The role of Ketotofen in the treatment of postoperative ileus.

No registrations found.

Ethical review	Positive opinion
Status	Recruitment stopped
Health condition type	-
Study type	Interventional

Summary

ID

NL-OMON24494

Source
Nationaal Trial Register

Brief title
N/A

Intervention

Outcome measures

Primary outcome

Gastric emptying.

Secondary outcome

Symptoms.

Study description

Background summary

This is a randomized, placebo-controlled, double-blind study with patients who'll undergo

abdominal surgery. They will be treated with Ketotofen 4 mg or 12 mg or placebo per day. The patient will start his medication intake 3 days before surgery till 2 days after surgery. Post operative effectivity will be determined by scintigraphy to look at gastric retention and colontransit. This scintigraphy will take place 24 hours after surgery. The patients will be clinical evaluated by filling out a daily symptom score list.

Study objective

N/A

Study design

N/A

Intervention

Ketotofen 4 mg or 12 mg or placebo per day.

Contacts

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Eligibility criteria

Inclusion criteria

Laparotomy for (malignant) proces originaty from colon or female reproduction organs.

Exclusion criteria

Pre-operative radiation, intake of medication with effect on the Gastrointestinal motility, intake of immunosuppressiva, intake of anti-allergy medication, intra-abdominal inflammation (cholecystitis or absces included).

Study design

Design

Study type:	Interventional
Intervention model:	Parallel
Allocation:	Randomized controlled trial
Masking:	Double blinded (masking used)
Control:	Placebo

Recruitment

NL

Recruitment status:	Recruitment stopped
Start date (anticipated):	01-04-2004
Enrollment:	48
Type:	Actual

Ethics review

Positive opinion

Date:	02-06-2005
Application type:	First submission

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
NTR-new	NL16
NTR-old	NTR38
Other	: N/A
ISRCTN	ISRCTN34927143

Study results

Summary results
N/A