

Gastric emptying and protein absorption of caprine and bovine casein protein

Published: 19-09-2019

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The study aims to quantify differences in gastric digestion and absorption of caprine and bovine casein protein

Ethical review	Approved WMO
Status	Recruitment stopped
Health condition type	Other condition
Study type	Interventional

Summary

ID

NL-OMON48463

Source

ToetsingOnline

Brief title

vision study

Condition

- Other condition

Synonym

normal diagection

Health condition

gezonde deelnemers

Research involving

Human

Sponsors and support

Primary sponsor: Wageningen Universiteit

Source(s) of monetary or material Support: Ausnutria BV

Intervention

Keyword: absorption, gastric emptying, protein digestion

Outcome measures

Primary outcome

gastric emptying t50

amino acids concentration over time in the blood

Secondary outcome

intragastric layering over time

Study description

Background summary

A suggested benefit of caprine (CAP) over bovine milk (BOV) has been hypothesized to be an accelerated gastric emptying of CAP. Due to more κ -casein and less κ s1-casein in caprine milk less coarse coagulates are thought to be formed in the stomach, which may facilitate enzymatic digestion and gastric emptying.

This may provide benefits for specific groups, for example patients who struggle to meet protein intakes as protein. However, thus far, no in vivo clinical data on digestion of caprine versus bovine casein is available.

Study objective

The study aims to quantify differences in gastric digestion and absorption of caprine and bovine casein protein

Study design

Intervention study with a balanced cross-over design

Intervention

300mL vanilla flavoured drink with either 30g of caprine casein or bovine casein

Study burden and risks

The risks associated with participation are negligible, as both phlebotomy and MRI are eminently safe medical techniques, and the stimuli consist of normally consumed food products. The burden associated with participation consists of two visits, which both require an overnight fast, 8 blood draws (10 mL per draw, totalling 80 mL) and multiple MRI scans over the period of 1 hours. These may all cause minimal discomfort. There is no benefit to participation for the participants, the group is only related insofar as they are healthy males.

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

- * Male
- * 18 - 55 yr old
- * In self-reported health
- * BMI between 18 and 25kg/m²

Exclusion criteria

- * Bovine milk allergy or intolerance (self-reported)
- * Caprine milk allergy or intolerance (self-reported)
- * Lactose intolerance (self-reported)
- * Disorders of the upper ingestive tract resulting in difficulties chewing/swallowing.
- * Unexplained weight change
- * Gastric disorders or regular gastric complaints, heart burn for example
- * Use of proton pump inhibitors or other medication which alters the normal function-ing of the stomach
- * Smoking
- * Having a contra-indication to MRI scanning (including, but not limited to):
- * Pacemakers and defibrillators
- * Intraorbital or intraocular metallic fragments
- * Ferromagnetic implants
- * Claustrophobia

Study design

Design

Study type:	Interventional
Intervention model:	Crossover
Masking:	Single blinded (masking used)
Control:	Uncontrolled
Primary purpose:	Other

Recruitment

NL	
Recruitment status:	Recruitment stopped
Start date (anticipated):	22-11-2019
Enrollment:	18

Type:

Actual

Ethics review

Approved WMO

Date: 19-09-2019

Application type: First submission

Review commission: METC Wageningen Universiteit (Wageningen)

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

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

ID: 28580

Source: NTR

Title:

In other registers

Register	ID
CCMO	NL70239.081.19
Other	nog geen nummer toegewezen
OMON	NL-OMON28580