# A randomised controlled trial of cognitive therapy for post-traumatic stress disorder related to civil conflict in Northern Ireland

Submission date	<b>Recruitment status</b> No longer recruiting	<ul><li>Prospectively registered</li></ul>		
30/03/2006		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
30/03/2006	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
20/12/2012	Mental and Behavioural Disorders			

## Plain English summary of protocol

Not provided at time of registration

# Contact information

## Type(s)

Scientific

#### Contact name

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# Additional identifiers

# Protocol serial number

037158 and 069777

# Study information

Scientific Title

#### Study objectives

To determine whether cognitive therapy is an effective treatment for Post-Traumatic Stress Disorder (PTSD) resulting from civil conflict.

#### 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)

Treatment

#### Health condition(s) or problem(s) studied

Post-traumatic Stress Disorder

#### **Interventions**

Patients suffering from PTSD who are referred to the Northern Ireland Centre for Trauma and Transformation (NICTT) in Omagh are invited to participate in the trial. Patients who agree to be included are initially randomly allocated to:

- 1. Immediate cognitive therapy
- 2. A 12 week wait-list control condition, followed by cognitive therapy

The treatment programme is the same as that used in NICTT's recently published audit of cognitive therapy in the treatment of PTSD following the 15th August 1998 car bomb in Omagh (Gillespie, Duffy, Hackman & Clark, 2002, Behaviour Research and Therapy, 40, 345-357). Patients allocated to immediate cognitive therapy receive up to 12 weekly sessions in 12 weeks, followed by a review.

Further treatment sessions are provided, if appropriate. Patients initially allocated to the wait list condition receive no treatment in the first 12 weeks but are then offered the same course of treatment. The main assessments are at pre-treatment/wait, 12 weeks, post-treatment and three and 12 month post-treatment follow-up.

# Intervention Type

Other

#### Phase

**Not Specified** 

### Primary outcome(s)

The primary outcome measures for the trial are:

1. The Post-traumatic Diagnosis Scale (PDS: Foa et al, 1997, Psychological Assessment, 9, 445-

451)

- 2. The Beck Depression Inventory (BDI: Beck et al, 1979, Cognitive Therapy for Depression, Guilford Press, New York)
- 3. The Sheehan Disability Scale (SDS: Sheehan, 1983)

## Key secondary outcome(s))

No secondary outcome measures

#### Completion date

01/03/2006

# **Eligibility**

#### Key inclusion criteria

- 1. Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (DSM-IV) (American Psychiatric Association, 1994) criteria for PTSD
- 2. Have experienced trauma in the context of civil conflict in Northern Ireland or elsewhere
- 3. PTSD considered to be the patient's main problem
- 4. Age 18 to 70 years
- 5. Willing to accept random allocation

#### Participant type(s)

**Patient** 

### Healthy volunteers allowed

No

#### Age group

Adult

#### Lower age limit

18 years

#### Sex

**Not Specified** 

#### Key exclusion criteria

- 1. Unable to travel to Northern Ireland Centre for Trauma and Transformation (NICTT) for regular treatment sessions
- 2. PTSD mainly related to childhood sexual abuse
- 3. Other severe psychiatric or physical disorder that requires immediate treatment in its own right

#### Date of first enrolment

01/08/2003

#### Date of final enrolment

01/03/2006

# Locations

#### Countries of recruitment

**United Kingdom** 

Study participating centre
The Northern Ireland Centre for Trauma and Transformation
Omagh
United Kingdom
BT79 OHW

# Sponsor information

#### Organisation

The Northern Ireland Centre for Trauma and Transformation (UK)

# Funder(s)

# Funder type

Research organisation

#### Funder Name

The Northern Ireland Office (UK)

#### Funder Name

The Wellcome Trust (UK) (refs: 037158 and 069777)

# **Results and Publications**

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	02/06/2007		Yes	No