

# A study aiming to reduce diagnostic delays of motor neurone disease within primary care in the UK using an automated electronic red flag alert tool

<b>Submission date</b> 18/11/2025	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 02/03/2026	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 02/03/2026	<b>Condition category</b> Nervous System Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input checked="" type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

Motor neurone disease (MND) is a rare, incurable condition affecting about 5,000 UK people. It causes muscle weakness over time, making it harder for people to move, speak, and swallow, and eventually leads to difficulty breathing. Because MND is rare, general practitioners (GPs) may not always recognise it right away, leading to delays in diagnosis. On average, it can take 9 to 27 months from first symptoms to a confirmed diagnosis.

To help GPs identify MND earlier, the MND Association and the Royal College of General Practitioners (RCGP) developed the MND Red Flag Tool in 2014. This tool lists common signs and symptoms that may indicate MND, helping GPs decide when to refer patients to specialists more quickly. The MND Alert system builds on this tool as an automated aid that can be added to patient electronic health records in GP practices. It scans patient records for Red Flag Tool signs linked to MND and sends notifications to GPs when these signs are detected, helping them identify possible cases earlier and make faster referrals. Currently, the MND Alert system can detect symptoms recorded as codes in patient records, but some important information may be written in free-text notes, such as in consultation letters. To ensure nothing is missed, some patient records may need additional manual review.

This study has two stages: (1) implementing the MND Alert system in GP practices for 6 months and (2) evaluating how well it works and if GPs find it helpful. In stage 1, the MND Alert system will be used in routine GP patient consultations. In stage 2, we will interview GPs from the participating practices to understand their experiences with the tool and assess how effective and practical it is for everyday clinical use.

### Who can participate?

GPs at participating sites in South Yorkshire

What does the study involve?

The tool will alert the GP during consultations to service users who may have MND-related symptoms. GPs will be invited to participate in a 60-minute semi-structured interview to explore the acceptability and usability of the MND Alert toolkit.

What are the possible benefits and risks of participating?

GPs may find it beneficial to participate in a study that aims to develop a toolkit to speed up the diagnosis of MND. There are no risks of participating.

When is the study run from?

Sheffield Institute for Translational Neuroscience (SITraN) (UK)

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

August 2025 to March 2026

Who is funding the study

1. Lifearc (UK)
2. MND Association (UK)

Who is the main contact?

Cara Gates, c.gates@sheffield.ac.uk

## Contact information

### Type(s)

Scientific

### Contact name

Dr Liam Knox

### Contact details

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Sheffield

United Kingdom

S10 2HQ

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L.Knox@sheffield.ac.uk

### Type(s)

Principal investigator

### Contact name

Prof Chris McDermott

### ORCID ID

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School of Medicine and Population Health  
385a Glossop Road  
Sheffield  
United Kingdom  
S10 2HQ  
+44 (0)114222295  
c.j.mcdermott@sheffield.ac.uk

### **Type(s)**

Scientific, Public

### **Contact name**

Dr Cara Gates

### **Contact details**

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School of Medicine and Population Health  
385a Glossop Road  
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United Kingdom  
S10 2HQ  
-  
c.gates@sheffield.ac.uk

## **Additional identifiers**

### **Central Portfolio Management System (CPMS)**

66185

### **Integrated Research Application System (IRAS)**

351043

## **Study information**

### **Scientific Title**

Motor Neuron Disease (MND) Alert: a mixed-method feasibility and acceptability pilot study

### **Acronym**

MND Alert

### **Study objectives**

There are two stages in this study, each with its own research question:

Stage 1: MND Alert implementation

Can the MND Alert toolkit help GPs identify early signs of MND in patients' health records?

Stage 2: MND Alert evaluation

Is the MND alert toolkit useful and acceptable for GPs in their everyday practice?

## **Ethics approval required**

Ethics approval required

## **Ethics approval(s)**

approved 29/05/2025, Health and Care Research Wales (Floor four, North, Welsh Government Offices, Crown Building, King Edward VII Avenue, Cardiff, CF10 3NQ, United Kingdom; +44 (0) 2920230457; HCRW.approvals@wales.nhs.uk), ref: 24/NE/0231

## **Study design**

Non-randomized study

## **Primary study design**

Interventional

## **Study type(s)**

Treatment

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

Motor neurone disease

## **Interventions**

We will conduct a mixed-methods study to test the MND Alert toolkit in up to 30 GP practices. The aim is to see if this automated alert system can help identify potential cases of MND in patient records over a 6-month period. We want to find out if using the alert system and assessment tools can help reduce delays in diagnosing MND.

Once the toolkit is in place, it will trigger an alert when it detects signs or symptoms related to MND in a patient's record. The toolkit will generate reports that include both pseudonymised and identifiable data; however, the identifiable data will only be used within the GP practices for verification. After this verification, GP practices will share only the pseudonymised data with the central research team through secure emails. This data will include unique row numbers that correspond to individual patients, which researchers will use to access patient records during case note reviews.

## **Intervention Type**

Other

## **Primary outcome(s)**

1. Acceptability of the MND Alert measured using thematic analysis of qualitative interviews at after at least 6 weeks of using the toolkit

## **Key secondary outcome(s)**

There are no secondary outcome measures

## **Completion date**

31/03/2026

## **Eligibility**

### **Key inclusion criteria**

## Stage 1: MND Alert implementation

### GP practices:

1. Located in South Yorkshire
2. Willing to integrate the MND Alert toolkit into their electronic health records
3. Using either the SystmOne or EMIS Web clinical information systems
4. Consenting to share pseudo-anonymised patient data generated through the implementation of the MND Alert toolkit
5. Willing to provide researchers access to complete patient records for conducting case note reviews of patients who triggered an alert
6. Willing and able to regularly provide feedback and share monthly pseudo-anonymised data generated through the MND Alert toolkit with the central research team

## Stage 2: MND Alert evaluation

1. English-speaking individuals who can provide informed consent
2. GPs from participating practices
3. GPs willing to participate in semi-structured interviews to share their experiences with the MND Alert toolkit
4. Individuals with access to a telephone or video-calling service (if semi-structured interviews are conducted remotely)

### **Participant type(s)**

Patient

### **Healthy volunteers allowed**

No

### **Age group**

Adult

### **Sex**

All

### **Total final enrolment**

0

### **Key exclusion criteria**

#### Stage 1: MND Alert implementation

##### GP practices:

1. Located outside of the NHS South Yorkshire ICB region
2. Unwilling or unable to integrate the MND Alert toolkit into their electronic health records
3. Not using either the SystmOne or EMIS Web clinical information systems
4. Not consenting to share pseudo-anonymised patient data generated through the implementation of the MND Alert toolkit
5. Unwilling or unable to provide researchers access to complete patient records for conducting case note reviews of patients who triggered an alert
6. Unwilling or unable to regularly provide feedback and share monthly data generated through the MND Alert toolkit with the central research team

#### Stage 2: MND Alert evaluation

1. GPs unwilling to participate in semi-structured interviews to share their experiences with the MND Alert toolkit

**Date of first enrolment**

20/08/2025

**Date of final enrolment**

30/03/2026

## Locations

**Countries of recruitment**

United Kingdom

England

**Study participating centre****Broom Lane Medical Centre**

70 Broom Lane

Broom

Rotherham

England

S60 3EW

**Study participating centre****Buchanan Road Surgery**

72 Buchanan Road

Sheffield

England

S5 8AL

**Study participating centre****Clifton Medical Centre**

Doncaster Gate

Rotherham

England

S65 1DA

**Study participating centre****Clover City Practice**

1 Mulberry Street

Sheffield

England

S1 2PJ

**Study participating centre**  
**Clover Group Practice**  
Darnall Primary Care Centre  
290 Main Road  
Sheffield  
England  
S9 4QH

**Study participating centre**  
**Highgate Surgery**  
Highgate  
Sheffield  
England  
S9 1WN

**Study participating centre**  
**Conisbrough Group Practice**  
The Stone Castle Centre  
Gardens Lane  
Conisbrough  
Doncaster  
England  
DN12 3JW

**Study participating centre**  
**Devonshire Green Medical Centre**  
126 Devonshire Street  
Sheffield  
England  
S3 7SF

**Study participating centre**  
**Dono Valley Healthcare - Sprotbrough Site**  
Newton Lane  
Doncaster  
England  
DN5 8DA

**Study participating centre**

**The Dove Valley Pms Practice**  
Worsbrough Healthcare Centre  
Powell Street  
Worsbrough  
Barnsley  
England  
S70 5NZ

**Study participating centre**  
**Dykes Hall Medical Centre**  
156 Dykes Hall Road  
Sheffield  
England  
S6 4GQ

**Study participating centre**  
**Far Lane Medical Centre**  
1 Far Lane  
Sheffield  
England  
S6 4FA

**Study participating centre**  
**Forge Health Group**  
Pitsmoor Surgery  
151 Burngreave Road  
Sheffield  
England  
S3 9DL

**Study participating centre**  
**Gleadless Medical Centre**  
636 Gleadless Road  
Sheffield  
England  
S14 1PQ

**Study participating centre**  
**Greasbrough Medical Centre**  
Munsbrough Rise  
Greasbrough

Rotherham  
England  
S61 4RB

**Study participating centre**  
**Greystones Medical Centre**  
33 Greystones Road  
Sheffield  
England  
S11 7BJ

**Study participating centre**  
**Harold Street Medical Centre**  
2 Harold Street  
Sheffield  
England  
S6 3QW

**Study participating centre**  
**Heeley Green Surgery**  
302 Gleadless Road  
Sheffield  
England  
S2 3AJ

**Study participating centre**  
**Clover North Darnall Health Centre**  
2 York Road, Darnall  
Sheffield  
England  
S9 5DH

**Study participating centre**  
**Oughtibridge Surgery**  
Church Street  
Oughtibridge  
Sheffield  
England  
S35 0FW

**Study participating centre**  
**Sothall & Beighton Health Centres**  
24 Eckington Road  
Beighton  
Sheffield  
England  
S20 1HQ

**Study participating centre**  
**Swallownest Health Centre**  
Worksop Road  
Swallownest  
Sheffield  
England  
S26 4WD

**Study participating centre**  
**The Crookes Practice**  
203-205 School Road  
Sheffield  
England  
S10 1GN

**Study participating centre**  
**The Lakeside Practice**  
The White Wings Centre  
Spa Pool Road  
Askern  
Doncaster  
England  
DN6 0HZ

**Study participating centre**  
**The Thorpe Practice**  
Moorthorpe Bank  
Owlthorpe  
Sheffield  
England  
S20 6PD

**Study participating centre****Woodhouse Health Centre (pcn Sheffield) - COVID Local Vaccination Service**

Woodhouse Health Centre  
5-9 Skelton Lane  
Woodhouse  
Sheffield  
England  
S13 7LY

**Study participating centre****Woodseats Medical Centre**

The Roddick Building  
900 Chesterfield Road  
Sheffield  
England  
S8 0SH

**Study participating centre****Woodstock Bower Group Practice**

1 Kimberworth Road  
Rotherham  
England  
S61 1AH

## Sponsor information

**Organisation**

University of Sheffield

**ROR**

<https://ror.org/05krs5044>

## Funder(s)

**Funder type**

Charity

**Funder Name**

Motor Neurone Disease Association

**Alternative Name(s)**

MND Association, Motor neurone disease (MND), Mndassos, MNDA

**Funding Body Type**

Private sector organisation

**Funding Body Subtype**

Associations and societies (private and public)

**Location**

United Kingdom

**Funder Name**

LifeArc

**Alternative Name(s)****Funding Body Type**

Private sector organisation

**Funding Body Subtype**

Other non-profit organizations

**Location**

United Kingdom

## Results and Publications

**Individual participant data (IPD) sharing plan**

The datasets generated during and/or analysed during the current study will be available upon request from Dr Liam Knox (L.knox@sheffield.ac.uk).

Type of data: anonymised interview data or analysis of interviews, practice-level based on Alert Tool usage, suggest qualitative patient-level data from case notes is neither appropriate nor useful to be made available.

When the data will be available and for how long: Anonymised data will be kept for at least five years and will be available after the completion of the study.

For what types of analyses: secondary analysis of interviews.

Interview participant consent will be taken for data sharing (when anonymised).

Consent not obtained from patients to access records and view consultation details (this was approved via CAG), therefore it is not appropriate to share this data.

**IPD sharing plan summary**

Available on request