

# Investigating the effectiveness of inpatient mental health rehabilitation services in the NHS and independent sector

<b>Submission date</b> 08/07/2022	<b>Recruitment status</b> Recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
<b>Registration date</b> 01/11/2022	<b>Overall study status</b> Ongoing	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 04/09/2025	<b>Condition category</b> Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

Mental health rehabilitation services provide specialist treatment to people with particularly severe and complex problems. These services include inpatient units and supported accommodation in the community. Over half of the 4400 mental health inpatient rehabilitation beds in England are provided by the independent sector, but there have been no studies investigating the effectiveness of inpatient rehabilitation services that have included the independent sector. Our study aims to address this gap.

### Who can participate?

Any patient with an adequate understanding of English at one of the mental health inpatient rehabilitation units selected for this study.

### What does the study involve?

A research interview which takes about 30 min where participants will be asked a series of questions about how satisfied they are with different aspects of their life, how they spend their time, the freedom they have to make day-to-day decisions, and how they feel about the care you receive from the inpatient rehabilitation service. The research team will also invite participants to a brief 5 min interview at 6, 12, and 18 months after the first interview. These interviews will be done over the telephone or video call (e.g. Zoom or Microsoft Teams).

The research team will also ask participants for their permission to ask a staff member about their abilities, needs, any specific difficulties they may have, their activities in the community, and whether they have been at risk or posed any risk to others. The research team will then contact staff involved in the participant's healthcare at regular intervals over the next 18 months to see if they have been discharged from the inpatient rehabilitation service, and, if they have been discharged, details of where they have been discharged to and whether they have had any readmissions. Participants may ask the researcher to see a copy of these questions.

The research team would also ask participants for their permission to collect some information from their healthcare records about the care they have received from mental health services in the past.

What are the possible benefits and risks of participating?

By participating in this study, participants will help the research team to assess the effectiveness of NHS and independent sector inpatient mental health rehabilitation services. It is hoped that the information collected from this study will help to improve inpatient rehabilitation services in the future. Each participant will be offered £20 in recognition of the time they have given to be involved in the study.

Where is the study run from?

University College London (UK)

When is the study starting and how long is it expected to run for?

From December 2019 to October 2025

Who is funding the study?

National Institute for Health and Care Research (NIHR) Health Services Delivery Research programme (UK)

Who is the main contact?

Christian Dalton-Locke (Project Manager)

c.dalton-locke@ucl.ac.uk

### **Study website**

<https://www.ucl.ac.uk/psychiatry/acer>

## **Contact information**

### **Type(s)**

Principal Investigator

### **Contact name**

Prof Helen Killaspy

### **ORCID ID**

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### **Type(s)**

Scientific

**Contact name**

Dr Christian Dalton-Locke

**Contact details**

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## Additional identifiers

**EudraCT/CTIS number**

Nil known

**IRAS number**

311434

**ClinicalTrials.gov number**

Nil known

**Secondary identifying numbers**

CPMS 52629, IRAS 311434, award ID: NIHR130693

## Study information

**Scientific Title**

Assessing the Clinical and cost-Effectiveness of inpatient mental health Rehabilitation services provided by the NHS and independent sector (ACER)

**Acronym**

ACER V1.0

**Study objectives**

1. Do sociodemographic and clinical characteristics of patients differ between people receiving inpatient rehabilitation in the NHS and the independent sector?
2. Does service quality differ between inpatient rehabilitation units provided by the NHS and the independent sector?
3. Do the experiences of treatment and care, from the perspectives of patients, informal carers and staff, differ between inpatient rehabilitation services provided by the NHS and the independent sector?
4. Is inpatient rehabilitation clinically more effective at preventing readmission when provided by the independent sector or the NHS, after adjusting for differences between the sectors in terms of patient characteristics and length of stay?

5. Is inpatient rehabilitation more cost-effective when provided by the independent sector or the NHS, after adjusting for differences between the sectors in terms of key predictors of costs such as patient characteristics and length of stay?

### **Ethics approval required**

Old ethics approval format

### **Ethics approval(s)**

Approved 08/06/2022, North East - Newcastle & North Tyneside 2 Research Ethics Committee (NHS BT Blood Donor Centre, Holland Drive, Newcastle upon Tyne, Tyne and Wear, NE2 4NQ; +44 (0)2071048086; newcastlenorthtyneside2.rec@hra.nhs.uk), ref: 22/NE/0067

### **Study design**

Observational cohort study

### **Primary study design**

Observational

### **Secondary study design**

Cohort study

### **Study setting(s)**

Hospital

### **Study type(s)**

Treatment

### **Participant information sheet**

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

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

Mental health

### **Interventions**

As we cannot compare NHS and independent sector services through a randomised trial, our programme will draw together findings from four components to assess the effectiveness of these services:

1. Survey of 60 inpatient mental health rehabilitation services across England (30 NHS and 30 independent sector)
2. In-depth interviews with users, relatives/carers, staff, and commissioners of these services to explore their experiences and perspectives
3. Cohort study to compare outcomes for 600 patients of the NHS and independent sector rehabilitation services surveyed in Component 1, using statistical methods that take account of any differences between them such as the severity of their mental health problems
4. Health economic evaluation to assess the cost-effectiveness of inpatient rehabilitation services provided by the NHS and the independent sector

### **Survey:**

We aim to recruit 500 patients from 60 inpatient mental health rehabilitation units across England (30 NHS and 30 independent sector) over a 12-month period to participate in a survey

(patients at this stage will be asked to consent to take part in Components 1 and 3). Once selected services have agreed to participate, researchers will visit the unit and describe the project to staff. The staff will be asked to introduce the patients to the researchers who will then provide the patient with a Participant Information Sheet. The study will be fully explained to the patient, including the purpose of the study, what their participation would involve, what data will be collected and how it will be collected, and that they are free to withdraw from the study at any point without it affecting their clinical care. The patient will have at least 24 hours to consider whether they would like to participate and have the opportunity to ask the researcher any questions. Patients who provide their informed consent by completing the consent form will then participate in a research interview where data will be collected using standardised measures regarding their quality of life (DIALOG, Recovering Quality of Life, EQ-5D-5L), autonomy (Resident Choice Scale), engagement in activities (Time Use Diary), satisfaction with care (Client Assessment of Treatment), and healthcare costs (Client Receipt Inventory). In addition, staff will be asked to complete standardised measures on the patient's social functioning (Life Skills Profile), challenging behaviours (Special Problems Rating Scale), alcohol and drug use (Clinical Alcohol and Drug Scale), needs (Camberwell Assessment of Needs), and activities (Time Use Diary). Sociodemographic, healthcare service use, current and previous risk, and engagement in community-based activities will be collected from the patient's healthcare records. Also, researchers will complete the QuIRC with the managers of each service. The QuIRC is a quality assessment tool developed specifically for inpatient mental health rehabilitation units.

#### In-depth interview (qualitative) study:

We will select and invite 10 inpatient rehabilitation units (5 NHS and 5 independent sector) participating in Component 1 to participate in Component 2. We will invite all staff from these services to participate in a focus group and ask about their experience and perspective of working in these services. We will also conduct additional staff interviews, including interviews with senior service managers, commissioners, and community care coordinators, to expand on any themes not adequately covered in the focus group, including the decision-making process about where an individual receives their inpatient rehabilitation.

From these participating services, we will also aim to interview between three and five patients from each service (between 30 and 50 in total) and ask about their experiences and perspectives on the care they are receiving. We will ask these patient participants for permission to contact and invite their relatives/carers to participate in an in-depth interview. We aim to recruit around 20 relatives/carers in total. Interviews with relatives/carers will take place via telephone or videoconferencing.

#### Cohort study:

Participants recruited into Component 1 will comprise the cohort for Component 3. The researchers will contact the services managers of the inpatient rehabilitation units participating in Component 1 once a month for 18 months to ask if any of the participants have been discharged from the unit, and if they have been discharged, information about where they have been discharged to including contact details of relevant staff (i.e. their 'key informant') who we may complete future monthly follow-ups with. We expect on most occasions this will be the participant's community care coordinator. We will invite all participants to a short research interview via telephone or video call (Teams or Zoom) at six, 12, and 18 months to complete two quality of life measures (Recovering Quality of Life and EQ-5D-5L). In addition, at six, 12, and 18 months we will ask key informants of participants who have been discharged from the inpatient rehabilitation service they were recruited from about any subsequent inpatient service use, and social care costs (use of supported accommodation and any individual 'care packages').

Health economic evaluation:

Using data collected in Components 1 and 3 analyses will be conducted to investigate the cost-effectiveness of NHS and independent inpatient rehabilitation services by calculating quality-adjusted life-years.

## **Intervention Type**

Other

## **Primary outcome measure**

Cohort study:

Successful discharge from inpatient rehabilitation, defined as being discharged to the community without readmission, measured using staff reports collected at 6, 12, and 18 months

Health economic evaluation:

Incremental cost-effectiveness ratio (ICER) comparing rehabilitation in the independent sector and rehabilitation in an NHS service calculated using cost-utility analysis (CUA) where the QALYs are measured using patient responses to the EuroQol 5-dimension 5-level (EQ-5D-5L) questionnaire collected at baseline, 6, 12, and 18 months

## **Secondary outcome measures**

Cohort study:

Total inpatient days over the follow-up period measured using staff reports collected at 6, 12, and 18 months

## **Overall study start date**

20/12/2019

## **Completion date**

31/10/2025

# **Eligibility**

## **Key inclusion criteria**

Patients:

1. Inpatients at rehabilitation units that are participating in the study
2. Provide informed consent to participate in the study. All efforts will be made to maximise the capacity of each patient to be able to provide informed consent for the study (for example, by explaining the purpose and process of the study in simple terms and over a number of meetings), and any patient with capacity who declines participation will not be included. However, for eligible patients who lack capacity to give informed consent we will seek advice from a consultee on their participation. Further detail on the procedure for seeking informed consent and assessing capacity is provided in section 8. Research data for participants lacking capacity will be collected via staff interview and healthcare records (as described above), but these participants will not be asked to participate in an interview themselves. The researchers expect only a minority of eligible patients to lack capacity (around 8% based on the REAL study) but it is important they are included in the study in order to prevent sample bias.

**Services:**

1. High dependency, community, or longer-term high dependency rehabilitation units
2. Community rehabilitation units will only be included if they are registered with CQC as an inpatient unit (rather than supported housing or residential care)

**Participant type(s)**

Patient

**Age group**

Adult

**Sex**

Both

**Target number of participants**

Planned Sample Size: 500; UK Sample Size: 500

**Key exclusion criteria****Patients:**

1. Unavailable during the researcher's visit, because they are on leave from the ward during the visit or are otherwise unavailable
2. Do not have an adequate understanding of English to understand the main purpose of the research and what participating would involve for them

**Services:**

1. Highly specialist units that focus on sub-groups (such as people with a diagnosis on the autism spectrum or those with neurodegenerative disease)
2. Specialist forensic mental health units (i.e. low secure rehabilitation units) as they do not form part of the standard mental health rehabilitation care pathway and are subject to specialist commissioning by NHS England

**Date of first enrolment**

08/06/2022

**Date of final enrolment**

30/09/2025

**Locations****Countries of recruitment**

England

United Kingdom

**Study participating centre**

**Camden and Islington NHS Foundation Trust**

St Pancras Hospital

4 St Pancras Way

London  
United Kingdom  
NW1 0PE

**Study participating centre**

**Avon and Wiltshire Mental Health Partnership NHS Trust**

Bath NHS House  
Newbridge Hill  
Bath  
United Kingdom  
BA1 3QE

**Study participating centre**

**Birmingham and Solihull Mental Health NHS Foundation Trust**

Unit 1  
50 Summer Hill Road  
Birmingham  
United Kingdom  
B1 3RB

**Study participating centre**

**Central and North West London NHS Foundation Trust**

Trust Headquarters  
350 Euston Road  
Regents PLACE  
London  
United Kingdom  
NW1 3AX

**Study participating centre**

**Coventry and Warwickshire Partnership NHS Trust**

Wayside House  
Wilsons Lane  
Coventry  
United Kingdom  
CV6 6NY

**Study participating centre**

**Cumbria, Northumberland, Tyne and Wear NHS Foundation Trust**

St Nicholas Hospital  
Jubilee Road



Gosforth  
Newcastle upon Tyne  
United Kingdom  
NE3 3XT

**Study participating centre**  
**Devon Partnership NHS Trust**  
Wonford House Hospital  
Dryden Road  
Exeter  
United Kingdom  
EX2 5AF

**Study participating centre**  
**Greater Manchester Mental Health NHS Foundation Trust**  
Prestwich Hospital  
Bury New Road  
Prestwich  
Manchester  
United Kingdom  
M25 3BL

**Study participating centre**  
**Herefordshire and Worcestershire Health and Care NHS Trust**  
Unit 2 Kings Court  
Charles Hastings Way  
Worcester  
United Kingdom  
WR5 1JR

**Study participating centre**  
**Leeds and York Partnership NHS Foundation Trust**  
2150 Century Way  
Thorpe Park  
Leeds  
United Kingdom  
LS15 8ZB

**Study participating centre**  
**Lincolnshire Partnership NHS Foundation Trust**  
St George's

Long Leys Road  
Lincoln  
United Kingdom  
LN1 1FS

**Study participating centre**  
**Mersey Care NHS Foundation Trust**  
V7 Building  
Kings Business Park  
Kings Drive  
Prescot  
United Kingdom  
L34 1PJ

**Study participating centre**  
**North Staffordshire Combined Healthcare NHS Trust**  
Lawton House  
Bellringer Road  
Trentham  
Stoke-on-trent  
United Kingdom  
ST4 8HH

**Study participating centre**  
**Nottinghamshire Healthcare NHS Foundation Trust**  
The Resource, Trust Hq  
Duncan Macmillan House  
Porchester Road  
Nottingham  
United Kingdom  
NG3 6AA

**Study participating centre**  
**Pennine Care NHS Foundation Trust**  
225 Old Street  
Ashton-under-lyne  
United Kingdom  
OL6 7SR

**Study participating centre**

**Rotherham Doncaster and South Humber NHS Foundation Trust**

Woodfield House  
Tickhill Road  
Doncaster  
United Kingdom  
DN4 8QN

**Study participating centre**

**South London and Maudsley NHS Foundation Trust**

Bethlem Royal Hospital  
Monks Orchard Road  
Beckenham  
United Kingdom  
BR3 3BX

**Study participating centre**

**South West London and St George's Mental Health NHS Trust**

Springfield Hospital  
61 Glenburnie Road  
London  
United Kingdom  
SW17 7DJ

**Study participating centre**

**Sussex Partnership NHS Foundation Trust**

Trust Hq  
Swandean  
Arundel Road  
Worthing  
United Kingdom  
BN13 3EP

**Study participating centre**

**Tees, Esk and Wear Valleys NHS Foundation Trust**

Trust Headquarters  
West Park Hospital  
Edward Pease Way  
Darlington  
United Kingdom  
DL2 2TS

**Study participating centre**

**St Andrew's Healthcare**

Billing Road  
Northampton  
United Kingdom  
NN1 5DG

**Study participating centre**

**Priory Group**

Priory Head Office Floor 5  
Hammersmith Road  
London  
United Kingdom  
W14 8UD

**Study participating centre**

**Barnet, Enfield and Haringey Mental Health NHS Trust**

Trust Headquarters Block B2  
St Ann's Hospital  
St Ann's Road  
London  
United Kingdom  
N15 3TH

**Study participating centre**

**Oxleas NHS Foundation Trust**

Pinewood House  
Pinewood PLACE  
Dartford  
United Kingdom  
DA2 7WG

**Study participating centre**

**Lancashire and South Cumbria NHS Foundation Trust**

Sceptre Point  
Sceptre Way  
Walton Summit  
Preston  
United Kingdom  
PR5 6AW

**Study participating centre**  
Hampshire and Isle of Wight Healthcare NHS Foundation Trust  
Tatchbury Mount Hospital  
Calmore  
Southampton  
United Kingdom  
SO40 2RZ

## Sponsor information

### Organisation

Camden and Islington NHS Foundation Trust

### Sponsor details

St Pancras Hospital  
4 St Pancras Way  
London  
England  
United Kingdom  
NW1 0PE  
+44 (0)20 7685 5949  
sponsor.noclor@nhs.net

### Sponsor type

Hospital/treatment centre

### Website

<http://www.candi.nhs.uk/>

### ROR

<https://ror.org/03ekq2173>

## Funder(s)

### Funder type

Government

### Funder Name

National Institute for Health and Care Research

### Alternative Name(s)

National Institute for Health Research, NIHR Research, NIHRresearch, NIHR - National Institute for Health Research, NIHR (The National Institute for Health and Care Research), NIHR

**Funding Body Type**  
Government organisation

**Funding Body Subtype**  
National government

**Location**  
United Kingdom

## Results and Publications

**Publication and dissemination plan**  
Planned dissemination of findings through peer-reviewed scientific journal publications by the end of 2025, hosting two webinars for participating services at key stages during the study to report on the progress of the study, and also presentation of the main findings at conferences.

**Intention to publish date**  
30/06/2026

**Individual participant data (IPD) sharing plan**  
The data sharing plans for the current study are unknown and will be made available at a later date

**IPD sharing plan summary**  
Data sharing statement to be made available at a later date

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Protocol file</a>	version 1.3	07/06/2022	05/08/2022	No	No
<a href="#">HRA research summary</a>			26/07/2023	No	No
<a href="#">Protocol article</a>		06/02/2024	07/02/2024	Yes	No