

Subserosal carbon nanoparticles for gastrectomy

Submission date 29/07/2023	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 02/08/2023	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 21/05/2025	Condition category Cancer	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Lymph node metastasis (tumour spreading into lymph nodes) is common in gastric (stomach) cancer. Extensive lymph node dissection is the fundamental step in radical gastrectomy (complete or near-complete removal of the stomach). Improving the visualization of lymph nodes can help surgeons identify and dissect lymph nodes during surgery. Carbon nanoparticles are an approved drug for visualization. However, it is unclear which approach of administration is best (subserosal injection or submucosal injection, i.e. injected under different layers of tissue). This study is designed to compare the effectiveness and safety of these two approaches in radical gastrectomy.

Who can participate?

Patients between 18 and 75 years old with gastric cancer

What does the study involve?

This study involves radical gastrectomy and the administration of carbon nanoparticles. Participants will be randomly allocated into two groups: Group 1: Subserosal injection of carbon nanoparticles for lymph node visualization. Group 2: Submucosal injection of carbon nanoparticles for lymph node visualization.

What are the possible benefits and risks of participating?

Participants may benefit from an increased number of retrieved lymph nodes from the surgery, which will help with the precise staging of the tumour. Any risks related to the surgery itself (such as bleeding, obstruction, or anastomotic leakage) might occur during this study. There is a very limited risk of participating since this is a safe and approved drug for surgery.

Where is the study run from?

Nanjing Drum Tower Hospital, the Affiliated Hospital of Nanjing University (China)

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

September 2022 to December 2025

Who is funding the study?
National Natural Science Foundation of China

Who is the main contact?
Dr Song Liu, liusong@njglyy.com

Contact information

Type(s)

Principal investigator

Contact name

Dr Song Liu

ORCID ID

<https://orcid.org/0000-0002-4780-9697>

Contact details

321 Zhongshan RD
Nanjing
China
210008
+86 (0)25 83106666 60931
liusong@njglyy.com

Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

DANCE-04

Study information

Scientific Title

Efficacy and safety of subserosal carbon nanoparticles navigated radical gastrectomy

Acronym

DANCE-04

Study objectives

Subserosal carbon nanoparticles is safer and more efficient than submucosal carbon nanoparticles in radical gastrectomy

Ethics approval required

Ethics approval required

Ethics approval(s)

approved 08/12/2022, Ethics committee of Nanjing Drum Tower Hospital, the Affiliated Hospital of Nanjing University Medical School (321 Zhongshan Rd, Nanjing, 210008, China; +86 (0)25 83106666; irb_dth@outlook.com), ref: 2023-372-02

Study design

Randomized interventional prospective single-center clinical trial

Primary study design

Interventional

Study type(s)

Treatment, Safety, Efficacy

Health condition(s) or problem(s) studied

Radical gastrectomy for patients with gastric cancer

Interventions

All participants will be randomized into two groups. Group 1: Subserosal injection of carbon nanoparticles for lymph node visualization. Group 2: Submucosal injection of carbon nanoparticles for lymph node visualization.

Method of randomization: simple randomization method using a computer-generated random number to assign each participant to a specific group.

Intervention Type

Procedure/Surgery

Primary outcome(s)

Number of retrieved lymph nodes measured by pathological examination immediately after radical gastrectomy

Key secondary outcome(s)

1. Number of metastatic lymph nodes measured by pathological examination immediately after radical gastrectomy
2. Sensitivity and specificity of carbon nanoparticles staining for the diagnosis of metastatic lymph nodes, measured by pathological examination immediately after radical gastrectomy
3. 3-year disease-free survival (DFS), 3-year relapse-free survival (RFS), 3-year overall survival (OS), measured by outpatient clinic or telephone follow-up at 3 years after surgery
4. Postoperative complications measured by Clavien-Dindo classification at 1, 3, and 7 days after surgery

Completion date

31/12/2025

Eligibility

Key inclusion criteria

1. Between 18 and 75 years old
2. cT1-4a, cNany, cM0 according to the AJCC classification (version 7.0) before surgery
3. Absence of adjacent organ invasion
4. Performance status (ECOG score) 0 or 1, ASA I-III
5. Consent of participation
6. Distal or total radical gastrectomy

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

75 years

Sex

All

Total final enrolment

99

Key exclusion criteria

1. Pregnancy or lactation period
2. Severe mental illness
3. Previous upper abdomen surgery excluding laparoscopic cholecystectomy
4. Previous gastric ESD or EMR history
5. Refusal to laparoscopy
6. Previous iodine allergy
7. Peri-gastric LNs larger than 3cm in preoperative CT or MR
8. Other malignancy in recent 5 years
9. Neoadjuvant therapy for gastric cancer
10. Unstable angina pectoris or myocardial infarction in recent 6 months
11. FEV1 less than 50% of predicted value
12. Systemic corticosteroid therapy in recent 1 month
13. Emergent surgery due to acute bleeding, perforation or obstruction
14. Convert to open surgery
15. Concomitant surgery excluding appendectomy or cholecystectomy

Date of first enrolment

01/08/2023

Date of final enrolment

31/12/2024

Locations

Countries of recruitment

China

Study participating centre

Nanjing Drum Tower Hospital, the Affiliated Hospital of Nanjing University Medical School

321 Zhongshan Rd

Nanjing

China

210008

Sponsor information

Organisation

Nanjing Drum Tower Hospital

ROR

<https://ror.org/026axqv54>

Funder(s)

Funder type

Government

Funder Name

National Natural Science Foundation of China

Alternative Name(s)

Chinese National Science Foundation, Natural Science Foundation of China, National Science Foundation of China, NNSF of China, NSF of China, National Nature Science Foundation of China, Guójiā Zìrán Kēxué Jījīn Wěiyuánhùi, , NSFC, NNSF, NNSFC

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

China

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated and/or analysed during this study will be published as a supplement to the results publication.

IPD sharing plan summary

Published as a supplement to the results publication

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article		19/05/2025	21/05/2025	Yes	No