

# Beer study

No registrations found.

<b>Ethical review</b>	Not applicable
<b>Status</b>	Recruiting
<b>Health condition type</b>	-
<b>Study type</b>	Interventional

## Summary

### ID

NL-OMON29218

### Source

Nationaal Trial Register

### Health condition

Neural responses

Beer

Reward

## Sponsors and support

### Primary sponsor: Wageningen University

Division of Human Nutrition (Bode 62)

PO Box 8129

6700 EV Wageningen

Phone: 0317 - 485302

Kennisinstituut Bier

PO Box 590

6700 AN Wageningen

Phone: 0317 - 750620

### Source(s) of monetary or material Support: Wageningen University

Division of Human Nutrition (Bode 62)

PO Box 8129

6700 EV Wageningen

Phone: 0317 - 485302

Kennisinstituut Bier

PO Box 590

6700 AN Wageningen

Phone: 0317 - 750620

## **Intervention**

## **Outcome measures**

### **Primary outcome**

Classified information

### **Secondary outcome**

-

## **Study description**

### **Background summary**

classified information

### **Study objective**

Classified information

### **Study design**

-

### **Intervention**

The study has a randomized crossover design (within subject design) in which participants taste a fixed amount of beer during an fMRI scan session, in a setting where beer is expected.

## **Contacts**

### **Public**

Paul A.M. Smeets  
Division of Human Nutrition

Bomenweg 2  
Biotechnion (307), room 323  
Wageningen 6703 HD  
The Netherlands  
0317 – 484681

### **Scientific**

Paul A.M. Smeets  
Division of Human Nutrition  
Bomenweg 2  
Biotechnion (307), room 323  
Wageningen 6703 HD  
The Netherlands  
0317 – 484681

## **Eligibility criteria**

### **Inclusion criteria**

Age: 18-35 years

- BMI: 20 – 25 kg/m<sup>2</sup>
- Healthy (as judged by the participant)
- Being right handed
- Used to drinking beer
- Willing to comply with the study procedures
- Willing to be informed about incidental findings of pathology
- Having given written informed consent
- Successful completion of the training session
- Successful completion of the sensory test

### **Exclusion criteria**

- Restraint eating (men: DEBQ score > 2.25)

- Lack of appetite
- Having difficulties with swallowing/eating
- Usage of an energy restricted diet during the last two months
- Weight loss or weight gain of 5 kg or more during the last two months
- Stomach or bowel diseases
- Diabetes, thyroid disease, other endocrine disorders
- Having a history of neurological disorders
- Having taste or smell disorders
- Usage of daily medication other Paracetamol
- Smoking more than one cigarette/cigar a day
- Being allergic/intolerant for products under study
- Working at the Division of Human Nutrition (WUR)
- Current participation in other research from the Division of Human Nutrition (WUR)
- Drinking on average more than 14 alcoholic beverages a week
- Family history of alcoholism (loss of control, tolerance or withdrawal symptoms towards alcohol in direct family)

## Study design

### Design

Study type:	Interventional
Intervention model:	Crossover
Allocation:	N/A: single arm study
Masking:	Single blinded (masking used)
Control:	N/A , unknown

## Recruitment

NL  
Recruitment status: Recruiting  
Start date (anticipated): 01-04-2014  
Enrollment: 20  
Type: Anticipated

## Ethics review

Not applicable  
Application type: Not applicable

## 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	NL4438
NTR-old	NTR4560
Other	13/28 : METC

## Study results