# Letting the Future In: therapeutic intervention for child sexual abuse

Submission date	Recruitment status  No longer recruiting	Prospectively registered		
21/02/2013		☐ Protocol		
Registration date 25/02/2013	Overall study status Completed	<ul><li>Statistical analysis plan</li></ul>		
		[X] Results		
<b>Last Edited</b> 29/01/2018	Condition category  Mental and Behavioural Disorders	[] Individual participant data		

## Plain English summary of protocol

Background and study aims

The NSPCC has developed a course of therapy called 'Letting the Future In' (LTFI) for children and young people aged between 4 and 18 who have been affected by sexual abuse and are living with a carer who has been assessed as safe. The aims of this study are to determine the effectiveness, costs, strengths and weaknesses of LTFI, and to find out which groups of children benefit the most.

## Who can participate?

Children aged between 4-17 years who have made a disclosure of sexual abuse and who have a safe carer who is willing to participate.

## What does the study involve?

Participants are randomly allocated to either receiving the therapy immediately or after waiting for 6 months. They are offered up to 24 sessions with a trained social worker or therapist. LTFI also emphasises work with the child's safe carer, who may receive up to six individual sessions as well as joint sessions with the child. Participants complete questionnaires when they join the study and after 6 and 12 months.

What are the possible benefits and risks of participating?

Participants allocated to the waiting list may experience an increase in anxiety and decline in mental health and if necessary can leave the study for immediate treatment.

Where is the study run from?

The School for Policy Studies, University of Bristol (UK).

When is the study starting and how long is it expected to run for? February 2013 to February 2015.

Who is funding the study? NSPCC (UK).

Who is the main contact?
Tricia Jessiman
Tricia.Jessiman@bristol.ac.uk

## **Contact information**

## Type(s)

Scientific

#### Contact name

Ms Patricia Jessiman

#### Contact details

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tricia.jessiman@bristol.ac.uk

## Additional identifiers

**EudraCT/CTIS** number

**IRAS** number

ClinicalTrials.gov number

Secondary identifying numbers 13718

# Study information

#### Scientific Title

Letting the Future In: therapeutic intervention for child sexual abuse - randomised trial

#### **Acronym**

Letting the Future In

## **Study objectives**

The study will evaluate a therapeutic intervention for children affected by sexual abuse. The impact of the intervention on children and their 'safe carer' will be measured using validated measures in a before and after design with follow-up. In teams where demand for the service exceeds the team's capacity to deliver, the research design will take the form of a randomised trial with a waiting list control (RCT). Service use data will be collected from the safe carers as part of a cost-effectiveness analysis.

More details can be found at: http://public.ukcrn.org.uk/Search/StudyDetail.aspx?StudyID=13718

## Ethics approval required

Old ethics approval format

## Ethics approval(s)

University of Bristol Faculty of Social Sciences and Law, 15/08/2012, ref: FSSLREC150812

## Study design

Randomised trial with a waiting list control

### Primary study design

Interventional

#### Secondary study design

Randomised controlled trial

## Study setting(s)

Other

### Study type(s)

Treatment

#### Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

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

Anxiety

#### **Interventions**

Letting the Future In: The intervention offers a therapeutic assessment followed by 20 individual sessions with the child or young person. (An additional 10 sessions may be negotiated with the team manager) and sessions with the 'safe carer'.

Follow Up Length: 6 month(s)

#### Intervention Type

Other

#### Phase

Not Applicable

#### Primary outcome measure

Trauma Symptoms Checklist (8-16) 54 items, self report (Briere et al. 1996); Timepoint(s): Baseline (T1), 6 months (T2), 12 months (T3)

## Secondary outcome measures

- 1. Juvenile Victimisation Questionnaire (10 questions) (Finklehor et al. 2005); Timepoint(s): T1, T3
- 2. Parenting Stress Index short form 36 items (Abidin, 1995); Timepoint(s): T1, T2, T3

## Overall study start date

01/02/2013

## Completion date

30/04/2013

# **Eligibility**

## Key inclusion criteria

- 1. Male & Female; upper age limit 17 years, lower age limit 6 years
- 2. Intra- or extra-familial sexual abuse
- 3. Safe carer willing to collaborate in treatment

## Participant type(s)

**Patient** 

## Age group

Child

## Lower age limit

6 Years

## Upper age limit

17 Years

#### Sex

Both

## Target number of participants

Planned Sample Size: 210

## Key exclusion criteria

Severe learning disability

## Date of first enrolment

01/02/2013

### Date of final enrolment

30/04/2013

## Locations

### Countries of recruitment

England

United Kingdom

## Study participating centre

## **University of Bristol**

Bristol United Kingdom BS8 1TZ

# Sponsor information

## Organisation

University of Bristol (UK)

## Sponsor details

Level 6 Education Centre Upper Maudlin Street Bristol England United Kingdom BS2 8DJ

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research-governance@bristol.ac.uk

#### Sponsor type

University/education

#### Website

http://www.bris.ac.uk/

#### **ROR**

https://ror.org/0524sp257

# Funder(s)

## Funder type

Charity

## **Funder Name**

NSPCC (UK) - Grant Codes: ORCA 38414

## **Results and Publications**

## Publication and dissemination plan

Not provided at time of registration

# Intention to publish date

01/06/2015

## 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?
Funder report results	results			No	No