

Depression in patients with heart failure

Submission date 04/09/2022	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 07/03/2023	Overall study status Ongoing	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 28/01/2026	Condition category Circulatory System	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

Many people in the North East and North Cumbria in England (UK) have heart failure, some of whom will also suffer with depression. Having both conditions is particularly challenging. It means you are more likely to have a worse quality of life, feel more fatigued and more likely to need hospital treatment for your heart failure. It also increases the chances of poor outcomes, such as heart transplant or death. It is much harder to diagnose depression in people with heart failure, so you may be less likely to access specialist mental health support. Also, the usual treatments for depression (talking therapies and medication) do not appear to be helpful in patients with heart failure.

The autonomic nervous system controls the unconscious activity in our body. It has an important role in regulating heart rate. Normally there is variation to the heart rate: sometimes the heart beats a little faster and sometimes a little slower. We call this 'heart rate variability' and it is a sign of a healthy heart. Heart rate variability is often reduced in heart failure and in depression. Therefore, we wonder if dysregulation of the autonomic nervous system is an important link between heart disease and mood, leading to worse outcomes.

Who can participate?

Adults over 18 years, with heart failure.

What does the study involve?

We want to learn more about depression in people with heart failure. We will interview people with heart failure and collect information about their mood, fatigue, quality of life and autonomic nervous system function. Many people with heart failure have implanted heart monitors. They allow us to measure their heart rates, and we will use these data to study heart rate variability – this will be an indicator of autonomic nervous system function. A single blood test will be taken to understand how well the heart is functioning, so that we can correlate heart disease severity with our findings.

What are the possible benefits and risks of participating?

Participants will not have any direct benefit from participating in our study. However, the results of our study might one day help improve the way we identify and treat depression in people with heart failure. We hope that this will help patients and families in this situation to improve their quality of life. If you choose to participate, you will be a fundamental part of making things better.

Apart from asking participants some questions and taking a blood test, we are not doing anything to them that wouldn't normally happen. Because of this, we are not expecting any side-effects to develop. As a result of getting a blood sample collected, participants might experience pain, bruising, light-headedness and on rare occasions, infection. We will help them get assistance if that happens. The usual NHS indemnity scheme will apply to all activities of this study.

We will talk about sensitive topics such as a low mood and depression. The person who will interview the participants is a psychiatrist, a medical doctor who is specially trained to do this. We will do our very best to be sensitive in our approach. Participants might still find it distressing and will offer help if that is the case. If we do find that participants have depression and if they feel that they don't have enough help for this, we can help them get in touch with their GP. We will not give them any treatment for depression as part of the study, but we can tell them how to get help.

There will be no changes to the heart failure treatment as a part of this study. If participants feel that they need any extra help, we can help them speak to the doctors and nurses at Newcastle Hospitals who look after their hearts.

Where is the study run from?

Newcastle upon Tyne Hospitals NHS Foundation Trust (UK)

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

February 2020 to June 2026

Who is funding the study?

Newcastle upon Tyne Hospitals Charity and Join Research Executive Scientific Committee Research Grant (UK)

Who is the main contact?

Dr Alan Bagnall, alan.bagnall@nhs.net

Contact information

Type(s)

Principal investigator

Contact name

Dr Alan Bagnall

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Integrated Research Application System (IRAS)
312832

Study information

Scientific Title

Quality of life, fatigue and autonomic dysfunction in patients with heart failure: association with symptoms of low mood and depression.

Study objectives

Autonomic nervous system (ANS) dysregulation may underpin the link between depression, fatigue and poor quality of life in comorbid heart failure (HF). This would help explain why so many people with heart failure also have depression, why it has such an impact on their quality of life and why depression in heart failure patients is so difficult to treat.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 20/11/2022, Newcastle North Tyneside 1 Research Ethics Committee (NHSBT Newcastle Blood Donor Centre, Holland Drive, Newcastle upon Tyne, NE2 4NQ, UK; +44 2071048255; newcastlenorthtyneside1.rec@hra.nhs.uk), ref: 22/NE/0209.

Study design

Observational cross-sectional

Primary study design

Observational

Study type(s)

Screening

Health condition(s) or problem(s) studied

Adult patients with heart failure and cardiac implantable electronic devices (CIED)

Interventions

We will interview people with heart failure and collect information about their mood, fatigue, quality of life and autonomic nervous system function. Many people with heart failure have implanted heart monitors. They allow us to measure their heart rates, and we will use these data to study heart rate variability – this will be an indicator of autonomic nervous system function. A single blood test will be taken to understand how well the heart is functioning, so that we can correlate heart disease severity with our findings.

Intervention Type

Other

Primary outcome(s)

Measured at a single time point:

1. Beck Depression Inventory II (BDI-II) to assess mood.
2. The Quick Inventory of Depressive Symptomatology - Self-report (QIDS-SR) to assess mood.
3. Structured interview, informed by the Mini-International Neuropsychiatric Interview (MINI), to obtain descriptive data about mood.

Key secondary outcome(s)

Measured at a single time point:

1. 5-level EuroQol 5D version (EQ-5D-5L) scale for quality of life.
2. 21-item Minnesota Living with Heart Failure (MLHF) questionnaire for quality of life in people with heart failure.
3. Multi-dimension Fatigue Inventory (MFI) to assess fatigue.
4. Composite Autonomic Symptom Scale-31 (COMPASS-31) to assess symptoms of autonomic function.
5. THINC-it cognitive screening tool for cognition.
6. Blood sample to test for N-terminal pro-brain natriuretic peptide (NT-proBNP) level, to estimate heart function.
7. Heart rate data will be downloaded from participants' cardiac implantable electronic device.
8. Information from patients' clinical records: Data will be collected on: Demographics, including age, sex, gender identity, sexuality, ethnicity, marital status and occupation; Previous physical health and mental health diagnosis; Current medication and medication taken over the previous year, including doses.

Completion date

30/06/2026

Eligibility

Key inclusion criteria

1. ≥ 18 years of age.
2. Open to the heart failure clinics at the Freeman Hospital or Royal Victoria Infirmary (RVI), NuTH.
3. Diagnosis of heart failure with reduced ejection fraction (HFrEF) - severe left ventricle systolic dysfunction (LVSD) with ejection fraction (EF) $< 35\%$.
4. Cardiac implantable electronic device (CIED) in place.
5. Able to provide written informed consent.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Mixed

Lower age limit

18 years

Upper age limit

99 years

Sex

All

Total final enrolment

30

Key exclusion criteria

1. Previous diagnosis of bipolar affective disorder, psychotic disorder or personality disorder.
2. Previous diagnosis of dementia.
3. Previous diagnosis of primary neurological injury (eg, anoxic injury, stroke or traumatic brain injury) or disorder (eg, Parkinson's disease).
4. Myocardial infarction (MI) within the previous 3 months.
5. Not fluent in English.

Date of first enrolment

01/02/2023

Date of final enrolment

20/01/2026

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

The Newcastle upon Tyne Hospitals NHS Foundation Trust

Freeman Hospital

Freeman Road

High Heaton

Newcastle upon Tyne

England

NE7 7DN

Sponsor information**Organisation**

Newcastle upon Tyne Hospitals NHS Foundation Trust

ROR

Funder(s)

Funder type

Charity

Funder Name

Newcastle upon Tyne Hospitals Charity and Join Research Executive Scientific Committee
Research Grant

Results and Publications

Individual participant data (IPD) sharing plan

All data will be stored for up to 5 years after the last data is collected. Data will be available to other researchers on reasonable request, made to the study PI (Dr Alan Bagnall) via email (alan.bagnall@nhs.net).

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
HRA research summary			28/06/2023	No	No
Participant information sheet	version 5.0	02/12/2022	22/12/2022	No	Yes
Protocol file	version 7.0	02/12/2022	22/12/2022	No	No