

# Safety and feasibility of neural transplantation in early to moderate Huntington's disease in the UK

**Submission date**  
23/10/2000

**Recruitment status**  
No longer recruiting

**Registration date**  
23/10/2000

**Overall study status**  
Completed

**Last Edited**  
27/08/2019

**Condition category**  
Mental and Behavioural Disorders

- ☐ Prospectively registered
- ☐ Protocol
- ☐ Statistical analysis plan
- ☒ Results
- ☐ Individual participant data

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

Prof Steve Dunnett

### Contact details

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## Additional identifiers

### Protocol serial number

G9825903 and 071659

## Study information

**Scientific Title**

Safety and feasibility of neural transplantation in early to moderate Huntington's disease in the UK

**Study objectives**

To determine the safety, feasibility and efficacy of foetal tissue transplantation as a restorative therapy for Huntington's disease

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Not provided at time of registration

**Study design**

Multicentre, randomised controlled trial

**Primary study design**

Interventional

**Study type(s)**

Treatment

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

Neuroscience, psychiatry

**Interventions**

Patients in matched pairs randomly allocated to experimental and control conditions; the pairs are selected from a larger research cohort undergoing longitudinal assessment but not (yet) selected for participation in the surgical trial.

All subjects will be examined using Positron Emission Tomography (PET) to assess survival and functional integration of the implanted grafts. The imaging component of the study, entitled 'Pre- and post-operative PET studies of Huntington's disease patients receiving human fetal striatal cell implants' (RTF - 071659), is funded by Wellcome trust, as part of the research training fellowship (for Y F Tai).

As of September 2007, this trial is still open, but temporarily suspended following introduction of EU Tissue directive and pending upgrading of facilities to GMP compliance and MHRA accreditation.

The end date of the follow up period will be extended to two years after the last patient was operated on.

Please also note that as of 22nd January 2007 the Sponsor institution of this trial changed. The previous sponsor was:

Cardiff University (UK)

Cardiff

CF10 3XQ

United Kingdom  
+44 (0)29 2087 4000  
[Http://www.cardiff.ac.uk](http://www.cardiff.ac.uk)

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome(s)**

Safety trial:

1. Presence/absence of adverse events
2. Feasibility:
  - a. co-ordination of an effective and efficient programme for foetal tissue collection
  - b. biosafety assessment
  - c. preparation and implantation into identified patients and their long term evaluation on the relevant structural and functional criteria of graft survival and efficacy

Efficacy trial:

Retardation or reversal of the progress of the disease in the affected patients as determined by the neurological, neuropsychological and neuropsychiatric and imaging criteria of neuropsychological tests of cognition using the CAmbridge Neuropsychological Test Automated Battery (CANTAB) test battery.

**Key secondary outcome(s)**

No secondary outcome measures

**Completion date**

30/06/2008

**Reason abandoned (if study stopped)**

This trial is still open, but temporarily suspended following introduction of EU Tissue directive and pending upgrading of facilities to GMP compliance and MHRA accreditation

**Eligibility****Key inclusion criteria**

1. Genetically confirmed Huntington's disease
2. Early to moderate stage of disease
3. Presence of motor signs
4. Availability of one close primary informant

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

Not Specified

**Total final enrolment**

17

**Key exclusion criteria**

1. Advanced disease
2. Still employable in usual occupation
3. Other concurrent major illness
4. Current serious psychiatric disturbance
5. Inadequate social support
6. Below the age of 18

**Date of first enrolment**

01/10/1999

**Date of final enrolment**

30/06/2008

## **Locations**

**Countries of recruitment**

United Kingdom

Wales

**Study participating centre**

**The Brain Repair Group**

Cardiff

United Kingdom

CF10 3US

## **Sponsor information**

**Organisation**

Medical Research Council (UK)

## **Funder(s)**

**Funder type**

Charity

**Funder Name**

The Wellcome Trust (UK) (grant ref: 071659)

**Funder Name**

Medical Research Council (UK) (grant ref: G9825903)

**Alternative Name(s)**

Medical Research Council (United Kingdom), UK Medical Research Council, MRC

**Funding Body Type**

Government organisation

**Funding Body Subtype**

National government

**Location**

United Kingdom

## 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?
<a href="#">Results article</a>	results	01/12/2002		Yes	No
<a href="#">Results article</a>	results	01/06/2013		Yes	No
<a href="#">Results article</a>	results	01/12/2018	27/08/2019	Yes	No