# Anal fistula plug versus endorectal advancement flap for the closure of high criptoglandular fistula-in-ano: a randomised study

Submission date	Recruitment status	Prospectively registered
29/10/2008	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
20/11/2008	Completed	Results
Last Edited	Condition category	[] Individual participant data
20/11/2008	Digestive System	<ul><li>Record updated in last year</li></ul>

# Plain English summary of protocol

Not provided at time of registration

# Contact information

## Type(s)

Scientific

#### Contact name

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# Additional identifiers

**EudraCT/CTIS** number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

# Study information

## Scientific Title

A randomised study to compare the results of the anal fistula plug (AFP) with the endorectal advancement flap (ERAF) in the treatment of high fistula-in-ano of cryptoglandular origin

## **Study objectives**

The use of lyophilised porcine submucosal plugs (Cook Surgisis®, AFPTM) has been proposed as an alternative to conventional surgical techniques for the treatment of anal fistulas. Rates of favourable outcomes are highly variable in the literature (between 13.9% and 87%). Unfortunately, some reports are retrospective studies, others are prospective cohort studies, and only one study has compared the efficacy of this technique with a retrospective review of patients treated with endorectal advancement flap (ERAF). Additionally, these studies include simple and complex anal fistulas, anovaginal fistulas and patients with inflammatory bowel disease. So far, there is not a randomised study comparing the AFP with other surgical procedures suitable for high fistulas. Therefore, the objective of this randomised study was to compare the results of AFP with ERAF in the treatment of high fistula-in-ano of cryptoglandular origin.

## Ethics approval required

Old ethics approval format

## Ethics approval(s)

The Comité Ético de Investigación Clínica of the Health Department of the Government of Navarra (Spain) gave approval on the 2nd April 2007 (ref: Pyto. 14\_07).

## Study design

Interventional randomised single-centre trial

## Primary study design

Interventional

## Secondary study design

Randomised controlled trial

## Study setting(s)

Hospital

## Study type(s)

Treatment

## Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

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

Fistula-in-ano of cryptoglandular origin

## **Interventions**

All patients underwent full mechanical bowel preparation and received antibiotic and antithromboembolic prophylaxis. The patients were operated on under general anaesthesia in the lithotomy position. Surgical procedures were performed by two surgeons with accredited degrees in Coloproctology (ESBQ Coloproctology).

## Anal fistula plug (AFP):

The plug was submerged in saline for two minutes; the internal fistula orifice was located by injecting hydrogen peroxide. Curettage of the track was not performed. A probe was inserted in the fistula track. The AFP was placed into the tract until resistance was felt and then fixed in place with a 2-0 polyglactin suture (Vicryl®) which included the internal sphincter. The suture was tied in order to close the internal opening of the fistula over the plug. Care was taken to ensure that the external orifice of the fistula was not completely occluded in order to allow the track to drain. The remaining plug was cut at the level of the external opening.

## Endorectal advancement flap:

The tract of the fistula was completely excised, including the internal opening. A rectal flap above the internal opening was mobilised, including the mucosa and submucosa, with a 3 to 4 cm broad base. The rectal flap was mobilised sufficiently to cover the internal opening. Exhaustive haemostasis was performed to avoid a haematoma under the flap. Finally, the flap was sutured to the edge of the anal canal, covering the internal opening.

Average duration of treatment:
Anal Fistula Plug: 30 minutes
Endorectal Advancement Flap: 90 minutes
Total duration of follow-up for all arms of your trial: one year

## Intervention Type

Other

#### Phase

**Not Specified** 

## Primary outcome measure

Fistula closure rate; recurrence is defined as the presence of an abscess arising in the area, or by obvious evidence of fistulation. Evaluated at 2, 4, 8, 12 weeks after surgery, and at 6, 9 and 12 months.

#### Secondary outcome measures

Continence, evaluated pre- and post-operatively (after one year) using the Wexner score.

## Overall study start date

01/05/2007

## Completion date

30/04/2009

# **Eligibility**

# Key inclusion criteria

- 1. Aged above 18 years, either sex
- 2. High fistula-in-ano of cryptoglandular origin (the fistulas were defined as high when they

included the upper two-thirds of the external sphincter complex)
3. Informed consent

# Participant type(s)

**Patient** 

## Age group

Adult

## Lower age limit

18 Years

## Sex

Both

## Target number of participants

186 patients (93 in each group)

## Key exclusion criteria

- 1. Secondary tracts
- 2. Horseshoe fistulas
- 3. Anovaginal fistulas
- 4. Rectouretral fistulas
- 5. Human immunodeficiency virus (HIV)-positive patients
- 6. Diagnosed from Crohn's disease

## Date of first enrolment

01/05/2007

## Date of final enrolment

30/04/2009

# Locations

## Countries of recruitment

Spain

# Study participating centre C/Trinidad Fernandez Arenas

Pamplona Spain 31002

# Sponsor information

## Organisation

Hospital Virgen del Camino (Spain) - Public University of Navarra

## Sponsor details

C/Irunlarrea 4 Pamplona Spain 31008

## Sponsor type

Hospital/treatment centre

## **ROR**

https://ror.org/046wwv897

# Funder(s)

## Funder type

Hospital/treatment centre

## **Funder Name**

Hospital Virgen del Camino (Spain) - Public University of Navarra

# **Results and Publications**

## Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

# IPD sharing plan summary

Not provided at time of registration