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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

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Additional identifiers

Protocol serial number
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

Study type(s)
Not Specified

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(s)
Attendance at Triple P Parenting group sessions.

Key secondary outcome(s))
Not provided at time of registration

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

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Not Specified

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
United Kingdom

England

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

Sponsor information

Organisation

Department of Health

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

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration