Executive Health Department

Sponsor details

St Andrew's House
Regent Road
Edinburgh
United Kingdom
EH1 3DG
+44 (0)131 244 2077
Peter.craig@scotland.gsi.gov.uk

Sponsor type

Government

Website

<http://www.sehd.scot.nhs.uk/cso/>

ROR

<https://ror.org/01bw7zm61>

Funder(s)

Funder type

Government

Funder Name

Chief Scientist Office (UK) - Scottish Executive Health Department

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	01/12/2006		Yes	No