

Observation of the effects of predictive nursing in patients with HIV/AIDS undergoing anal fistula surgery

Submission date 24/10/2024	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 06/11/2024	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 05/11/2024	Condition category Other	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

Constipation caused by fear of defecation is a common complication of anal fistula surgery. HIV infection is common in patients with a variety of anorectal diseases, postoperative prognosis is poorer, and complication rates are relatively high. Predictive nursing, as one of the new clinical nursing models, can achieve the purpose of reducing postoperative complications by the comprehensive evaluation of patients by medical staff.

Who can participate?

Patients with HIV/AIDS aged 18 years and older undergoing surgical treatment of complex anal fistula.

What does the study involve?

The patients were randomly assigned to two groups. No patients had acute symptoms on presentation, and all underwent a series of standardised clinical assessments, including ultrasound, digital rectal examination and colonoscopy, to confirm the absence of contraindications to the surgical procedures used in this study; the types of surgery included cutting or non-cutting seton, ligation of the intersphincteric fistula tract, anorectal advancement flap, fibrin glue injection, stem cell treatment and laser treatment. After admission, patients in the control group received routine nursing, and those in the intervention group received predictive nursing based on routine nursing.

What are the possible benefits and risks of participating?

The ability of patients to care for themselves may improve, and the incidence of postoperative constipation may decrease. There are no direct risks to participants.

Where is the study run from?

Beijing Youan Hospital, Capital Medical University (China)

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

January 2021 to January 2023

Who is funding the study?

1. Beijing Hospitals Authority Youth Programme
2. Beijing You'an Hospital affiliated with Capital Medical University, 2021 Hospital Young and Middle-aged Talents Incubation Project (China).

Who is the main contact?

Li-Li Zhang, zhanglili@yeah.net

Contact information

Type(s)

Public, Scientific, Principal investigator

Contact name

Dr Li-Li Zhang

ORCID ID

<https://orcid.org/0009-0004-3610-9152>

Contact details

Department of Nursing, Beijing Youan Hospital, Capital Medical University
No.8, Xi Tou Tiao, Youanmen wai, Fengtai District

Beijing

China

100069

+86 10 8399 7143

zhanglili@yeah.net

Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

Protocol serial number

Nil known

Study information

Scientific Title

Observation of the effects of predictive nursing on postoperative constipation prevention and self-care ability in patients with HIV/AIDS undergoing anal fistula surgery

Study objectives

To explore the effects of predictive nursing on the prevention of postoperative constipation and self-care ability in patients with HIV/AIDS undergoing anal fistula surgery.

Ethics approval required

Ethics approval required

Ethics approval(s)

approved 07/12/2021, Ethics Committee of Beijing Youan Hospital, Capital Medical University (No.8, Xi Tou Tiao, Youanmen wai, Fengtai District, Beijing, 100069, China; +86 010-83997028; youanyuanban@163.com), ref: LL-2021-160-K

Study design

Randomized clinical trial

Primary study design

Interventional

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

Constipation prevention, self-care ability

Interventions

The patients were randomly assigned to two groups using block randomisation, with 100 patients in each group. The experimental group received predictive nursing based on routine nursing, which includes: risk assessment, diet care, psychological care, bowel care, and anal function exercise. Whereas the control group received routine nursing, including perioperative routine health education guidance.

Control group: routine nursing.

The whole intervention period lasted for 7 days. The main contents are perioperative routine health education guidance, including preoperative, intraoperative and postoperative health education guidance, psychological nursing, and medication nursing. These measures were designed to assist patients in handling psychological stress and emotional issues, thereby enhancing treatment compliance.

Experimental group: predictive nursing based on routine nursing.

The experimental group received predictive nursing based on routine nursing, which includes: risk assessment, diet care, psychological care, bowel care, and anal function exercise. The intervention period was 7 days.

Follow-up time point: The follow-up was performed on the 7th day after discharge. Follow-up included an assessment of the patient's constipation. For patients with constipation, the focus was on whether there is still constipation to check the recovery of the surgical site for complications, such as infection and adhesion. At the same time, the patients were asked about their quality of life and psychological status at each follow-up visit to understand the impact of the treatment on their overall life.

Follow-up: Outpatient review, telephone interview and online questionnaire were used to ensure a comprehensive collection of patient information.

Intervention Type

Behavioural

Primary outcome(s)

1. The effectiveness of constipation prevention was measured using outpatient review, telephone interview and online questionnaire before and after the intervention, according to the Consensus on Traditional Chinese Medicine Diagnosis and Treatment of Constipation (2017) as follows: 1) ineffective: bowel movements less than 3 times per week with hard stools, difficult defaecation or a feeling of incomplete evacuation; 2) effective: bowel movements more than once within 3 days; and 3) highly effective: bowel movements more than once within 2 days. The effective rate of constipation prevention (%) was calculated by dividing the sum of highly effective and effective cases by the total number of cases and multiplying by 100.
2. Self-care ability measured using the Exercise of Self-Care Agency (ESCA) scale before and after the intervention

Key secondary outcome(s)

The following secondary outcome measures are assessed using data recorded in patient medical records:

1. Preoperative history of constipation measured by asking the patient whether he has had constipation in the past month before surgery
2. Anal fistula duration
3. Operative time
4. Blood loss
5. Type of surgery
6. CD4 cell counts / viral loads measured using standard laboratory methods and a preoperative blood test
7. Bowel disorders and type of anal fistula

Completion date

31/01/2023

Eligibility

Key inclusion criteria

1. A confirmed HIV infection / AIDS diagnosis according to the 2021 edition of the Chinese Guidelines for the Diagnosis and Treatment of HIV/AIDS, having received antiretroviral therapy and with a current CD4+ T-cell count of ≥ 200 cells/ μ L
2. Aged ≥ 18 years
3. Ability to cooperate with the researchers and complete the study-related procedures and data collection
4. Voluntary participation in the study and signing of an informed consent form

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

All

Total final enrolment

200

Key exclusion criteria

1. Presence of other anal or colorectal diseases, such as perianal dermatitis or colorectal tumours
2. Recurrent transsphincteric anal fistula
3. Presence of severe diseases such as cardiac, hepatic or renal conditions
4. History of previous anal surgery
5. Presence of mental or psychological disorders

Date of first enrolment

01/01/2022

Date of final enrolment

31/01/2023

Locations**Countries of recruitment**

China

Study participating centre

Beijing Youan Hospital, Capital Medical University

No.8, Xi Tou Tiao, Youanmen wai, Fengtai District

Beijing

China

100069

Sponsor information**Organisation**

Beijing YouAn Hospital

ROR

<https://ror.org/04etaja30>

Funder(s)**Funder type**

Hospital/treatment centre

Funder Name

Beijing Hospitals Authority Youth Programme

Funder Name

Beijing You'an Hospital affiliated to Capital Medical University, 2021 Hospital Young and middle-aged Talents Incubation Project

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 Li-Li Zhang, Email: zzhaglili@yeah.net

IPD sharing plan summary

Available on request