

Continuous nursing at home for well-being after stomach cancer surgery

Submission date 31/10/2025	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 25/11/2025	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 05/11/2025	Condition category Cancer	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

Gastric cancer (GC) is the fifth most common malignant tumor in the world in terms of incidence rate and the second most common malignant tumor in terms of mortality. The postoperative condition of patients with GC metastasis is still poor. How to effectively improve the quality of life (QoL) of postoperative patients has gradually become the key target of current treatment. This study aimed to explore the impact of continuous nursing care on the QoL of GC lung metastasis patients, to better implement treatment for GC patients and improve their QoL.

Who can participate?

Patients who have a confirmed diagnosis of stomach cancer that has spread to the lungs.

What does the study involve?

This study will compare two interventions: routine nursing care, which involves regular telephone follow-ups and continuous nursing care, which involves a structured program of telephone follow-ups, scheduled home visits, priority appointments at a specialist clinic, and support through an online health platform.

These measurements will be taken at the time of hospital discharge, and again at 3 months, 6 months, and 1 year after discharge using a questionnaire.

What are the possible benefits and risks of participating?

By enrolling, participants might gain access to a new and potentially better treatment that isn't available to everyone, all while receiving very close medical care from a team of experts. The research team will explain all the known risks, like tiredness or disturbance, in detail.

Where is the study run from?

The First Hospital of China Medical University, Shenyang, China.

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

June 2022 to October 2025

Who is funding the study?
Investigator initiated and funded

Who is the main contact?
Yueyang Jiang, yueyang_jiang@hotmail.com

Contact information

Type(s)

Public, Scientific, Principal investigator

Contact name

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Additional identifiers

Study information

Scientific Title

Effect of continuous nursing care on postoperative quality of life in patients with gastric cancer lung metastasis

Study objectives

This study aimed to explore the impact of continuous nursing care on the quality of life (QoL) of gastric cancer (GC) lung metastasis patients, to better implement treatment for GC patients and improve their QoL.

Ethics approval required

Ethics approval required

Ethics approval(s)

approved 17/07/2025, Ethics Committee of the First Hospital of China Medical University (No. 155 Nanjing North Street, Heping District, Shenyang City, 110001, China; -; ethics@cmu1h.com), ref: 101022119

Study design

Randomized controlled trial

Primary study design

Interventional

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

Gastric cancer

Interventions

This was a 1-year follow-up study where 102 patients with GC lung metastasis in the First Hospital of China Medical University were randomized to routine telephone follow-ups (Control) or a comprehensive continuous nursing program involving phone calls, home visits, clinic services, and online support (Observation). The anxiety, depression and QoL scores of patients: physical function, role function, emotional function, cognitive function, and social function, were observed at the time of discharge, 3 months, 6 months and 1 year after discharge.

Control Group Protocol: the patients received regular telephone follow-ups conducted by a nurse specialist, which included: medication guidance, dietary counseling and mental health monitoring. All follow-up records were documented in a discharge follow-up logbook.

The randomization process was designed to ensure robustness and minimize bias, beginning with a computer-generated random number sequence created by an independent statistician uninvolved in patient care. Assignment occurred only after patients had provided informed consent and their eligibility was definitively confirmed. To enforce allocation concealment, sequentially numbered, opaque, sealed envelopes were used, and a research nurse opened each envelope solely after a participant's baseline assessment was complete, thereby ensuring the enrolling team remained blinded to the upcoming group assignment throughout implementation.

Intervention Type

Behavioural

Primary outcome(s)

The following primary outcome measures were assessed at discharge, 3 and 6 months, and 1 year after discharge:

1. Anxiety measured using a simplified version of the General Anxiety Disorder-7 (GAD-7) questionnaire
2. Depression measured using a simplified version of the Patient Health Questionnaire-9 (PHQ-9)

Key secondary outcome(s)

There are no secondary outcome measures

Completion date

31/10/2025

Eligibility

Key inclusion criteria

1. Pathologically confirmed diagnosis of gastric cancer with lung metastasis
2. Possessing self-awareness and ability to cooperate in answering questions
3. Patients who showed clinical improvement after surgical treatment

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

40 years

Upper age limit

80 years

Sex

All

Total final enrolment

102

Key exclusion criteria

1. Comorbidities with other severe systemic diseases
2. Patients with cognitive dysfunction
3. Patients unwilling to cooperate with home visits
4. Other reasons preventing participation

Date of first enrolment

01/06/2022

Date of final enrolment

01/06/2025

Locations**Countries of recruitment**

China

Study participating centre

First Hospital of China Medical University

No. 155 Nanjing North Street

Heping District

Shenyang

China

110001

Sponsor information

Organisation

First Hospital of China Medical University

ROR

<https://ror.org/04wjghj95>

Funder(s)**Funder type**

Other

Funder Name

Investigator initiated and funded

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 Mrs Yueyang Jiang (Yueyang_jiang@hotmail.com)

IPD sharing plan summary

Available on request