

Effects of isoenergetic overfeeding of either carbohydrate or fat in young men

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Ten pairs of normal men were overfed by 5 MJ/d for 21 d with either a carbohydrate-rich or a fat-rich diet (C- and F-group). The two subjects in each pair were requested to follow each other throughout the day to ensure similar physical activity and were otherwise allowed to maintain normal daily life. The increase in body weight, fat free mass and fat mass showed great variation, the mean increases being 1.5 kg, 0.6 kg and 0.9 kg respectively. No significant differences between the C- and F-group were observed. Heat production during sleep did not change during overfeeding. The RQ during sleep was 0.86 and 0.78 in the C- and F-group respectively. The accumulated faecal loss of energy, DM, carbohydrate and protein was significantly higher in the C- compared with the F-group (30, 44, 69 and 51 % higher respectively), whereas the fat loss was the same in the two groups. N balance was not different between the C- and F-group and was positive. Fractional contribution from hepatic *de novo* lipogenesis, as measured by mass isotopomer distribution analysis after administration of [1-¹³C]acetate, was 0.20 and 0.03 in the C-group and the F-group respectively. Absolute hepatic *de novo* lipogenesis in the C-group was on average 211 g per 21 d. Whole-body *de novo* lipogenesis, as obtained by the difference between fat mass increase and dietary fat available for storage, was positive in six of the ten subjects in the C-group (mean 332 (SEM 191) g per 21 d). The change in plasma leptin concentration was positively correlated with the change in fat mass. Thus, fat storage during overfeeding of isoenergetic amounts of diets rich in carbohydrate or in fat was not significantly different, and carbohydrates seemed to be converted to fat by both hepatic and extrahepatic lipogenesis.

Overfeeding: *De novo* lipogenesis: Energy expenditure: Leptin: Faecal loss

Energy consumed as carbohydrate, fat or protein, is either lost, used for internal or external work or stored if in excess. Several studies indicate that human subjects have a preference for oxidation of protein and carbohydrate, and that fat ingested in surplus of energy demand is deposited (Jebb *et al.* 1996; Jéquier, 1998; Stubbs, 1998; Wolfe, 1998). In several recent reviews on weight gain and energy intake in man the importance of the fat : carbohydrate ratio in the food has been stressed (for example see Flatt, 1987, 1995; Swinburn & Ravussin, 1993; Astrup & Raben, 1995; Lissner & Heitmann, 1995; Stubbs, 1998). Such a correlation between weight gain and the fat : carbohydrate ratio has been tentatively explained by a higher palatability of fat, a higher energy expenditure with carbohydrate (originating from a larger thermogenic effect of carbohydrates including

a larger storage cost of deposition in comparison with fat), an increased activity of the sympathetic nervous system, and an induced higher physical activity level (for example see Astrup & Raben, 1995; Lissner & Heitmann, 1995). However, it is not clear whether overconsumption of isoenergetic amounts of fat or carbohydrate will result in different fat deposition. Fat deposition induced by overfeeding with a high-carbohydrate diet can occur either as a result of an increased direct storage of ingested triacylglycerol in adipose tissue and/or as a result of *de novo* lipogenesis (DNL) from ingested carbohydrate. The control of the relative contribution of these two processes in man is not well understood. DNL takes place in the liver and in adipose tissue, but is generally considered to be of too low a capacity to explain observed weight gain (Hellerstein *et al.* 1996).

Abbreviations: DNL, *de novo* lipogenesis; SEE, sleeping energy expenditure.

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Short-term ingestion of a large amount of carbohydrate resulted in increased storage of glycogen with no or only very limited DNL (Acheson *et al.* 1982, 1984). In longer experiments with extreme carbohydrate overfeeding, however, DNL was not negligible (Schutz *et al.* 1985; Acheson *et al.* 1988). In addition Pasquet *et al.* (1992) found an increase in body fat mass after massive overfeeding of a diet rich in carbohydrate, likely to be caused mainly by DNL (see calculations in Hellerstein *et al.* 1996). In more recent experiments Horton *et al.* (1995) studied effects of a 50% increase in energy intake, provided either as carbohydrate or as fat. Although the subjects given a surplus of fat showed an initially higher increase in body weight, the body weight after 14 d intervention was the same in the two groups. DNL was not measured. Aarsland *et al.* (1997) found a DNL of about 170 g/d in human subjects massively overfed with carbohydrate, predominantly in the form of intravenous glucose, where hepatic contribution accounted for only 2% of the total. Schwarz *et al.* (1995) fed 50% excess energy as carbohydrate and observed only modest increases in the absolute rate of hepatic DNL.

The aim of the present study was to investigate, in a period of 3 weeks, the effects of overfeeding of isoenergetic amounts of normal foods either high in carbohydrate or high in fat on body composition, heat production during sleep and hepatic and total DNL. The subjects of the study were non-obese young men and, in contrast to most previous studies, they were allowed to maintain a normal daily life. Similar physical activity of the carbohydrate- and the fat-group was ensured by letting the subjects follow each other throughout the experimental period, i.e. in the daytime they were free to engage in whatever activities they could agree upon ('Siamese' activity twins). In addition to changes in body composition, energy expenditure (direct and indirect) at sleep and DNL, energy loss in faeces, N balance, and the concentration of glucose, insulin and leptin in plasma were measured.

Design and methods

Study design

The study was designed to evaluate the effects of isoenergetic overfeeding of either a high-fat or a high-carbohydrate diet with an intended daily food excess of 5 MJ. Twenty normal young men were studied. They were divided into pairs, and the experiment was carried out with one pair at a time. The intervention period was planned for 21 d preceded by a 2-week period of recording habitual food intake. The study involved 3 phases. Phase 1, the prephase, lasted 10–14 d. The subjects lived at home and registered their habitual food intake by weighing all food items eaten, as

well as recording all drinks by content and volume. Phase 2, baseline measurements, lasted 2–3 d. The subjects moved into the department, living and acting as 'Siamese' activity twins. They ate their preferred meals while baseline data for the experiment were obtained. Detailed instructions for the daily routines regarding body composition measurement, food registration and consumption, and faeces and urine collections were given. During this period the baseline measurements were obtained. Phase 3, the intervention period, was scheduled for 21 d. The energy intake was increased by 5 MJ above the individual habitual energy intake as determined during phase 1.

Each person in a pair was randomly assigned to the high-carbohydrate (C-group) or the high-fat diet (F-group), and denoted C1–C10 and F1–F10 respectively. All faeces and urine were collected during phase 3. Daily measurements of body composition were obtained in the morning after voiding, and before any meals were taken. Once a week a full day was used for measurements of body composition by underwater weighing, hepatic DNL and blood variables. For logistic reasons phase 3 varied from 17 to 22 d for the ten pairs. All results presented are normalised to 21 d.

Subjects

Twenty apparently healthy subjects (male university students) with no previous record of obesity, weight loss or other metabolic illnesses participated in the study. They were between 20 and 25 years of age with BMI between 19.7 and 25.4 kg/m². Subjects were paired according to habitual activity and body size. There were no statistically significant differences between the two groups with regard to age, height, body weight or fat mass (Table 1).

While living in the quarters of the department, the subjects conformed with the following restrictions. The subjects acted as 'Siamese' activity twins in order to have similar physical activity within the pair. Thus the pair had to spend the entire 3 weeks together. All meals were prepared and consumed at the department. All nights were spent in the department's quarters. Every second night each subject slept in a whole-body calorimeter where both direct and indirect energy expenditure were measured. The subjects went to bed and woke up according to their own schedules. The subject who woke up first called the other subject. All twenty subjects completed the experiment.

Food intake

Food intake was calculated as energy and nutrients by means of the food composition database Dankost 2000 (Dansk Catering Center A/S, Copenhagen, Denmark),

Table 1. Subjects characteristics at the start of the experimental period (phase 2)
(Mean values with their standard errors)

Subjects	<i>n</i>	Age (years)		Body weight (kg)		Height (cm)		Fat mass (kg)	
		Mean	SEM	Mean	SEM	Mean	SEM	Mean	SEM
C1–C10	10	22.3	1.7	76.4	8.8	183	5	12.0	3.4
F1–F10	10	22.4	1.9	73.4	6.7	181	7	10.6	2.7

based on the Danish Food Composition Tables (Møller & Saxholt, 1996). These food tables present data on metabolizable energy. We consider them equal to total energy intake because of evidence that the values reported in the tables are too high (Nielsen *et al.* 1995). In phase 3, the C-group received 78 % of energy as carbohydrates, 11 % of energy as protein and 11 % of energy as fat, while the F-group received 58 % of energy as fat, 11 % of energy as protein and 31 % of energy as carbohydrate. A dietitian managed the diet planning and provided recipes with measures and explanations for each of the meals. The recipes were in general similar for all the subjects in the C-group and in the F-group, but were individually adjusted to provide approximately 5 MJ above the habitual intake, as well as to account for individual tastes and preferences. The subjects did the shopping themselves, and during the meal preparations they weighed and recorded all food items in accordance with the recipes and instructions given. The research staff regularly, but unannounced, confirmed that the meal preparations and the recording of intake of food, drinks, snacks and sweets took place in accordance with the instructions.

Analytical procedures

Anthropometry. Subjects were weighed every morning on a Tanita body fat analyser TBF-305 (Tanita Corp., Tokyo, Japan) (impedance method). Data for body weight, fat free mass, fat mass, and total body water were obtained. The subjects' standing height was measured on the first day in the department, and this measurement was used in all the body composition calculations.

As a validation of the impedance method fat mass and fat free mass were also measured once weekly in the morning by underwater weighing (densitometry). The subject was placed in a sitting position, and the residual volume of the lungs during the submergence was determined with the N₂-dilution method. The diluted air mixture was measured for its concentration of O₂ and CO₂ with a paramagnetic O₂ analyser (Servomex OOA 184; Servomex Controls Ltd, Crowborough, Sussex, UK) and an infrared CO₂ analyser (Beckman LB2; Sormedics Corp., Anaheim, CA, USA). Both analysers were calibrated with air which had its content determined using micro-Scholander technique (Scholander, 1947). Density measurements were repeated at approximately 12 min intervals until at least three of them agreed within 0.0015 g/ml.

Energy expenditure. Sleeping energy expenditure (SEE) was measured directly (SEE_{dir}) and indirectly (SEE_{indir}). The latter was calculated from measurements of the CO₂ production and O₂ uptake and converted to energy expenditure using equation 19 from Elia & Livesey (1992). Measurements are given as the mean value of all the measured SEE during the last 3 h of sleep. SEE_{dir} was corrected for the nightly drop in body temperature by measurements of rectal temperature before entering the chamber and on the following morning. The drop in core temperature was assumed to be linear during the sleeping period, and the distribution between core and shell was considered constant overnight. The calorimeter chamber was kept at 28°C. Details of the chamber are given elsewhere (Faber *et al.* 1998). Daily calibrations of the direct measurements and the flow through the chamber were

performed with an electrical dummy and evaporation from a water tank placed on a scale in the chamber. Calibrations of the indirect measurements were performed daily with calibrations of the O₂ and CO₂ analysers using air with a known content as measured by the micro-Scholander technique (Scholander, 1947).

Faeces and urine. All faeces and urine were collected during phase 3. The urine was kept cold (0°C), and once a day urine from the last 24 h was weighed and a representative sample frozen for later analysis. Faeces were stored in plastic bags at -18°C after being weighed. Before analysis, faeces were thawed and homogenised by a Stomacher Lab Blender® (Seward Medical Ltd, London, UK), and representative samples from each 24 h period were collected. The samples were freeze-dried and analysed for energy content by bomb calorimetry and for N content by the Kjeldahl method respectively. Analysis of the energy content of faeces for each day was determined for one subject, and showed an approximately constant energy content per g DM after day 3. Therefore the analysis of faeces energy content on all other subjects was performed on pooled faeces samples from day 1-2, day 3-7, day 8-14, and day 15 to the end of phase 3. The energy content of protein in faeces was calculated from the N content (g) multiplied by 6.25 and by 23.6 kJ/g. Carbohydrate and fat content were calculated from residual energy assuming an energy value of 17.4 kJ/g for carbohydrate and 39.3 kJ/g for fat. The values reported are the mean of duplicate determinations.

Urine samples were analysed for N by the Kjeldahl method. Protein oxidation (g) was calculated as urine N excretion (g) multiplied by 6.25.

Hepatic de novo lipogenesis. Fractional hepatic DNL, i.e. the percentage of total plasma VLDL-fatty acids synthesised *de novo*, was determined once in phase 2 and three times (i.e. once weekly) in phase 3 by mass isotopomer distribution analysis for pairs no. 2-10 as described by Hellerstein *et al.* (1991) and Faix *et al.* (1993). In order to reach a near-steady state of [¹³C]acetyl-CoA labelling, 400 mg [1-¹³C]acetate, 99 % enrichment, in approximately 50 ml water was given orally every 60 min from 10.00-19.00 hours. Blood samples were obtained at 10.00 hours while the subject was fasting and at 18.00, 19.00, 20.00 and 21.00 hours (non-fasting values) and were immediately centrifuged at -5°C in order to separate plasma and erythrocytes. Plasma was stored at -21°C until analysis. Before VLDL isolation from the plasma, chylomicrons were removed by centrifugation at 20°C for 20 min at 112 500 g of 2 ml plasma overlaid by 1.5 ml salt solution (Mills *et al.* 1984). The lower 2.2 ml was removed and layered under 2 ml salt solution and centrifuged again at 20°C for 22 h at 300 000 g. The VLDL fraction (the top layer) was aspirated and used for determination of the concentration of VLDL and for mass isotopomer distribution analysis. The concentration of VLDL was measured by enzymatic glycerol determination (Wieland, 1984) after hydrolysis in ethanolic KOH (Kates, 1986). Mass isotopomer distribution analysis was carried out on methyl esters of the VLDL fatty acids, prepared by transesterification of a lipid extract as previously described (Hellerstein *et al.* 1991).

The rate constant of hepatic DNL was determined for pairs no. 7-10 in phase 2 and at the end of phase 3 by

measuring the incorporation of [$1-^{13}\text{C}$]acetate in VLDL-palmitate over time (Hellerstein & Neese, 1993). For that analysis 400 mg [$1-^{13}\text{C}$]acetate, 99% enrichment, in approximately 50 ml water was given orally every 45 min from 10.00–18.00 hours, and blood samples were obtained every 45 min from 10.00–18.00 hours and at 19.00, 20.00 and 21.00 hours. The fractional hepatic DNL (F) was determined at each time point as described above, and the rate constant determined by curve-fitting of the time-course of F to the equation $F = A(1 - e^{-K_s(t-L)})$, where A is the asymptotic value of F at isotopic steady-state, K_s the rate constant for VLDL synthesis, and L the lag-time for ^{13}C -incorporation in VLDL-fatty acids (Hellerstein & Neese, 1993). Total hepatic DNL was calculated as $0.75 \times K_s \times F \times \text{VLDL} \times \text{MM}_{\text{TG}}$ (Hellerstein & Neese, 1993), where VLDL is the size of the VLDL-pool (in mol), calculated as the mean of the non-fasting concentration of triacylglycerol in VLDL multiplied by the plasma volume (assumed to be 0.04 times the body weight), and MM_{TG} set at 850 as an average molecular mass for triacylglycerol to obtain the results in g lipid. The factor 0.75 takes into account the fraction of VLDL-fatty acids (25%) that are essential fatty acids and thus cannot be synthesised *de novo* (Hudgins *et al.* 1996). Total hepatic DNL calculated in this way was multiplied by 1.33 to correct for less than complete isotopic equilibration due to the relatively short time of isotope administration (L Hudgins, M Hellerstein, R Neese, J Hirsch, unpublished results). No correction was attempted for the probably lower contribution of DNL to fatty acids other than palmitate.

Blood glucose, insulin and leptin. The plasma used for the study of hepatic DNL was also used for determination of glucose (Kunst *et al.* 1984), insulin and leptin by radioimmunoassays according to the protocol of the manufacturer (Coat-A-Count[®] Insulin, Diagnostic Products Corporation, Los Angeles, CA, USA and Linco Research Inc., St Charles, MO, USA).

Statistical procedures

Results are presented as mean values with standard errors of the means, and *P*-values were calculated by Student's *t* test

or the Mann-Whitney test as appropriate. Curve fittings were performed by GraphPad Prism, version 2.01 (GraphPad Software, Inc., San Diego, CA, USA).

Ethics

Each subject received a written as well as verbal instruction concerning the study. The study was approved by the ethical committee of Fyns and Vejle Amt, and a letter of consent was signed by each subject.

Results

Energy intake

Habitual food intake is shown in Table 2. Alcohol consumption in some of the subjects was high in phase 1. The energy intake in phase 3 was in accordance with that intended; 78% from carbohydrates, 11% from fat and 11% from protein for the C-group and 31%, 58% and 11% for the F-group respectively. Alcohol was not included in the diet in phase 3, but if the subjects occasionally had visitors, a few drinks were allowed. The total intake is given as the mean value per day in Table 3. Energy intake was very similar in the C- and F-group during phase 3: 19.3 and 19.2 MJ/d respectively. On average the C-group had a surplus energy intake of 5.62 (SEM 0.50) MJ/d and the F-group of 4.78 (SEM 0.70) MJ/d, corresponding to an average increase of 37% in total energy intake.

The large amount of carbohydrate consumed by the C-group led to a substantial consumption of purified sucrose, from 20 to 36% of the total intake of carbohydrate. The percentage of energy from sucrose was 15–29 and thus within the range of sucrose consumption registered in Denmark (Danskernes kostvaner, 1996).

Anthropometry

Curve fitting of the time course of body weight and fat mass to linear-, power- or exponential functions showed that the time course was best described by a linear equation. Accordingly, changes in body weight, fat mass, fat free

Table 2. Energy intake per day during phase 1*

Subjects†	Energy intake (MJ/d)		Carbohydrate intake (MJ/d)		Fat intake (MJ/d)		Protein intake (MJ/d)		Alcohol intake (MJ/d)	
	C-group	F-group	C-group	F-group	C-group	F-group	C-group	F-group	C-group	F-group
Pair 1	18.6	18.5	9.3	8.7	5.0	0.2	2.0	2.1	2.2	0.8
Pair 2	15.2	15.2	8.9	6.9	4.0	4.8	2.0	2.6	0.3	1.5
Pair 3	12.2	14.5	6.3	6.1	4.5	3.6	1.6	1.4	1.1	3.5
Pair 4	12.2	15.6	7.0	6.8	3.4	6.4	1.5	2.4	0.2	0.0
Pair 5	8.9	11.2	5.1	5.5	2.2	3.2	1.1	1.3	0.5	1.3
Pair 6	13.5	13.0	7.1	6.9	4.4	3.9	1.7	1.7	0.2	0.4
Pair 7	15.1	11.4	10.4	5.9	2.7	3.8	2.0	1.5	0.0	0.2
Pair 8	10.3	10.3	5.0	4.1	2.5	2.0	1.2	1.0	1.6	3.1
Pair 9	14.4	20.1	8.3	11.2	4.0	6.6	1.6	2.3	0.6	0.0
Pair 10	16.2	13.9	8.7	6.8	4.1	3.7	2.1	2.1	1.3	1.4
Mean	13.7	14.4	7.6	6.9	3.7	3.8	1.7	1.8	0.8	1.2
SEM	0.9	0.9	0.5	0.6	0.3	0.6	0.1	0.2	0.2	0.4

*Phase 1 (prephase) lasted 10–14 d. For details of measurement of energy intake see pp. 234–235.

†Subjects (*n* 20) were paired according to habitual activity and body size. For details of subjects see Table 1.

Table 3. Total intake per day of energy, carbohydrate, fat, protein and alcohol in phase 3 and increase in energy intake from phase 1 to phase 3 (Δ Energy)*

Subjects†	Energy (MJ/d)		Carbohydrate (MJ/d)		Fat (MJ/d)		Protein (MJ/d)		Alcohol (MJ/d)		Δ Energy (MJ/d)	
	C-group	F-group	C-group	F-group	C-group	F-group	C-group	F-group	C-group	F-group	C-group	F-group
Pair 1	23.0	23.1	17.6	7.2	2.6	12.9	2.4	2.4	0.4	0.4	4.4	4.6
Pair 2	20.7	20.5	15.5	6.3	2.2	11.5	2.2	2.1	0.7	0.6	5.4	5.4
Pair 3	15.7	14.6	12.2	4.4	1.7	8.5	1.9	1.6	0.0	0.0	3.6	0.0
Pair 4	19.3	19.5	14.7	5.6	2.0	10.7	2.0	2.1	0.5	0.7	7.2	3.9
Pair 5	16.8	13.7	11.8	3.9	1.7	8.2	1.7	1.6	0.0	0.0	7.9	2.5
Pair 6	19.4	18.8	15.0	5.8	2.1	10.9	2.2	2.1	0.1	0.1	5.8	5.9
Pair 7	20.2	20.3	15.7	6.2	2.2	11.7	2.2	2.3	0.0	0.1	5.1	8.9
Pair 8	15.3	15.8	11.9	5.2	1.7	8.7	1.7	1.8	0.0	0.1	5.0	5.5
Pair 9	22.6	25.5	19.6	7.7	2.5	14.8	2.6	2.8	0.1	0.2	8.2	5.4
Pair 10	19.8	19.6	15.5	5.8	2.1	11.5	2.1	2.3	0.1	0.0	3.6	5.7
Mean	19.3	19.2	15.0	5.8	2.1	10.9	2.1	2.1	0.2	0.2	5.6	4.8
SEM	0.8	1.1	0.8	0.3	0.1	0.6	0.1	0.1	0.1	0.1	0.5	0.7

* Phase 1 (prephase) lasted 10–14 d; phase 3 (intervention period) lasted 21 d. For details of measurement of energy intake in phase 1 see pp. 234–235. In phase 3, energy intake was intended to be increased by 5 MJ above habitual intake as determined in phase 1 (see pp. 234–235).

† Subjects (n 20) were paired according to habitual activity and body size. For details of subjects see Table 1. For details of C- and F-group diets in phase 3 see pp. 234–235.

mass and total body water were calculated from the daily measurements of weight and impedance by linear regression analysis (Table 4). The C- and the F-group showed a significant increase in body weight and fat mass of about similar size, 1.5 and 0.9 kg respectively, with no statistically significant differences between the groups. A significant change of fat free mass was only observed in the C-group. No changes were observed in total body water.

Mean values of the changes in fat mass and fat free mass determined by weekly underwater weighing are also given in Table 4. These values were similar to those obtained by impedance measurements, although gains in fat free mass

were somewhat greater and gains in fat were somewhat lower than with impedance.

A considerable inter-individual variation in weight gain was observed in both groups, the CV being 97 and 82% in the C- and F-group respectively, despite an intended surplus energy intake of 5 MJ/d in all subjects. One reason for the variation in weight gain could be that some subjects under-reported energy consumption in phase 1. An indication of such under-reporting may be obtained by comparing the reported energy intake with the total energy expenditure as predicted by multiplication of BMR by a physical activity factor. With BMR calculated from age, height and weight of

Table 4. Changes in body composition (body weight, fat free mass, fat mass, and total body water) in phase 3 (intervention phase) using impedance and hydrostatic measurements†

Subjects‡	Δ Body weight (kg)		Δ Fat free mass (kg)		Δ Fat mass (kg)		Δ Total body water (kg)	
	C-group	F-group	C-group	F-group	C-group	F-group	C-group	F-group
Impedance measurements								
Pair 1	2.25	3.24	1.08	1.51	1.17	1.73	1.03	1.13
Pair 2	2.34	3.48	1.50	1.61	0.84	1.87	1.01	1.09
Pair 3	-0.10*	1.63	-0.02	0.46	0.12*	1.17	-0.25	0.15
Pair 4	2.58	2.91	1.25	1.25	1.33	1.66	0.95	0.99
Pair 5	-0.73*	-0.71	-0.79	-1.09	0.06*	0.38	-0.46	-0.57
Pair 6	-0.59*	1.33*	0.08	0.75	-0.66*	0.58*	-0.25	0.34
Pair 7	1.40	2.00	-0.03	0.50	1.42	1.50	0.00	0.00
Pair 8	2.05	-0.06*	1.07	-0.11	0.98	0.06*	1.05	-0.04
Pair 9	1.12	0.76*	0.49	0.32	0.64	1.08	0.27	0.04
Pair 10	3.17	1.22	1.51	0.43	1.66	0.79	1.28	0.61
Mean	1.35	1.58	0.61	0.50	0.76	1.08	0.46	0.37
SEM	0.42	0.41	0.23	0.25	0.22	0.19	0.20	0.17
P§	0.010	0.006	0.030	0.090	0.009	0.001	0.060	0.070
Hydrostatic measurements								
Mean			1.33	0.85	0.86	0.83		
SEM			0.33	0.28	0.49	0.32		

* Slope of regression line not significantly different from zero.

† For details of procedures see pp. 234–235.

‡ Subjects (n 20) were paired according to habitual activity and body size. For details of subjects see Table 1. For details of C- and F-group diets in phase 3 (21 d) see pp. 234–235.

§ P values for changes in impedance measurements were obtained by testing the changes for being significantly different from zero.

Table 5. Sleeping energy expenditure in phase 2 and in phase 3 (intervention phase)*

Subjects†	Energy expenditure (kJ/h)							
	Direct method				Indirect method			
	C-group		F-group		C-group		F-group	
	Phase 2	Phase 3	Phase 2	Phase 3	Phase 2	Phase 3	Phase 2	Phase 3
Pair 1	320	364	392	396	486	403	324	346
Pair 2	356	320	270	274	407	396	320	335
Pair 3	446	nd	266	nd	299	331	349	328
Pair 4	277	274	292	302	342	396	353	367
Pair 5	295	302	313	313	328	342	331	324
Pair 6	410	342	346	342	418	396	374	367
Pair 7	378	346	313	320	364	353	346	331
Pair 8	nd	302	nd	338	nd	331	nd	353
Pair 9	342	338	389	320	364	374	371	342
Pair 10	277	299	320	328	310	346	317	338
Mean	345	321	322	326	368	367	343	343
SEM	20	10	15	11	20	10	7	5

nd, not determined (technical problems).

* For details of procedures see pp. 234–235.

† Subjects (*n* 20) were paired according to habitual activity and body size. For details of subjects see Table 1. For details of C- and F-group diets in phase 2 (baseline phase, 2–3 d) and phase 3 (21 d) see pp. 234–235.

the subjects (Schofield, 1985) and a physical activity factor of 1.78 as determined on free-living subjects by the double-labelled water technique (Goldberg *et al.* 1991), there was no indication of under-reporting when the calculated means of the C- and the F-group were considered, but individual variation was large.

Energy expenditure

Mean SEE was measured by direct and indirect calorimetry and shown in Table 5. No changes over time in energy expenditure could be observed neither in the C-group nor in the F-group as a result of overfeeding (phase 2 *v.* phase 3). Direct and indirect measurements of energy expenditure in the experimental period were in agreement in the F-group, but in the C-group the results obtained by the indirect method were about 15 % higher in both phase 2 and phase 3 ($P < 0.002$).

Respiratory quotient

The calculated RQ of food, the measured RQ during sleep, and the calculated non-protein RQ are given in Table 6. Food RQ was slightly higher than the measured RQ during phase 3, both for the period as a whole and if comparisons were performed separately for week 1, 2 or 3 (data not shown). As expected, RQ was higher for the C-group than for the F-group. The measured values for RQ were stable from night-to-night and decreased on average by 0.01 ($P < 0.004$) during the night, when compared hour-by-hour.

Faeces, urine and nitrogen-balance

The total DM content of faeces in phase 3 was 44 % higher ($P = 0.001$) in the C-group than in the F-group (Table 7). Also, the energy loss in faeces was 30 % higher in the C-group ($P = 0.013$). On average the C-group lost 8.6 % and

Table 6. Respiratory quotients (RQ) in phase 3 (intervention phase) of ingested food, during sleep and during sleep corrected for protein oxidation*

Subjects†	RQ (food)		RQ (sleep)		Non-protein RQ	
	C-group	F-group	C-group	F-group	C-group	F-group
Pair 1	0.93	0.80	0.89	0.80	0.90	0.79
Pair 2	0.93	0.80	0.89	0.82	0.90	0.82
Pair 3	0.94	0.80	0.86	0.80	0.87	0.79
Pair 4	0.93	0.80	0.80	0.75	0.79	0.73
Pair 5	0.94	0.80	0.82	0.75	0.82	0.73
Pair 6	0.94	0.80	0.79	0.74	0.78	0.73
Pair 7	0.94	0.80	0.88	0.78	0.89	0.76
Pair 8	0.94	0.81	0.86	0.75	0.87	0.73
Pair 9	0.94	0.80	0.90	0.80	0.91	0.79
Pair 10	0.93	0.81	0.94	0.81	0.96	0.81
Mean	0.94	0.80	0.86	0.78	0.87	0.77
SEM	0.001	0.001	0.01	0.01	0.02	0.01

* For details of procedures see pp. 234–236.

† Subjects (*n* 20) were paired according to habitual activity and body size. For details of subjects see Table 1. For details of C- and F-group diets in phase 3 (21 d) see pp. 234–235.

Table 7. Loss in faeces of energy, dry matter, carbohydrate, fat and nitrogen during phase 3 (intervention phase)*

Subjects†	Energy (MJ/d)		Dry matter (g/d)		Carbohydrate (g/d)		Fat (g/d)		Protein (g/d)	
	C-group	F-group	C-group	F-group	C-group	F-group	C-group	F-group	C-group	F-group
Pair 1	2.05	1.86	97.00	74.62	61.24	37.10	8.76	21.29	27.02	16.21
Pair 2	1.86	1.12	83.67	46.62	48.81	21.71	12.05	10.10	22.79	14.79
Pair 3	1.32	0.97	59.67	40.71	32.00	19.90	7.29	8.38	20.39	12.44
Pair 4	1.71	1.66	77.48	62.86	32.76	25.00	18.00	21.43	26.68	16.40
Pair 5	1.23	0.75	53.90	32.00	28.52	15.62	8.86	5.81	16.47	10.57
Pair 6	1.66	1.32	72.43	51.95	38.29	22.05	12.24	14.48	21.92	15.42
Pair 7	1.66	1.47	76.38	61.00	43.86	29.14	8.24	13.43	24.30	18.46
Pair 8	1.43	0.96	67.90	43.05	42.76	22.71	5.76	5.43	19.33	14.88
Pair 9	1.80	1.25	82.19	55.38	48.19	30.52	10.14	8.29	23.84	16.59
Pair 10	1.35	1.02	63.38	42.90	36.67	20.90	4.95	9.00	21.78	12.99
Mean	1.61	1.24	73.40	51.11	41.31	24.47	9.63	11.76	22.45	14.88
SEM	0.08	0.11	4.04	3.99	3.10	1.96	1.20	1.84	1.03	0.73

* For details of procedures see p. 235.

† Subjects (n 20) were paired according to habitual activity and body size. For details of subjects see Table 1. For details of C- and F-group diets in phase 3 (21 d) see pp. 234–235.

the F-group 6.6% of the energy intake, a difference which was significant ($P=0.001$). The content of carbohydrate, fat and protein in faeces is also given in Table 7. The loss of carbohydrate and protein was significantly higher ($P=0.015$ and 0.0002 respectively) in the C-group, whereas the fat loss was the same in both groups.

The total intake of N, the loss in faeces and urine, and the derived N balance are shown in Table 8. The C- and F-groups had very similar protein intake. In the C-group the larger loss of N in faeces apparently was fully counterbalanced by a smaller loss in urine, resulting in a positive N balance of the same magnitude as in the F-group. Approximately 20% of the protein consumed was retained, resulting in an accumulation of about 430 g protein in both groups (cf. Table 8).

Hepatic de novo lipogenesis

Fractional hepatic DNL in phase 3 was 19.9 (SEM 1.5)% and 3.3 (SEM 0.8)% of total hepatic lipogenesis in the C-group

and F-group respectively (Table 9, column 1). The time-course of DNL in phase 3 is shown in Table 10. In the C-group, fractional hepatic DNL increased within the first week and remained at about 20% for the rest of the period. The fractional hepatic DNL in phase 2 was generally low except for C7, C10 and F10 (Table 10). These high values are not explicable by a habitual diet high in carbohydrate, except perhaps for C7 (Table 2), but may reflect large inter-individual differences in hepatic lipogenesis (Aarsland *et al.* 1996).

Absolute hepatic DNL was estimated from the rate constant (K_s) for VLDL synthesis as determined by the rate of [$1-^{13}C$]acetate incorporation in VLDL palmitate, the fractional hepatic DNL, and the plasma concentration of VLDL-triacylglycerol. K_s was 0.304 (SEM 0.051) h^{-1} as measured for C7 to C10 at the end and for C10 at the start of phase 3. This value was also used for the calculation of absolute DNL for C2 to C6. Furthermore, a constant rate of hepatic DNL equal to the rate in the measurement period (10.00–21.00 hours) was assumed throughout the day. Calculated in

Table 8. Nitrogen intake, loss in faeces and urine and nitrogen balance in phase 3 (intervention phase)*

Subjects†	Food nitrogen intake (g/d)		Faeces nitrogen loss (g/d)		Urine nitrogen loss (g/d)		N balance (g/d)‡	
	C-group	F-group	C-group	F-group	C-group	F-group	C-group	F-group
Pair 1	22.3	22.5	4.3	2.6	11.3	15.0	5.8	4.1
Pair 2	20.7	20.0	3.7	2.4	13.4	14.3	2.8	2.5
Pair 3	17.5	15.5	3.3	2.0	9.4	11.7	4.1	1.2
Pair 4	19.1	20.0	3.8	2.6	11.9	14.6	2.6	2.0
Pair 5	16.0	15.5	2.6	1.7	12.0	12.0	0.7	1.2
Pair 6	20.2	19.6	3.5	2.5	12.4	13.6	3.5	2.7
Pair 7	21.0	21.4	3.9	3.0	11.6	12.1	4.7	5.5
Pair 8	16.1	17.3	3.1	2.4	9.8	11.5	2.6	2.7
Pair 9	24.4	26.2	3.8	2.7	15.0	18.7	4.6	3.8
Pair 10	19.4	21.3	3.5	2.1	11.3	14.2	3.8	4.1
Mean	19.7	19.9	3.6	2.4	11.8	13.8	3.5	3.0
SEM	0.8	1.0	0.2	0.1	0.5	0.7	0.5	0.4

* For details of procedures see p. 235.

† Subjects (n 20) were paired according to habitual activity and body size. For details of subjects see Table 1. For details of C- and F-group diets in phase 3 (21 d) see pp. 234–235.

‡ Values are corrected for skin nitrogen loss (4% of protein intake; Bingham & Cummings, 1985).

Table 9. Fractional and absolute hepatic *de novo* lipogenesis (DNL) and calculation of total and extrahepatic lipogenesis*

Subjects†	DNL, hepatic fractional (%)		DNL, hepatic absolute (g/d) phase 3	DNL, hepatic absolute (g/d)† phase 3	Fat intake (g/d) phase 3	Faecal fat loss (g/d) phase 3	Fat oxidized (g/d) phase 3	Exogenous fat for storage (g/d) phase 3	Fat mass increase (g/d) phase 3	DNL total (g/d) phase 3	DNL extrahepatic (g/d) phase 3
	Phase 2	Phase 3									
C1	nd	24.5	nd	nd	68.7	8.8	22.2	37.7	55.7	18.0	nd
C2	4.0	28.1	10.2	13.6	59.2	12.0	20.2	26.9	40.2	13.3	-0.3
C3	4.9	19.6	6.7	8.9	45.5	7.3	20.4	17.8	5.6	-12.2	-21.1
C4	1.7	22.0	6.6	8.8	53.4	18.0	40.6	-5.2	63.4	68.6	59.7
C5	3.4	11.6	5.1	6.7	45.4	8.9	33.0	3.6	2.7	-0.9	-7.7
C6	2.6	20.4	9.8	13.0	56.3	12.2	47.8	-3.7	-31.5	-27.7	-40.7
C7	18.9	20.0	13.9	18.4	58.6	8.2	21.9	28.4	67.8	39.4	21.0
C8	8.4	19.2	8.5	11.3	44.7	5.8	23.9	15.0	46.6	31.6	20.3
C9	2.0	13.0	3.8	5.1	65.1	10.1	17.3	37.7	30.4	-7.3	-12.4
C10	17.3	20.9	3.4	4.6	55.8	5.0	7.1	43.7	79.0	35.3	30.8
Mean	7	20	7.6	10.1	55.3	9.6	25.4	20.2	36.0	15.8	5.5
SEM	2.1	1.5	1.1	1.4	2.6	1.2	3.8	5.6	10.9	9.1	9.6
F1-F10	5.6	3.3	-	-	287.8	11.8	43.2	232.8	51.4	-	-
Mean											

nd, not determined.

* For details of procedures see pp. 235–236.

† Values are those of preceding column corrected by 1.33 (correction for short isotope administration period); see p. 236.

‡ For details of subjects see Table 1. For details of C- and F-group diets in phase 2 (baseline phase, 2–3 d) and phase 3 (intervention phase, 21 d) see pp. 234–235.

this way, absolute hepatic DNL for phase 3 varied from 96 to 387 g (Table 9, column 3). These values are most likely overestimates, since the rate of hepatic DNL falls in the post-absorptive period (Davies *et al.* 1992; Hudgins *et al.* 1996).

Total de novo lipogenesis

At the whole-body level, DNL must have taken place if the amount of fat deposited is larger than the amount of fat available for storage. The latter can be calculated as the amount consumed minus the amount lost in faeces and minus the amount oxidised (Table 9, columns 4, 5 and 6). Fat oxidation was estimated from total non-protein energy expenditure and the non-protein RQ. In the calculations we have assumed that no oxidation of fat took place during the daytime (i.e. 16 h), since energy expenditure and RQ were only measured during sleep. This assumption is very unlikely to be true (even after heavy overfeeding with carbohydrate, fat oxidation did not stop totally (Clore *et al.* 1995)) and causes underestimation of total DNL. The possible overestimation of N balance (see p. 235) will give rise to an overestimation of total DNL of about 5%. Total DNL was positive in six and negative in four of the subjects in the C-group with a mean value of 332 g/21 d (Table 9, column 9).

Extrahepatic DNL (Table 9, column 10), calculated as total (column 9) minus hepatic DNL (column 3), was positive in four of the nine subjects with a mean value of 115 g/21 d.

Very-low density lipoprotein

In the C-group both the fasting and non-fasting values of blood VLDL concentration (μmol triacylglycerol/ml plasma) were significantly higher in phase 3 than in phase

Table 10. Fractional hepatic *de novo* lipogenesis (DNL) for the C- and F-group*

Subjects†	Phase 2	Phase 3		
		Week 1	Week 2	Week 3
C2	4.0	21.5	32.0	19.5
C3	4.9	21.0	14.2	23.6
C4	1.7	12.2	29.1	24.6
C5	3.4	8.0	16.6	10.3
C6	2.6	18.9	29.4	12.9
C7	18.9	18.2	18.7	23.2
C8	8.4	18.3	25.1	14.4
C9	2.0	15.9	12.4	10.8
C10	17.3	17.9	24.3	20.4
F2	9.3	2.8	3.3	2.2
F3	2.8	2.3	3.5	3.8
F4	4.2	3.4	4.7	6.1
F5	2.3	1.0	0.4	0.9
F6	1.6	1.9	2.1	2.5
F7	1.8	1.4	1.6	2.0
F8	3.0	0.6	0.7	0.0
F9	6.1	0.9	4.0	4.5
F10	19.6	2.4	2.5	9.7

* Fractional hepatic DNL was determined as described on p. 235 in phase 2 and at the end of each week of phase 3 (week 1, 2 and 3). Results are mean of values determined at 18.00, 19.00, 20.00 and 21.00 hours.

† For details of subjects see Table 1. For details of C- and F-group diets in phase 2 (baseline phase, 2–3 d) and phase 3 (intervention phase, 21 d) see pp. 234–235.

2 (fasting values 1.08 (SEM 0.18) and 1.88 (SEM 0.30), and non-fasting values 1.64 (SEM 0.30) and 2.11 (SEM 0.29) respectively, $P=0.007$ and 0.0006 respectively). The corresponding values for the F-group were 1.46 (SEM 0.12) and 1.07 (SEM 0.22), and 2.07 (SEM 0.22) and 1.95 (SEM 0.33) respectively, i.e. no significant changes from phase 2 to phase 3.

Glucose, insulin and leptin

Fasting and non-fasting glucose concentrations were within the range of 4.5–5.1 and 4.8–5.2 $\mu\text{mol/ml}$ plasma respectively. No significant differences between the C- and the F-group or between phase 2 and phase 3 were observed.

The fasting and non-fasting insulin concentrations were 9–12 and 23–31 $\mu\text{U/ml}$ plasma respectively. No significant differences between the C- and the F-group or between phase 2 and phase 3 were observed.

Leptin concentration in each subject was calculated as the mean value of measurements at 10.00, 18.00, 19.00, 20.00 and 21.00 hours in order to eliminate the effects of diurnal variation. The leptin concentration (ng/ml plasma) in the C-group was 2.4 (SEM 0.2) in phase 2 and increased significantly ($P=0.028$) in the intervention period to 3.2 (SEM 0.4). In the F-group the phase 2 values were similar to those of the C-group (2.5 (SEM 0.3)) but no significant changes could be observed in phase 3 (2.8 (SEM 0.3) at the end of phase 3). When data from both groups were pooled a significant positive correlation ($P=0.0046$) was observed between the increase in fat mass and the concentration of leptin (Fig. 1). This is in agreement with the findings of Considine *et al.* (1996), who found a strong positive correlation

between serum leptin concentrations and the percentage of body fat.

Discussion

The present study demonstrates that overfeeding of isoenergetic amounts of carbohydrate and fat led to nearly the same increase in body weight and fat mass; that carbohydrate overeating resulted in a 2–10-fold increase in the fractional hepatic DNL; that whole-body DNL in the subjects from the C-group was larger than hepatic DNL although highly variable among individuals, and that the amount of energy lost in faeces was about 30 % higher in the C-group than in the F-group.

Design of the study

The present study introduces a design with pairs of subjects acting as ‘Siamese’ twins in order to keep the physical activity similar in each pair during the non-sleeping period of the intervention. Subjects complied well to the twin-design. Staff members controlled the twin-condition both during the daytime, in the evenings and at the weekends, and only very seldom was one absent. Consequently, the differences in physical activity between the subjects of each pair can be considered eliminated as a factor influencing the general conclusions of the investigation.

The data on energy and macronutrient intake, energy expenditure during sleep, urine and faeces composition, body weight, fat mass and fat free mass during the intervention period were all collected on a daily basis over the course of the 3-week experimental period (phase 3). With

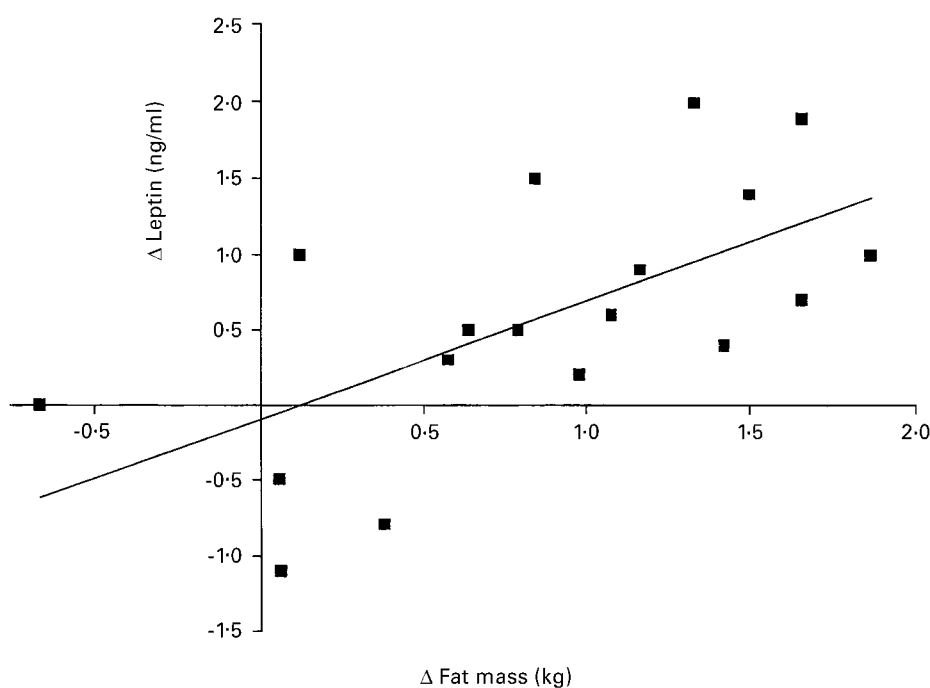


Fig. 1. Correlation between change in fat mass and change in plasma leptin concentration in normal male subjects (n 18) overfed by approximately 5 MJ/d for 21 d ($y = 0.79x - 0.10$, r^2 0.40, $P < 0.005$). For details of subjects see Table 1. Fat mass and leptin concentrations were determined as described on pp. 234–235. Leptin was not measured in two subjects (C1 and F1).

exception of the collection of urine and faeces, any random variation in these variables will tend to cancel out, thereby not affecting the overall results. However, the advantage of determination of many values for each data point does not apply to the baseline data on energy intake in phase 1 (the weighing and registering of all food eaten) and to the data obtained in phase 2, where body composition, energy expenditure at sleep, urine, faeces, and blood variables were only determined once.

Body composition

Weight gain. In the present investigation the increase in body weight (1.5 kg) showed a considerable variation, the CV being high (97 and 82 % in the C- and F-group respectively). No statistically significant difference in weight gain (1.5 kg) could be demonstrated between the C- and F-group. Similar results were obtained by Horton *et al.* (1995) (see p. 234). The results are in accordance with the concept that in the long-term only the energy content of the diet counts, and that the quality of the surplus of energy consumed is of little importance for the weight change. Supporting this notion are the findings of Golay *et al.* (1996) that underfeeding of normal or obese persons on a high-fat or an isoenergetic high-carbohydrate diet resulted in the same weight loss, and those of Schrauwen *et al.* (1997) that a change from a low-fat to a high-fat diet in normal persons did not change the body weight, provided that energy intake was unaltered. In contrast, it has been claimed that chronic imbalance between intake and oxidation is more likely if the surplus is given as fat (Bennett *et al.* 1992; Swinburn & Ravussin, 1993). This view is in accordance with the notion that recently ingested macronutrients are oxidised in the order protein and carbohydrate followed by fat (Jebb *et al.* 1996; Stubbs, 1998). Nevertheless, since there appears to be no hard evidence for a carbohydrate induced 'luxuskonsumption' (Stubbs, 1998; present study), an increased intake of carbohydrate would be expected to result in a correspondingly increased carbohydrate oxidation, which in turn would cause a larger part of the ingested fat to be deposited. A carbohydrate intake larger than the amount which can be oxidised will cause deposition of glycogen and/or fat by DNL.

In earlier studies of overfeeding most authors have found a CV in the weight gain of 15–25 % (Ravussin *et al.* 1985; Forbes *et al.* 1986; Bandini *et al.* 1989; Roberts *et al.* 1990; Diaz *et al.* 1992; Pasquet *et al.* 1992) whereas Horton *et al.* (1995) and Dallosso & James (1984) found variations similar to those of the present study. One reason for the large CV of the measured weight gain in the present study could be that weighing of the food eaten in phase 1 (*ad libitum* period) caused some of the subjects to eat less than usual. Furthermore, the relatively small weight gain in the present study combined with individual variation in tendency to deposit fat during overconsumption (Bouchard *et al.* 1990) may contribute to the large variation.

Fat and fat free mass. Fat mass increased significantly in both groups (Table 4), while only the C-group showed a significant increase in fat free mass, corresponding to 45 % of the increase in body weight, which is in accordance with earlier investigations (Forbes *et al.* 1986; Diaz *et al.* 1992;

Pasquet *et al.* 1992; Tremblay *et al.* 1992). In addition, the CV for measurements of fat mass and fat free mass were high. From the measured gain in fat free mass (by impedance) of about 0.5 kg, a N balance of 17 g N is expected, disregarding accumulation of glycogen. This is much less than found (Table 8). The reason for the discrepancy may be, at least partly, that collection of urine during phase 3 was not quantitative, in accordance with a large CV (40 %) for the daily volume of urine in the single individual (data not shown). This factor might well account for the approximately 3 g N/d losses not found here. Agreement between the measured gain in fat free mass and the N balance data would otherwise require a 30 % higher protein oxidation rate than was actually found.

Energy accounting

SEE did not change from phase 2 to phase 3. Roberts *et al.* (1990) in an overfeeding study found an increase of 24 h resting metabolic rate of 7 %, whereas the resting metabolic rate was unchanged during fasting. Other investigations (Dallosso & James, 1984; Ravussin *et al.* 1985) have estimated that overfeeding can be expected to give a 2–3 % increased BMR per kg of weight gain. In the present study this would correspond to BMR changes of 3–5 %, which is of the same order of magnitude as the accuracy of the energetic measurements and would therefore not be detected. There was a disagreement in data for energy expenditure by the indirect and direct method for the C-group, which presently is not understood. Since the data for the F-group agreed, the discrepancy is unlikely to stem from the analytical procedures.

Respiratory quotient. In the present experiment RQ was determined during sleep only. As expected RQ in the C-group was significantly higher than in the F-group. RQ_{sleep} did not change during the 21 d intervention, and remained smaller than RQ_{food} both in the C- and in the F-group. This is in contrast to the observation of Schrauwen *et al.* (1997) that the average daily RQ in subjects in energy balance equalled RQ_{food} 8 d after switching to a high-fat diet. This discrepancy can probably be related to the fact that RQ is lower during sleep than in the daytime (Dallosso & James, 1984; Lean & James, 1988) and that subjects in the present study were not in energy balance.

Loss of energy in faeces

Although the two diets were isoenergetic, the total loss of energy in faeces was significantly higher in the C-group (7.8 MJ/21 d; if not lost this would correspond to an increase in body mass of about 0.23 kg (7800/33.8; Forbes *et al.* 1986)).

There was no simple correlation between intake and loss in faeces of the various food components. Total intake of carbohydrate was 258 % of energy higher in the C-group than in the F-group. However, the loss in faeces of carbohydrate was only 169 % of energy higher in the C-group. The larger loss of carbohydrate in the C-group is in accordance with a larger intake of fibres in that group (5.25 g/MJ per d v. 2.66 g/MJ per d in the F-group). No significant difference was observed in the loss of lipid between the two

groups although the total lipid intake was 519 % of energy higher in the F-group. Furthermore, although protein intake was identical in the two groups, the C-group had a 51 % higher loss of protein in faeces than the F-group.

Hepatic de novo lipogenesis

The fraction of fatty acids in VLDL synthesised by DNL was about 20 % in the C-group and as expected very low in the F-group (Table 9) and did not change significantly during the last 2 weeks of phase 3 (Table 10). Fractional hepatic DNL was increased in all nine investigated C-subjects, even in C6 who lost fat mass, and the increase seems to be a response to consumption of carbohydrate, especially simple sugars (Hudgins *et al.* 1996, 1998). Most reported values of fractional hepatic DNL under conditions of 'eucaloric' energy intakes are below 5 % (Hellerstein, 1999), and under conditions of carbohydrate overfeeding are 20–40 % (Neese *et al.* 1995; Schwarz *et al.* 1995; Aarsland *et al.* 1997) as reported here.

Absolute hepatic DNL varied between 96 and 387 g per 21 d with a mean value of 211 g (Table 9). These values are maximum values (see p. 240). Aarsland *et al.* (1996) reported values of similar but somewhat lower magnitude (average value for five subjects corresponding to 60 g per 21 d, range 15–144 g).

Total and extrahepatic de novo lipogenesis

Total DNL was positive for six of the ten C-subjects (mean 332 g per 21 d) and showed large inter-individual variation (Table 9). As discussed on p. 240 the calculated values for whole-body DNL provide minimum values. Aarsland *et al.* (1997) estimated whole-body DNL after 4 d of massive enteral plus parenteral carbohydrate overfeeding, representing 2.5 times normal energy expenditure, and found whole-body DNL to be about 170 g/d, and on the basis of data from Pasquet *et al.* (1992) under massive overfeeding (i.e. 6000 kcal (25.1 MJ) as carbohydrate per d) a total DNL of at least 123 g/d can be inferred. The degree of carbohydrate overfeeding in the present study was not as extreme as in the studies of Aarsland *et al.* (1997) or Pasquet *et al.* (1992), however.

The values for total DNL are net values, i.e. the balance between oxidation and *de novo* synthesis. For these reasons the true rate of total and extrahepatic DNL is probably higher than indicated in Table 9. It can be concluded that DNL is sufficient to account for about 40 % of the increase in fat mass, and that extrahepatic DNL may account for about one-third of total DNL. Measurements of energy expenditure, RQ etc. during daytime would be desirable and would allow more precise conclusions regarding total and extrahepatic DNL.

It has been claimed that human subjects have a low capacity for DNL because of low activities of the enzymes involved in fatty acid synthesis (Acheson *et al.* 1982). Reported activities of fatty acid synthesis at 25°C are about 1 mU/mg protein in liver and adipose tissue of human subjects on a normal diet and somewhat higher in patients with long-term fat-free nutrition (Weiss *et al.* 1986). Assuming 1.5 kg liver with 200 mg protein/g wet weight

and 10 kg adipose tissue with 40 mg protein/g wet weight and that the enzyme activity at 37°C is twice that at 25°C, it can be calculated that 1 mU/mg protein corresponds to a total capacity for fatty acid synthesis of 16 g/d in the liver and 24 g/d in adipose tissue, i.e. more than sufficient to account for the values measured in the present study.

By the process of DNL from carbohydrate there is an obligatory energy loss as heat of about 19% as compared to about 3% by the direct storage of dietary fat (Flatt, 1978). Therefore an interesting question is whether heat production associated with the glucose-to-fatty acid conversion is in excess, i.e. causes a corresponding increase in the overall metabolic turnover, or whether this heat is compensated for by a correspondingly smaller metabolic rate. If the latter mechanism is in operation weight gain could occur on a 'per kilojoule' basis, independent of whether the source is fat or carbohydrate. If not, high carbohydrate intake may cause relatively less fat deposition. It should be noted that this contribution will be very small compared with the total energy turnover, and it would have been too small to detect. However, the present data indicate that weight gain and fat deposition caused by overconsumption may not be avoided by consuming a diet rich in carbohydrate and poor in fat.

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