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Dose-dependent satiating effect of whey relative to casein or soy

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ABSTRACT

Dietary protein plays a role in body weight regulation, partly because of its effects on appetite. The objective was to compare the effects of high or normal casein-, soy-, or whey-protein breakfasts on appetite, specific hormones, amino acid responses and subsequent energy intake. Twenty-five healthy subjects (mean±SEM BMI:23.9±0.3 kg/m²; age:22±1 years) received standardized breakfasts: custards with either casein-, soy, or whey-protein with either 10/55/35 (normal) or 25/55/20 (high)En% protein/carbohydrate/fat in a randomized, single-blind design. Appetite profile (Visual Analogue Scales) and amino acid concentrations were determined for 4 h whereas plasma glucose, insulin, active Glucagon-like Peptide 1 (GLP-1), and active ghrelin concentrations were determined for 3 h; the sensitive moment for lunch was determined. Subjects returned for a second set of experiments and received the same breakfasts, *ad lib* lunch was offered 180 min later; energy intake (EI) was assessed. At 10En%, whey decreased hunger more than casein or soy ($p<0.05$), coinciding with higher leucine, lysine, tryptophan, isoleucine, and threonine responses ($p<0.05$). At 25En% there were no differences in appetite ratings. Whey triggered the strongest responses in concentrations of active GLP-1 ($p<0.05$) and insulin ($p<0.05$) compared with casein and/or soy. There were no differences in EI. In conclusion, differences in appetite ratings between different proteins appeared at a normal concentration; at 10En% whey-protein decreased hunger more than casein- or soy-protein. At 25En% whey-protein triggered stronger responses in hormone concentrations than casein- or soy-protein. The results suggest that a difference in appetite ratings between types of protein appears when certain amino acids are above and below particular threshold values.

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1. Introduction

Obesity is the result of a positive energy balance, which arises when energy intake exceeds energy expenditure. Since body weight regulation involves several pathways, weight management requires a multi-factorial approach [1]. Recent findings suggest that a relatively high protein intake plays a role in weight loss as well as in weight maintenance thereafter, partly through increased postprandial and post-absorptive satiety [1–4]. Weigle et al. showed that satiety is of major importance, in an experiment in which a high protein diet reduced *ad lib* food intake while sustaining satiety at a comfortable level during a 12-week period [4]. In the present study we focused on short-term satiety effects, i.e. those induced by a single meal. It has been shown that protein is more satiating than carbohydrates or fat [5], and in previous experiments we found differences in appetite ratings between different concentrations of the same protein type [6–8]. It is,

however, less clear whether there are differences between different types of protein offered at fixed concentrations.

A limited number of human studies have compared different protein types in terms of their effects on satiety. Although Hall et al. found whey to be more satiating than casein [9], their results could not be replicated by others [10]. A study by Bowen et al. found no differences in postprandial responses after a whey, soy, or gluten protein preload [11]. Anderson et al. nevertheless showed that whey as well as soy protein, but not egg albumen, suppressed food intake at a meal 1 h later [12]. A comparison of the effects of beef, chicken, and fish protein revealed that fish protein increased satiety more than the other protein types [13]. Lang et al. did not observe significantly different effects of egg albumin, casein, gelatin, soy, pea, and wheat gluten on appetite scores or energy intake [14], and in another experiment, casein, soy, and gelatin protein did have weak but inconsistent effects on satiety and did not affect food intake at dinner [15]. Thus, results on the satiating properties of different types of protein have been inconclusive.

We investigated differences in appetite between three different protein types, namely casein, soy, and whey, all offered in two

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concentrations. The amounts of protein represented the highest recommended protein intake per day in energy balance, i.e. 25% of energy from protein, or the lowest, normal, protein intake per day, 10% of energy from protein [16]. Casein is considered to be a 'slow' protein, whereas whey protein is a relatively 'fast' protein [9,17–19]. Soy is a high quality vegetable protein that is often used in food products. Hence, the proteins offered differed in amino acid composition as well as in kinetics. Active Glucagon-like Peptide 1 (GLP-1) and active ghrelin were measured since previous research showed that GLP-1 may inhibit appetite and reduce food intake in humans [20,21], whereas ghrelin is an orexigenic hormone that has been suggested to be involved in meal initiation [22].

The aim of the present study was to compare the effects of casein, soy, or whey containing breakfasts on appetite ratings, plasma amino acid, glucose, insulin, active GLP-1, and active ghrelin concentrations and subsequent energy intake in two dosages. Since the timing of a test meal plays an important role [12], first the moment in time that may be sensitive to show a possible difference in food intake was determined by assessing appetite ratings and blood parameters for 4 h. Accordingly, in a subsequent experiment energy intake was measured at the pre-determined moment in time.

2. Subjects and methods

2.1. Subjects

Thirty healthy male and female volunteers (Body Mass Index 22–30 kg/m², age 18–40 years) were recruited by advertisements in local newspapers and on notice boards at the university. They underwent a screening procedure including medical history taking, measurement of body weight and height and cognitive restrained eating, using a Dutch translation of the Three Factor Eating Questionnaire (TFEQ) [23,24]. Twenty-five subjects (11 male, 14 female) were selected on the basis of being in good health, non-smokers, non-vegetarian, not cognitively dietary restraint (TFEQ Factor 1 score ≤9), not using medication apart from oral contraceptives and at most moderate alcohol users (≤10 alcoholic consumptions per week). Their mean age was 22 ± 1 years, and their body weight was 74.4 ± 1.8 kg (BMI: 23.9 ± 0.3 kg/m²). Written informed consent was obtained from these participants and the study protocol was approved by the Medical Ethics Committee of the University Hospital Maastricht.

2.2. Study design

A randomized, single-blind, within-subject experimental study was performed. All subjects came to the university on six occasions, separated by at least one week. On each test day, subjects received a subject-specific standardized breakfast. Appetite ratings and blood parameters were obtained for 4 h after breakfast.

The sensitive moment in time to offer lunch was determined by the latest time point after breakfast where there still were statistically significant differences in the changes of concentrations of the orexigenic hormone ghrelin between treatments. After two months, when the sensitive moment in time had been determined, subjects returned to the university on six occasions in a randomized, single-blind design, separated by at least one week. On each test day subjects received a subject-specific standardized breakfast, after which an *ad lib* lunch was offered at the pre-determined sensitive moment in time.

2.2.1. Breakfast

Breakfast was offered as a custard with either casein (Calcium Caseinate S, DMV International, Veghel, The Netherlands), soy (Supro® 590, The Solae Company, St. Louis, MO, United States of America), or whey (Ultra Whey 90, Volactive Functional Food Products, Orwell, United Kingdom) as a single protein source, with either protein/carbohydrate/fat: 10/55/35 En% (normal protein) or protein/carbohydrate/fat: 25/55/20 En% (high protein). Protein was exchanged with fat;

carbohydrate content was kept constant because of its effect on protein metabolism [25]. All custards had an energy density of 4 kJ/g. The breakfast contained 20% of daily energy requirements, calculated as basal metabolic rate (BMR), according to the equations of Harris–Benedict, multiplied by an activity index of 1.75 which is the average value reported for the general population in The Netherlands [26,27]. The mean energy content of the breakfast was 2.52 ± 0.07 MJ and the provided breakfasts were finished within 15 min.

The custards were produced by NIZO Food Research bv. (Ede, The Netherlands) and had tapioca starch (Farinex VA50T, AVEBE, Veendam, The Netherlands and Perfectamyl 3108 AVEBE, Veendam, The Netherlands) and sunflower oil (Reddy, NV Vandemoortele, Roosendaal, The Netherlands) respectively as the carbohydrate and fat sources and were citrus–vanilla (Citrus, J.B. de lange, Belfeld, The Netherlands; Vanilla, J.B. de lange, Belfeld, The Netherlands) flavored. Extensive product development and use of a taste panel lead to custards not different in color, taste, or viscosity. The amino acid composition of the custards is presented in Table 1.

2.2.2. Lunch

According to a normal Dutch lunch consisting of bread and a filling, lunch consisted of Turkish bread (400 g) with egg salad (400 g) with 13/41/46 En% protein/carbohydrate/fat with an energy density of 11.4 kJ/g. Beforehand it was tested whether all subjects liked the lunch sufficiently. Subjects were instructed to eat till they were comfortably full.

2.2.3. Study protocol

The protocol started at 08.00 h after an overnight fast from 22.00 h. A Venflon catheter was placed in a superficial dorsal vein of the hand for blood sampling. To obtain arterialized venous blood samples the hand was placed in a thermostatically controlled hot box at 60 °C for 20 min before the sampling time. A basal blood sample was taken and appetite ratings were scored. After 5 min a second basal blood sample was obtained and breakfast was offered (*t*=0 min). After the first and the last bite, taste perception was scored. Appetite ratings were completed just before breakfast and at 20, 40, 60, 80, 100, 120, 180, and 240 min after breakfast. Blood samples for urea and amino acid determination were obtained at –5 min and subsequently just after the appetite ratings; blood samples for determination of glucose, insulin, and active ghrelin concentrations were obtained before and 40, 60, 120, and 180 min after breakfast. In order to be able to observe

Table 1
Amino acid content of the breakfasts given as a custard with either 10% or 25% of energy from casein, soy, or whey protein (g amino acid/100 g custard)

	Casein 10%	Soy 10%	Whey 10%	Casein 25%	Soy 25%	Whey 25%
Glutamic acid ^a	0.477	0.328	0.381	1.127	0.816	0.957
Aspartic acid ^b	0.150	0.200	0.230	0.355	0.497	0.579
Cysteine	0.009	0.022	0.055	0.021	0.054	0.139
Serine	0.120	0.089	0.099	0.283	0.220	0.249
Histidine	0.064	0.048	0.039	0.152	0.119	0.097
Glycine	0.040	0.071	0.035	0.094	0.177	0.088
Threonine	0.090	0.066	0.150	0.214	0.164	0.378
Arginine	0.092	0.139	0.055	0.218	0.345	0.139
Alanine	0.064	0.073	0.106	0.150	0.182	0.266
Tyrosine	0.120	0.069	0.061	0.283	0.171	0.154
Valine	0.141	0.085	0.123	0.333	0.212	0.309
Methionine	0.064	0.022	0.048	0.152	0.056	0.121
Isoleucine	0.112	0.089	0.141	0.265	0.222	0.355
Phenylalanine	0.110	0.094	0.062	0.259	0.234	0.156
Tryptophan	0.027	0.023	0.039	0.064	0.057	0.099
Leucine	0.204	0.145	0.226	0.483	0.360	0.567
Lysine	0.172	0.110	0.201	0.405	0.274	0.504
Proline	0.230	0.087	0.128	0.544	0.216	0.321

^a Glutamic acid = glutamine + glutamate.

^b Aspartic acid = asparagine.

possible differences at 30 and 90 min between meals that were observed previously [28], venous blood samples for determination of active GLP-1 concentration were obtained separately before, and at 30, 60, 90, 120, and 180 min after breakfast by means of a Venflon catheter placed in an antecubital vein [28]. Subjects were allowed to drink maximally two glasses of water spread over the morning.

In the second set of experiments, the protocol started after an overnight fast from 22.00 h at 8.30 h with scoring appetite ratings. Breakfast was offered ($t=0$ min) and completed within 15 min. Subjects stayed in the laboratory till lunch was offered at the previously determined sensitive moment in time. The laboratory was a large room, and subjects were sitting in such a position that they were not able to see each other or each others meals. Maximally eight subjects were tested at the same time. They were sitting behind each other in a row at least 2 m apart, with room dividers in between subjects. Reminders of lunch were collected at the end, when all subjects had finished their lunch. They were not allowed to talk to each other, and background music prevented sound-signals that would indicate finishing meals. Subjects were allowed to drink three glasses of water spread over the entire test period.

2.3. Measurements

2.3.1. Appetite profile

To determine the appetite profile, hunger, fullness, satiety, and desire to eat were rated on 100 mm Visual Analogue Scales (VAS), anchored with 'not at all' and 'extremely' during the test day. VAS are often used to measure subjective appetite sensations and the validity and reproducibility has been shown in several studies [29,30]. Subjects were instructed to rate themselves by marking the scale at the point that was most appropriate to their feeling at that time. The distance from this point to the left end of the scale was measured in mm; changes from baseline (Δ) were calculated by subtracting the baseline score (-5 min) from the score at a certain time point.

2.3.2. Taste perception

Taste perception profiles of the custards were assessed after the first and the last bite of the breakfast using 100 mm Visual Analogue Scales (VAS), anchored with 'not at all' and 'extremely' on the aspects: pleasantness, sweetness, sourness, saltiness, bitterness, savouriness, crispiness, and creaminess.

2.3.3. Blood parameters

Blood was distributed into EDTA tubes for glucose, insulin, and active ghrelin measurement. For active GLP-1 measurement blood was collected in EDTA tubes with added dipeptidyl peptidase IV inhibitor. For amino acid and urea determination, blood was collected in lithium heparin tubes. Blood samples were centrifuged at 4 °C for 10 min at 3000 rpm. Hydrochloric acid and phenylmethylsulfonyl fluoride were added to plasma for active ghrelin determination. For amino acid analysis, 250 μ l plasma was deproteinized by mixing it with 20 mg dry sulfosalicylic acid. For analysis of urea, 200 μ l plasma was deproteinized by mixing it with 20 μ l of a 500 g/l trichloroacetic acid solution. All samples were stored at

-80 °C until further analysis. Plasma glucose concentrations were determined using the hexokinase method (Glucose HK 125 kit, ABX diagnostics, Montpellier, France). Insulin concentrations were measured by RIA (Linco Research Inc., St. Charles, Missouri, USA). Plasma active ghrelin concentrations were measured by ELISA (Linco Research Inc., St. Charles, Missouri, USA). Plasma active GLP-1 samples were analyzed using ELISA (EGLP-35K; Linco Research Inc., St. Charles, Missouri, USA). Plasma concentrations of amino acids were determined with the use of a fully automated HPLC (Pharmacia, Woerden, The Netherlands), after precolumn derivatization with *o*-phthalaldehyde [31]. Plasma urea was analyzed spectrophotometrically on a COBAS Mira S (Roche Diagnostica, Hoffman-La Roche, Basel, Switzerland).

2.3.4. Energy intake

The food provided for lunch was weighed before and after eating and energy intake was calculated by multiplying the amount of food consumed with the energy value of the food as indicated by the product labels (11.4 kJ/g).

2.4. Statistical analysis

Data are presented as mean changes from baseline \pm standard error to the mean (SEM), unless otherwise indicated [32]. The area under the curve (AUC) or area above the curve (AAC) of changes from baseline over time for appetite ratings and glucose, insulin, active GLP-1, active ghrelin, amino acid and urea concentrations was calculated using the trapezoidal method. To determine possible differences between the different types of protein at a concentration of 10% and 25% of energy from protein, a repeated measures ANOVA between factors with protein level as factor was carried out. When there was no effect of protein level a repeated measures ANOVA with Fisher's PLSD correction for multiple comparisons within one protein type was carried out. After the second set of experiments, a repeated measures ANOVA between factors with protein level as factor and a repeated measures ANOVA with Fisher's PLSD correction for multiple comparisons was carried out to determine possible differences in energy intake. A p -value <0.05 was regarded as statistically significant. Statistical procedures were performed using StatView 5.0 (SAS Institute Inc., USA, 1998).

3. Results

3.1. Appetite profile

Baseline appetite ratings were not different between treatments. The changes in appetite ratings per type of protein did not differ depending on protein level. Within one protein level, namely at 10% of energy from protein, the AAC of hunger ratings was increased more after a breakfast with whey than after a breakfast with casein (8643 ± 814 mmVAS. h vs. 6099 ± 1066 mmVAS. h, $p < 0.05$, Table 2, Fig. 1). Hunger suppression was increased more after a breakfast with whey than after a breakfast with casein at 20, 40, 60, 80, 120, and 240 min after breakfast ($p < 0.05$ for each time point, Fig. 1) and after a breakfast with whey than after a breakfast with soy at 20 min after breakfast ($p < 0.05$, Fig. 1). At the level of 25% of

Table 2

Hunger, glucose, insulin, GLP-1, and ghrelin responses expressed as area above the curve (hunger, ghrelin) or area under the curve (glucose, insulin, GLP-1) after a breakfast given as a custard with either 10% or 25% of energy from casein, soy, or whey protein in 25 subjects (men and women)

	Casein	Soy	Whey	Casein	Soy	Whey
	10%	10%	10%	25%	25%	25%
Hunger (mmVAS. h)	6099 \pm 1066 w	7348 \pm 1199	8643 \pm 814 c	8217 \pm 1082	9210 \pm 1011	7613 \pm 1101
Glucose (mmol/l. h)	124 \pm 14	120 \pm 21	99 \pm 17	68 \pm 18 s	122 \pm 13 c	95 \pm 11
Insulin (mU/l. h)	6530 \pm 621 s	4936 \pm 469 c	5820 \pm 386	4792 \pm 980 sw	7520 \pm 929 c	9159 \pm 692 c
GLP-1 (pmol/l. h)	218 \pm 78	216 \pm 94	266 \pm 71	161 \pm 90 w	195 \pm 72	425 \pm 135 c
Ghrelin (pmol/l. h)	708 \pm 140 s	399 \pm 108 c	439 \pm 106	546 \pm 184	430 \pm 128	722 \pm 145

ANOVA repeated measures with Fisher's PLSD correction.

Within one protein level, c indicates a significant difference with casein, s indicates a significant difference with soy, w indicates a significant difference with whey.

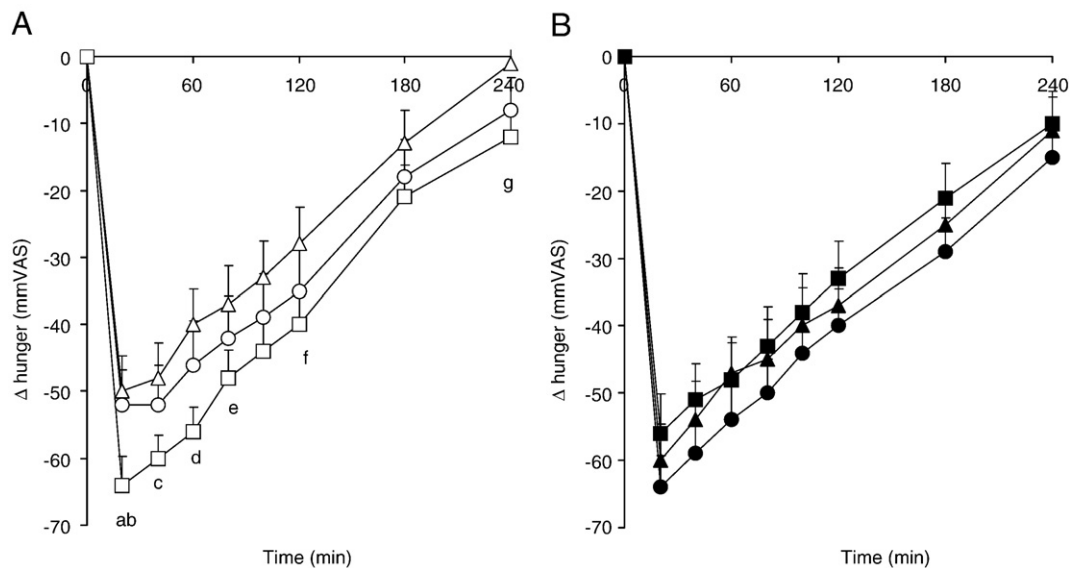


Fig. 1. Changes in hunger ratings (mmVAS) after a breakfast offered as a custard with either 10% (A) or 25% (B) of energy from casein, soy, or whey protein expressed as delta compared to baseline in 25 subjects (men and women). Values are means+SEM. Δ 10% of energy from casein, \circ 10% of energy from soy, \square 10% of energy from whey, \blacktriangle 25% of energy from casein, \bullet 25% of energy from soy, \blacksquare 25% of energy from whey. ANOVA repeated measures with Fisher's PLSD correction * $p < 0.05$, ** $p < 0.01$; a whey < casein **, b whey < soy *, c whey < casein *, d whey < casein **, e whey < casein *, f whey < casein *, g whey < casein *; area above the curve hunger 10% casein < whey *.

energy from protein there were no differences in hunger ratings between casein, soy, or whey (Fig. 1). The other appetite ratings were similar with respect to AUC or AAC (fullness, satiety, desire to eat) (data not shown).

3.2. Taste perception

Pleasantness of taste of the custards with the first bite was sufficient with a mean score of 56 ± 4 mmVAS without differences between custards.

3.3. Glucose

Baseline plasma glucose concentrations were not different between treatments. The changes in glucose concentration per type of protein did not differ depending on protein level. Within one protein level there were no differences in changes in glucose concentration between casein, soy, or whey after a breakfast with 10% of energy from protein, however, after a breakfast with 25% of energy from protein, glucose concentrations were increased more after a breakfast with soy than after a breakfast with casein (122 ± 13 mmol/l. h vs. 68 ± 18 mmol/l. h, $p < 0.05$, Table 2).

3.4. Insulin

Baseline plasma insulin concentrations were not different between treatments. The changes in insulin concentration per protein type differed depending on the level of protein. At the level of 10% of energy from protein, insulin concentrations were increased more after a breakfast with casein than after a breakfast with soy (6530 ± 621 mU/l. h vs. 4936 ± 468 mU/l. h, $p < 0.05$, Table 2, Fig. 2). At the level of 25% of energy from protein, insulin concentrations were increased more after a breakfast with soy or whey than after a breakfast with casein (7520 ± 929 mU/l. h or 9159 ± 692 mU/l. h, vs. 4792 ± 980 mU/l. h, $p < 0.05$ and $p < 0.01$ respectively, Table 2, Fig. 2).

3.5. Active GLP-1

Baseline plasma active GLP-1 concentrations were not different between treatments. The changes in active GLP-1 concentration per

type of protein did not differ depending on protein level. Within one protein level there were no differences in changes in active GLP-1 concentration between casein, soy or whey after a breakfast with 10% of energy from protein, however, after a breakfast with 25% of energy from protein, active GLP-1 concentrations were increased more after a breakfast with whey than after a breakfast with casein (425 ± 135 pmol/l. h vs. 161 ± 90 pmol/l. h, $p < 0.05$, Table 2).

3.6. Active ghrelin

Baseline plasma active ghrelin concentrations were not different between treatments. The changes in active ghrelin concentration per type of protein did not differ depending on protein level. Within one protein level, namely 10% of energy from protein, active ghrelin concentrations were decreased more after a breakfast with casein than after a breakfast with soy (AAC 708 ± 140 pmol/l. h vs. 399 ± 108 pmol/l. h, $p < 0.05$, Table 2, Fig. 3).

3.7. Amino acids

Baseline plasma amino acid concentrations were not different between treatments. The changes in glutamate, asparagine, glycine, threonine, citrulline, arginine, valine, methionine, isoleucine, phenylalanine, tryptophan, leucine, and lysine concentration per type of protein differed depending on protein level. The AUC of the response of the different amino acids after the six different breakfasts is presented in Fig. 4; differences ($p < 0.05$) between treatments are indicated with C (different from the casein breakfast), S (different from the soy breakfast), or W (different from the whey breakfast).

At the level of 10% of energy from protein the amino acids threonine, alanine, alpha-aminobutyric acid, isoleucine, tryptophan, leucine, and lysine were increased more after a breakfast with whey than after a breakfast with casein ($p < 0.05$, Fig. 4). The amino acids threonine, alpha-aminobutyric acid, methionine, isoleucine, tryptophan, leucine, and lysine were increased more after a breakfast with 10% of energy from whey than after a breakfast with 10% of energy from soy ($p < 0.05$, Fig. 4).

At the level of 25% of energy from protein, the amino acids asparagine, threonine, alpha-aminobutyric acid, valine, isoleucine, tryptophan, leucine and lysine were increased more after a breakfast with whey than after a breakfast with casein ($p < 0.05$, Fig. 4). The

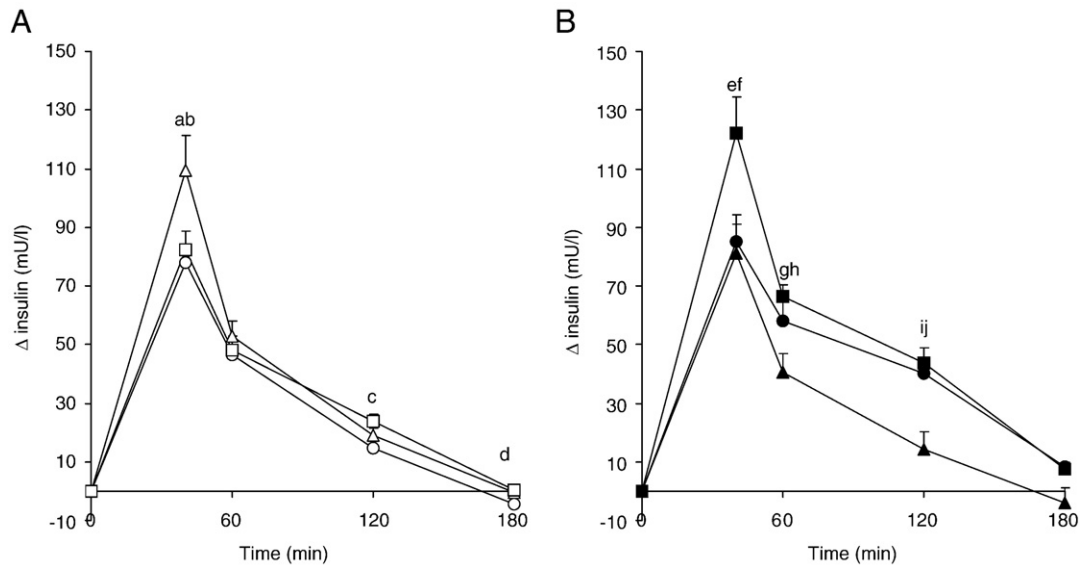


Fig. 2. Changes in insulin concentrations (mU/l) after a breakfast offered as a custard with either 10% (A) or 25% (B) of energy from casein, soy, or whey protein in 25 subjects (men and women). Values are means+SEM. Δ 10% of energy from casein, \circ 10% of energy from soy, \square 10% of energy from whey, \blacktriangle 25% of energy from casein, \bullet 25% of energy from soy, \blacksquare 25% of energy from whey. ANOVA repeated measures with Fisher's PLSD correction * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$; a soy < casein **, b whey < casein **, c soy < whey **, d soy < whey **, e casein < whey **, f soy < whey **, g casein < soy *, h casein < whey **, i casein < soy ***, j casein < whey ***, Area Under the Curve insulin soy 10% < casein 10% **, area under the curve insulin casein 25% < soy 25% *, casein 25% < whey 25% **.

amino acids threonine, alpha-aminobutyric acid, valine, isoleucine, tryptophan, leucine and lysine were increased more after a breakfast with whey than after a breakfast with soy ($p < 0.05$, Fig. 4).

3.8. Energy intake

Based on the significant differences in concentrations of the orexigenic hormone active ghrelin at 180 min, the *ad lib* lunch was offered at 180 min after breakfast.

At the level of 10% of energy from protein, energy intake at lunch was 3133 ± 226 kJ, 3098 ± 286 kJ and 2879 ± 239 kJ after the breakfast with casein, soy, or whey respectively (ns). At the level of 25% of energy from protein, energy intake at lunch was 3080 ± 229 kJ, $3212 \pm$

280 kJ and 2876 ± 243 kJ after the breakfast with casein, soy, or whey, respectively (ns).

4. Discussion

Based upon the appetite ratings, a breakfast with whey reduced hunger more than a breakfast with casein, and at short term also than soy, at the level of 10% of energy from protein, however, this did not affect subsequent energy intake at lunch. At the level of 25% of energy from protein, the breakfast with whey triggered the strongest response in insulin and active GLP-1, however, there were no differences in appetite ratings or energy intake at lunch.

The citrus-vanilla flavored custards were similar to custards widely available and often consumed in The Netherlands. It is therefore unlikely

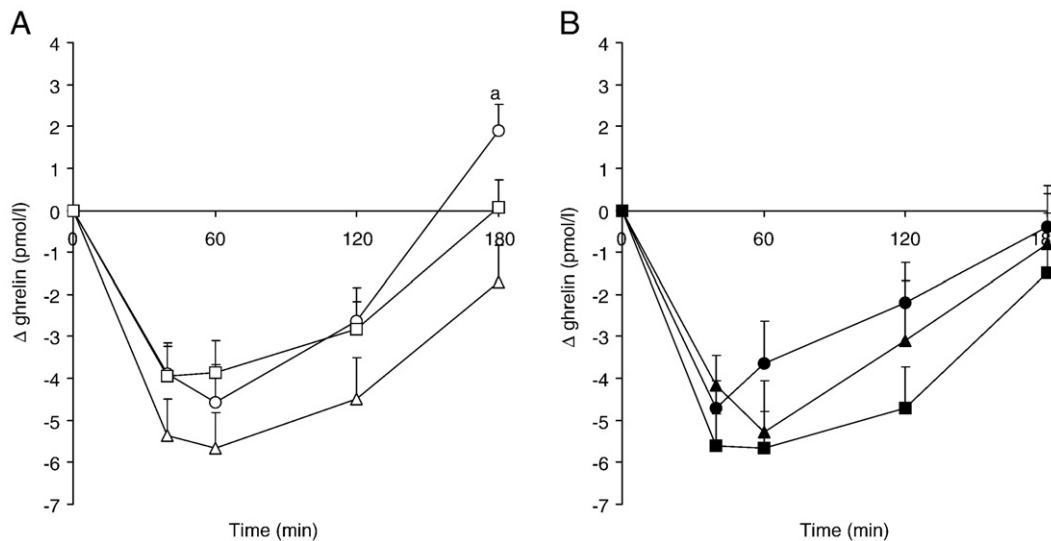


Fig. 3. Changes in active ghrelin concentrations (pmol/l) after a breakfast offered as a custard with either 10% (A) or 25% (B) of energy from casein, soy, or whey protein expressed as delta compared to baseline in 25 subjects (men and women). Values are means+SEM. Δ 10% of energy from casein, \circ 10% of energy from soy, \square 10% of energy from whey, \blacktriangle 25% of energy from casein, \bullet 25% of energy from soy, \blacksquare 25% of energy from whey. ANOVA repeated measures with Fisher's PLSD correction * $p < 0.05$, ** $p < 0.01$; a soy < casein **, area above the curve active ghrelin casein 10% < soy 10% *.

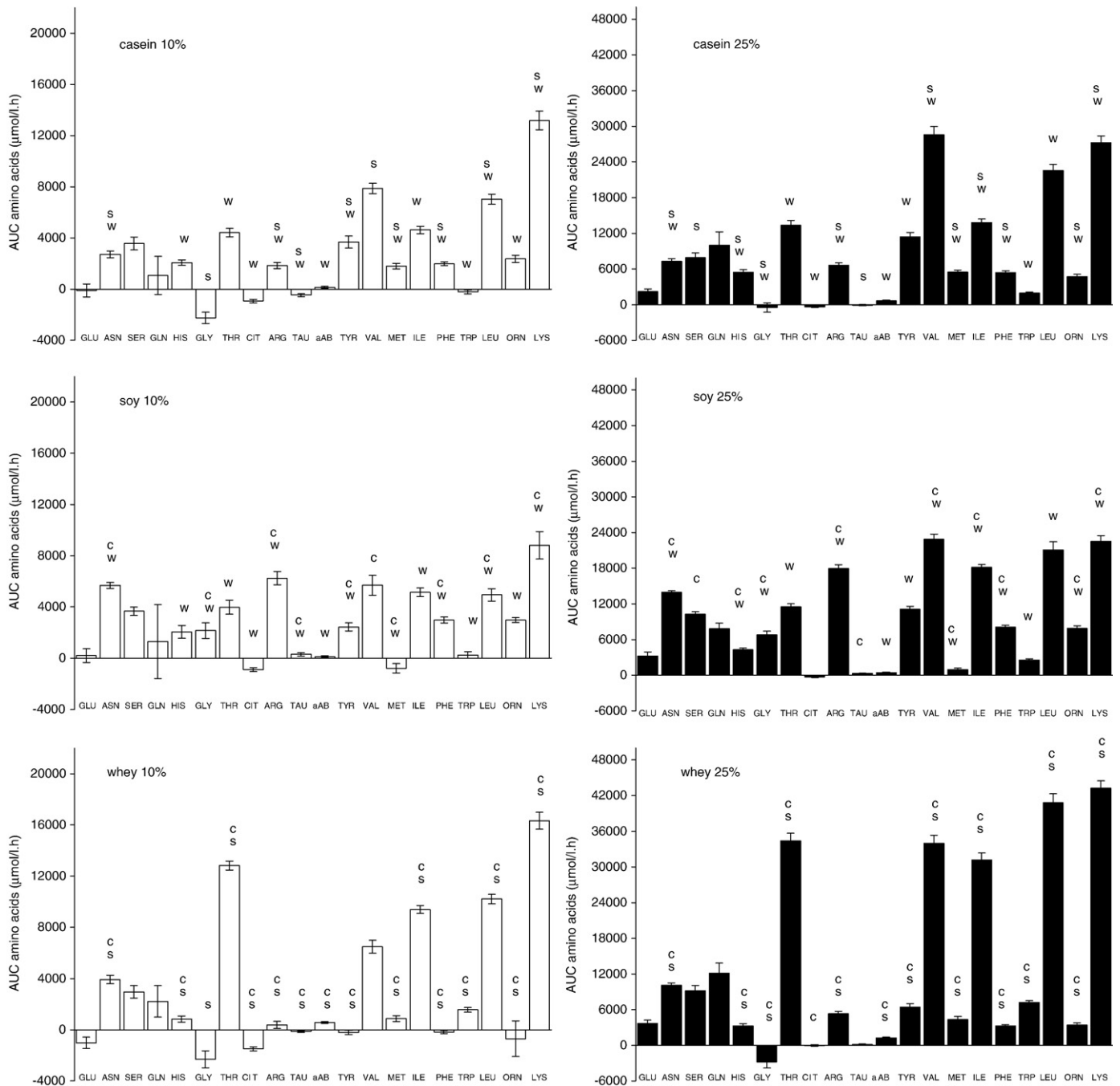


Fig. 4. Amino acid responses expressed as AUC from baseline ($\mu\text{mol/l h}$) after a breakfast offered as a custard with either 10% or 25% of energy from casein, soy, or whey protein in 25 subjects (men and women). Values are means+SEM. GLU: glutamate, ASN: asparagine, SER: serine, GLN: glutamine, HIS: histidine, GLY: glycine, THR: threonine, CIT: citrulline, ARG: arginine, TAU: taurine, aAB: alpha-aminobutyric acid, TYR: tyrosine, VAL: valine, MET: methionine, PHE: phenylalanine, TRP: tryptophan, LEU: leucine, ORN: ornithine, LYS: lysine. ANOVA repeated measures with Fisher's PLSD correction; C: different from the casein breakfast ($p < 0.05$), S: different from the soy breakfast ($p < 0.05$), W: different from the whey breakfast ($p < 0.05$).

that unfamiliarity with the breakfasts influenced satiety responses. To avoid any specific sensory effect of the iso-energetic custards, food technology was involved to optimize taste and hedonic value of the breakfasts. The custards were citrus-vanilla flavored and after being tested by a professional taste panel of NIZO Food Research, taste perception and hedonic values again were evaluated by the subjects and were excluded to affect appetite profile ratings differently.

The relatively stronger hunger suppression after a breakfast with 10% of energy from whey, compared with a breakfast with casein or soy, coincided with a greater increase in responses of leucine, lysine, tryptophan, isoleucine, and threonine; amino acids which may be

involved in the satiety response. Leucine and isoleucine are two of the three branched-chain amino acids that regulate protein synthesis and degradation, as well as insulin secretion and synthesis [33]. The concomitant high energy costs of these processes may be related to satiety [5,34]. Tryptophan has been suggested to be involved in satiety via brain serotonin; serotonin is synthesized from tryptophan and is an important regulator of appetite, macronutrient preference, and mood [35]. The results of the present study suggest that tryptophan may indeed be involved in the satiety process. Lysine has previously been shown to produce a moderate decrease in food intake in sheep [36]; excess levels of threonine added to a low protein diet resulted in

a reduced weight gain in rats [37]. The mechanisms via which these amino acids may influence satiety are not clear however and need to be further established.

The moment at which lunch was offered was based upon the latest moment in time when there were significant differences in ghrelin concentrations between treatments. Ghrelin has been suggested to play a physiological role in meal initiation in humans [22]. Differences in ghrelin concentrations may therefore result in differences in energy intake. Although there were differences in appetite ratings between the different types of protein at the level of 10% of energy from protein, there were no significant differences in energy intake. Apparently in this experiment the differences in appetite ratings were not large enough to induce effects on energy intake.

With respect to the proteins offered in a concentration with 25% of energy there were no differences in appetite ratings. Nevertheless there were significant differences in hormone responses between the breakfasts with 25% of energy from protein; after a breakfast with whey, increases in insulin and active GLP-1 were larger than after a breakfast with casein and/or soy. Previously it has been shown that casein coagulates in the stomach which delays gastric emptying [17,18,38], this resulted in slower and less pronounced physiological responses compared with soy and whey. The relatively larger insulin responses after the high whey breakfast is in accordance with the findings of Frid and others, reporting an insulinotropic effect of whey which partly may be explained by the involvement of certain amino acids that have insulinogenic properties [39,40]. The larger increase in active GLP-1 concentrations after a breakfast with whey can be explained by the finding that whey inhibits dipeptidyl peptidase IV activity, the enzyme involved in the breakdown of active GLP-1, thus prolonging the action of active GLP-1 [41]. Active GLP-1 enhances satiety and is an incretin hormone whereas insulin has been reported to inhibit active GLP-1 secretion, probably as a negative feedback loop [20,21,42]. Although insulin and active GLP-1 are considered as 'satiety' hormones, there was no larger increase in hunger ratings after a breakfast with 25% of energy from whey than after the other breakfasts. Here, a mathematical uncoupling of a satiating effect and increases in 'satiety' hormone concentrations takes place.

Since there were differences in appetite ratings between types of protein at the level of 10% but not at the level of 25% of energy, it seems that the concentrations of certain amino acids need to be above a particular threshold to promote a relatively stronger hunger suppression or greater satiety. The results suggest that certain proteins will reach these threshold concentrations earlier than other types of protein. After a breakfast with whey, sufficiently increased amino acid concentrations were reached at the level of 10% of energy, whereas concentrations were lower after a breakfast with casein or soy. Hence, discriminating between types of protein is probably not sensitive anymore at a higher level of protein, since the amino acid responses of all breakfasts were above the threshold.

This is the first study that investigated acute differences in appetite ratings between types of protein in concentrations within the normal range in realistic mixed meals. The relatively high amount of protein (≥ 50 En%) may have caused the lack of differences in satiety between different types of protein when comparing appetite after either a casein, whey, or carbohydrate preload or when comparing *ad lib* food intake after whey, soy, or gluten protein [10,11]. It may not be possible to distinguish satiating properties of different types of protein anymore when the concentration of amino acids is above a threshold level. In the present study the protein part of the breakfast consisted exclusively of the protein type to be investigated whereas in previous comparisons of the satiating capacities of egg albumin, casein, gelatin, soy, pea, and wheat gluten protein only 60 to 70% of the protein part was manipulated. This may have led to diminished results and consequently the absence of significant differences in appetite ratings between the different protein types [14,15]. Timing plays an important role in studying the effect of protein on food intake. An amount of

0.65 g/kg body weight of whey, soy, or egg albumen protein did induce significant differences in food intake 1 h after the preload compared with water as control, however, this is a rather irrelevant moment in time for a next meal in a normal, free-living situation [12]. Hall et al. observed a reduced desire to eat after a whey preload of 1.7 MJ with 48 g protein compared with a similar casein preload [9]. However, 90 min after the preload subjects already got a standard lunch with fixed energy intake. The reduced desire to eat was observed between 90 and 180 min; conclusions about the solely effect of the two preloads can hardly be drawn. Moreover, the conclusions by Hall et al. could not be confirmed in a similar study from Bowen and colleagues [10,11].

This study provides new information for the development of weight-loss diets. Whey-protein can be used, already with an amount of 10 En%, in a diet to help people comply to a diet. When people feel less hungry and desire to eat is suppressed, it is easier for them to comply to a diet because they really feel an effect of the diet. Although there were no short term differences in energy intake between casein, soy and whey in the present study, people may comply better to a high protein diet with whey and eventually eat less and lose weight.

In conclusion, hunger was decreased more after a breakfast with whey than after a breakfast with casein or soy in a concentration of 10% of energy from protein, which coincided with increased concentrations of the amino acids leucine, lysine, tryptophan, isoleucine, and threonine. Although there were no differences in appetite ratings between casein, soy, or whey at a level of 25% of energy from protein, the breakfast with whey triggered stronger responses in hormone concentrations than the breakfasts with casein or soy. The results suggest that a difference in appetite ratings between different types of protein may only appear when certain amino acids are above and below particular thresholds.

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