A randomized trial comparing the CORTRAK system with the endoscopic technique for duodenal feeding tube placement

Published: 02-04-2013 Last updated: 24-04-2024

Compare the success rate of duodenal feeding tube placement using the CORTRAK system (DFT-C) with the endoscopic technique (DFT-E).

Ethical review Approved WMO **Status** Recruitment stopped

Health condition type Gastrointestinal conditions NEC

Study type Interventional

Summary

ID

NL-OMON41400

Source

ToetsingOnline

Brief title

CORRECT study

Condition

Gastrointestinal conditions NEC

Synonym

duodenul feeding tube placement; feeding tube placement

Research involving

Human

Sponsors and support

Primary sponsor: Universitair Medisch Centrum Utrecht

Source(s) of monetary or material Support: CORPAK MedSystems, Inc, Illinois,

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USA.subsidie van CORPAK

Intervention

Keyword: COTRAK, DFT, duodenal feeding tube, endoscopy

Outcome measures

Primary outcome

Success rate of DFT placement; defined as the tip of the tube placed postpyloric and into the duodenum as confirmed by an abdominal X-ray.

Secondary outcome

Costs associated with DFT placement, procedure time, difficulty of the procedure, accuracy of tip location with CORTRAK, DFT tip location, safety, use of sedatives, reintervention rate within 10 days and patient acceptance (in unsedated patients).

Study description

Background summary

Increasing evidence confirms the important role of enteral feeding in (critically ill) patients. A substantial part of these patients have an indication for duodenal or jejunal feeding. However, duodenal feeding tube (DFT) placement can be difficult, time-consuming, or costly, depending on the technique used. Endoscopic tube placement has high success rates, but the availability of appropriate staff and specialized equipment is required for this technique, making this technique costly and difficult to provide consistently. Several other techniques for DFT placement have been assessed, all require a high level of expertise and most have not been compared with the endoscopic technique. An electromagnetic tube placement device, the CORTRAK system, is increasing in popularity and several observational studies have demonstrated this technique to be safe, efficient, and cost-effective. Only two small studies have compared the CORTRAK system with other placement techniques in a controlled setting.

Study objective

Compare the success rate of duodenal feeding tube placement using the CORTRAK system (DFT-C) with the endoscopic technique (DFT-E).

Study design

Randomized sequential prospective, multicenter, non-blind, controlled trial.

Intervention

Patients will be randomized to undergo DFT placement with the CORTRAK system or the endoscopic technique.

Study burden and risks

The risk for complications when performing DFT placement with the CORTRAK system is probably lower since fewer sedatives will be used. Possible complications are epistaxis, inadvertent passage of the feeding tube into the respiratory tract, aspiration and unsuccessful placement of the feeding tube. There are no guaranteed benefits of participation for the subject, though the burden of participating is negligible. Possible benefits of DFT placement with the CORTRAK system are less need for sedation and less complications due to sedation, cost reduction due to savings on staff and equipment and a simplified logistics that might result in more prompt initiation of enteral feeding with all the benefits of early start of enteral feeding.

Contacts

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years) Elderly (65 years and older)

Inclusion criteria

- All patients needing a duodenal feeding tube
- Written informed consent provided by patient or representative
- *18 years

Exclusion criteria

- Implantable pacing devices (potential interference with the signal transmission)
- Altered anatomy of the upper gastrointestinal tract due to surgery of the esophagus, stomach or duodenum
- High suspicion of stenosis or obstruction in the upper digestive tract
- Esophageal varices
- Signs of active upper gastrointestinal bleeding.
- Woman with known pregnancy (because of abdominal X-rays performed in order to confirm location of the DFT)

Study design

Design

Study phase: 3

Study type: Interventional

Intervention model: Parallel

Allocation: Randomized controlled trial

Masking: Open (masking not used)

Control: Active

Primary purpose: Treatment

Recruitment

NL

Recruitment status: Recruitment stopped

Start date (anticipated): 09-12-2013

Enrollment: 309

Type: Actual

Medical products/devices used

Generic name: CORTRAK system

Registration: Yes - CE intended use

Ethics review

Approved WMO

Date: 02-04-2013

Application type: First submission

Review commission: METC Universitair Medisch Centrum Utrecht (Utrecht)

Approved WMO

Date: 03-09-2014

Application type: Amendment

Review commission: METC Universitair Medisch Centrum Utrecht (Utrecht)

Approved WMO

Date: 04-06-2015

Application type: Amendment

Review commission: METC Universitair Medisch Centrum Utrecht (Utrecht)

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register ID

CCMO NL42753.041.12