Random controlled trial of an intervention to increase attendance at parenting group services

Submission date	Recruitment status	Prospectively registered
30/09/2005	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2005	Completed	Results
Last Edited	Condition category	Individual participant data
21/09/2017	Other	[] Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Ms Claire Halsey

Contact details

North Staffordshire Combined Healthcare Trust Department of Psychology Ward 2B, Westcliffe Hospital Turnhurst Road, Chell Stoke-on-Trent United Kingdom ST6 6LA +44 (0)1782 425883 Claire.Halsey@nsch-tr.wmids.nhs.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0158145193

Study information

Scientific Title

Random controlled trial of an intervention to increase attendance at parenting group services

Study objectives

To discover whether exposure to cognitive behavioural methods before an intervention increases the effectiveness of that intervention.

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)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Not Applicable: Service delivery

Interventions

This study applies psychological theories and practice to the problem of client non-attendance at parenting services.

The intervention is in the form of a brief twenty to thirty minute session of cognitive behavioural problem solving before clinical services commence, delivered during a routine home visit which is part of the usual service delivery pattern.

The control group will undergo a similar period of additional time with a staff member but this time will be spent in gaining additional information through naturalistic observation of the family.

Clients who have been referred or self referred for parenting group services will undergo the standard pre-intervention assessment in the form of paper and pencil psychometric tests and these will be again administered following service delivery.

Hypotheses will be tested using data on actual session attendance by clients and through the psychometric data routinely collected.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Attendance at Triple P Parenting group sessions.

Secondary outcome measures

Not provided at time of registration

Overall study start date

10/06/2004

Completion date

01/02/2007

Eligibility

Key inclusion criteria

Parents with one or more child age eleven years old or younger. Our parenting group service is designed for this client group only.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

10/06/2004

Date of final enrolment

01/02/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
North Staffordshire Combined Healthcare Trust
Stoke-on-Trent
United Kingdom
ST6 6LA

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Government

Funder Name

North Staffordshire Research and Development Consortium (UK)

Funder Name

North Staffordshire Combined Healthcare Trust

Funder Name

NHS R&D Support Funding

Results and Publications

Publication and dissemination plan

Added 21/09/17: Published as doctoral dissertation only. No publication intended as of 12/04/2011.

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration