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A Palaeolithic diet improves glucose tolerance more than a Mediterranean-like diet in individuals with ischaemic heart disease

Lindeberg S, Jönsson T, Granfeldt Y, Borgstrand E, Soffman J, Sjöström K, Ahrén B. Diabetologia 2007 Sep;50(9):1795-807. Epub 2007 Jun 22

Abstract

Aims/hypothesis Most studies of diet in glucose intolerance and type 2 diabetes have focused on intakes of fat, carbohydrate, fibre, fruits and vegetables. Instead, we aimed to compare diets that were available during human evolution with more recently introduced ones. Methods Twenty-nine patients with ischaemic heart disease plus either glucose intolerance or type 2 diabetes were randomised to receive (1) a Palaeolithic ('Old Stone Age') diet (n=14), based on lean meat, fish, fruits, vegetables, root vegetables, eggs and nuts; or (2) a Consensus (Mediterranean- like) diet (n=15), based on whole grains, low-fat dairy products, vegetables, fruits, fish, oils and margarines. Primary outcome variables were changes in weight, waist circumference and plasma glucose AUC (AUC Glucose₀₋₁₂₀) and plasma insulin AUC (AUC Insulin₀₋₁₂₀) in OGTTs. Results Over 12 weeks, there was a 26% decrease of AUC Glucose₀₋₁₂₀ (p=0.0001) in the Palaeolithic group and a 7% decrease (p=0.08) in the Consensus group. The larger (p=0.001) improvement in the Palaeolithic group was independent (p=0.0008) of change in waist circumference (-5.6 cm in the Palaeolithic group, -2.9 cm in the Consensus group; p=0.03). In the study population as a whole, there was no relationship between change in AUC Glucose_{0–120} and changes in weight (r=-0.06, p=0.9) or waist circumference (r=0.01, p=1.0). There was a tendency for a larger decrease of AUC Insulin₀₋₁₂₀ in the Palaeolithic group, but because of the strong association between change in AUC Insulin₀₋₁₂₀ and change in waist circumference (r=0.64, p=0.0003), this did not remain after multivariate analysis.

Conclusions/interpretation A Palaeolithic diet may improve glucose tolerance independently of decreased waist circumference.

Keywords Diet · Evolution · Glucose intolerance · Ischaemic heart disease · Palaeolithic diet · Type 2 diabetes

Abbreviations

BIA bioelectrical impedance analysis E% percentage of total energy intake HOMA-IR homeostasis model assessment of

insulin resistance

IFG impaired fasting glucose
IGT impaired glucose tolerance
IHD ischaemic heart disease
NGT normal glucose tolerance

Introduction

Impaired glucose tolerance (IGT) and type 2 diabetes are common risk factors for ischaemic heart disease (IHD) [1, 2]. which negatively affect the long-term prognosis after myocardial infarction [3, 4]. In fact, cross-sectional studies have found only 35–54% of IHD patients have normal glucose tolerance (NGT) [5–11]. Increased physical activity, healthy food choices and decreased waist circumference may help to lower the rate of progression from IGT to diabetes [12–14]. Standard dietary advice for patients with IHD and/ or IGT generally includes whole-grain cereals, low-fat dairy products, vegetables, fruits, legumes, oily fish and refined fats that are rich in monounsaturated fatty acids and alphalinolenic acid while low in transunsaturated fatty acids [15–17]. However, the optimal dietary treatment of IGT and insulin resistance is a matter of debate, including the preferred amounts and types of fat, carbohydrate and protein [16, 18–

21], and amounts of fruits [22] and sodium [23, 24].

Since nutritional science is hampered by confounders, an evolutionary approach has been suggested. It is postulated that foods that were regularly eaten during primate and human evolution, in particular during the Palaeolithic (the 'Old Stone Age,' 2.5-0.01 million years BP), may be optimal to prevent insulin resistance and glucose intolerance [25, 26]. A Palaeolithic diet includes lean meat, fish, shellfish, fruits, vegetables, roots, eggs and nuts, but not grains, dairy products, salt or refined fats and sugar, which became staple foods long after the appearance of fully modern humans. We found that traditional Pacific Islanders of Kitava, Papua New Guinea, had no signs of IHD, stroke or markers of the metabolic syndrome, possibly because of their traditional lifestyle [27–29]. In the present study, we examined the effect of dietary advice according to this Palaeolithic diet model on glucose tolerance and postchallenge insulin response in glucose-intolerant IHD patients recruited from a Coronary Care Unit, compared with dietary advice according to standard clinical practice. Our hypothesis was that the Palaeolithic diet would provide metabolic benefits beyond its nutrient composition.

Materials and methods

Participants The study was a 12 week controlled dietary intervention trial in 29 (out of 38 eligible) male IHD patients with waist circumference >94 cm and increased blood glucose or known diabetes, recruited from the Coronary Care Unit at Lund University Hospital, Sweden. We included patients with any of the following conditions: an ongoing acute coronary syndrome, a history of myocardial infarction diagnosed by creatine kinase MB isoenzyme or troponin elevation, percutaneous coronary intervention or coronary artery bypass surgery or

angiographically diagnosed coronary stenosis ≥30%. Exclusion criteria were BMI <20 kg/m2, serum creatinine >130 umol/l, poor general condition, dementia, unwillingness/inability to prepare food at home, participation in another medical trial, chronic inflammatory bowel disease, type 1 diabetes and treatment with hypoglycaemic agents, warfarin or oral steroids. Other drugs were not restricted, and treatment with statins and beta blockers was usually initiated and/or changed during the trial. Approval for the study was obtained from the regional Medical Ethics Committee, and all individuals gave written informed consent to participate in the study. In addition to the 29 patients who completed the trial, nine randomised subjects were excluded for the following reasons: worsening general condition (n=4), unwillingness to continue (n=3, all in the Palaeolithic group) or missing OGTT data (one in each group).

Procedure All eligible subjects were informed of the intention to compare two healthy diets and that it was unknown if either of them would be superior to the other with regard to weight reduction and improved glucose metabolism. Patients qualified for the study if they had known type 2 diabetes or, at a screening OGTT with 75 g glucose, a fasting capillary blood glucose ≥6.1 mmol/l or a 2 h capillary blood glucose ≥7.8 mmol/l. In 13 subjects, this screening OGTT was performed after an acute coronary episode (Table 1). The remaining 16 subjects, eight in each group, were recruited between 2 months and 2 vears after hospital discharge. Blood glucose concentrations were analysed in capillary whole blood immediately after collection with a HemoCue photometer (HemoCue, Ängelholm, Sweden). A second OGTT was performed within 2 weeks, when venous plasma samples were collected <5 min before and 30 and 120 min after ingestion of 75 g of glucose and analysed for glucose by the glucose

Table 1 Patient characteristics at baseline

	Group		p value
	Palaeolithic (<i>n</i> =14)	Consensus (<i>n</i> =15)	- · ·
Age (years)	65±10	57±7	0.01
Weight (kg)	92±11	96±12	0.3
$BMI (kg/m^2)$	29±4	30±2	0.3
Waist (cm)	106±8	107±8	0.8
Fasting plasma glucose (mmol/l)	6.8±1.3	7.1±1.8	0.6
2 h plasma glucose (mmol/l)	8.9 ± 1.8	8.8 ± 3.8	1.0
Glucose AUC (mmol/l × min)	1,104±116	1,145±298	0.6
HbA_{lc} (%)	4.8 ± 0.3	4.9 ± 0.8	0.6
IFG/IGT/diabetes (capillary), n	2/10/3	3/9/5	0.7
IFG/IGT/diabetes (OGTT), n	0/2/10	2/4/9	0.7
Fasting plasma insulin (pmol/l)	102±36	123±68	0.3
2 h plasma insulin (pmol/l)	988±570	674±532	0.14
ln HOMA-IR	0.62 ± 0.38	0.75 ± 0.53	0.5
Insulin AUC (nmol/l \times min)	81±41	70 ± 45	0.5
Systolic blood pressure (mmHg)	132±12	129±19	0.6
Diastolic blood pressure (mmHg)	77±9	78±11	0.7
Serum cholesterol (mmol/l)	4.2 ± 0.6	4.5±0.9	0.3
Serum triacylglycerols (mmol/l)	1.3±0.6	1.9 ± 0.8	0.06
C-reactive protein (µg/ml)	4.5 (0.8-88)	4.5 (0.8-43)	1.0
Study start <2 weeks			
after acute coronary syndrome (n)	6	7	0.8
number of days	4±2	4 ± 2	1.0
after statin treatment initiated (n)	6	6	0.9
after quitting smoking (n)	2	1	0.8
No statin treatment (n)	1	2	0.9
Smoking: never/ex- (n)	5/9	4/11	0.7

Values are means±SD for all continuous variables except C-reactive protein, for which values are geometric means (ranges)

oxidase technique and for insulin by RIA (Linco Research, St Charles, MO, USA). Normal plasma glucose was defined as a fasting venous plasma glucose <6.1 mmol/l and a 2 h venous plasma glucose <7.8 mmol/l. IGTwas defined as 2 h plasma glucose of 7.9−11.0 mmol/l and fasting plasma glucose <7 mmol/l, and diabetic levels as fasting plasma glucose ≥7.0 mmol/l or 2 h plasma glucose ≥11.1 mmol/l. Impaired fasting glucose (IFG) was defined as a fasting plasma glucose

6.1–6.9 mmol/l and a 2 h venous plasma glucose <7.8 mmol/l.

Diets Immediately after the second OGTT, subjects were randomised to one of two healthy diets: (1) a Consensus (Mediterranean-like) diet (n=15) based on whole-grain cereals, low-fat dairy products, potatoes, legumes, vegetables, fruits, fatty fish and refined fats rich in monounsaturated fatty acids and alphalinolenic acid; or (2) a Palaeolithic diet (n=14) based on lean meat, fish, fruits,

^aFor difference between groups

b<1 week between acute coronary syndrome (myocardial infarction and/or percutaneous coronary intervention) and dietary intervention

leafy and cruciferous vegetables, root vegetables (including restricted amounts of potatoes), eggs and nuts. All subjects were never-smokers or ex-smokers (Table 1), five of whom had stopped smoking ≤2 weeks prior to study start (three in the Palaeolithic group and two in the Consensus group). The others had stopped smoking >6 months ago. All subjects were informed individually (by S. Lindeberg, K. Sjöström or E. Borgstrand) during two 1 h sessions and were given written dietary advice and food recipes.

Only subjects in the Consensus group were informed of the possible benefits of Mediterranean-like diets rich in whole grains and about the Lyon Diet Heart Study [30]. The Consensus group was also educated by use of a dietary questionnaire for nutrition counselling ('20 questions') [31] used in a successful health promotion programme, 'Live For Life,' which led to lowered cardiovascular and total mortality in the Habo municipality, Sweden [32] (Supplementary Table 1). Only subjects in the Palaeolithic group were educated in the concept of evolutionary health promotion [33] and the potential benefits of a Palaeolithic diet. They were advised to increase their intake of lean meat, fish, fruits and vegetables and to avoid all kinds of dairy products, cereals (including rice), beans, sugar, bakery products, soft drinks and beer. The following items were

accepted in limited amounts for the Palaeolithic group: eggs (one or fewer per day), nuts (preferentially walnuts), potatoes (two or fewer medium-sized per day), rapeseed or olive oil (one or fewer tablespoons per day). The intake of other foods was not restricted and no advice was given with regard to proportions of food categories (e.g. animal vs plant foods).

The type of dietary advice given to the Consensus group was similar to the established programme at the Coronary Care Unit. Since the required increase in education intensity in order to match the Palaeolithic group was rather small, no 'usual care' control group was considered necessary. Advice about regular physical activity was given equally to the two groups. Both groups were advised not to consume more than one glass of wine per day.

Evaluation Changes in the AUC between 0 and 120 min during OGTT for plasma glucose (AUC Glucose₀₋₁₂₀) and plasma insulin (AUC Insulin₀₋₁₂₀) were predefined primary endpoints, along with changes in body weight and waist circumference. The base of the AUC was set at 0 mmol/l for glucose and 0 pmol/l for insulin. The computer-generated homeostasismodel assessment of insulin resistance (HOMA-IR) index, which has been suggested to provide a reasonable estimate of insulin

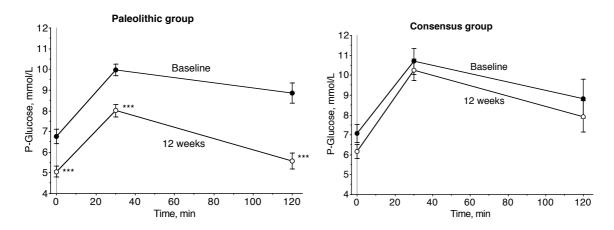


Fig 1 Plasma glucose during OGTTs at study start (baseline, closed circles) and after 12 weeks (open circles) in the Palaeolithic (a) and Consensus (b) groups. Values are means±SE. ***p<0.001.

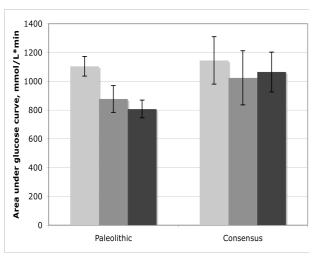


Fig. 2 Mean glucose AUCs (0–120 min) during OGTTs at study start (baseline, light grey columns) and after 6 weeks (dark grey columns) and 12 weeks (black columns) in the Palaeolithic and Consensus groups. Error bars denote 95% CIs.

resistance, was derived from fasting plasma glucose and insulin (www.dtu.ox.ac.uk) [34]. The early phase of post-challenge glucose and insulin responses were represented by the AUCs (Incremental AUC Glucose_{0–30} and Incremental AUC Insulin_{0–30}) during the first 30 min of the OGTT, using levels at 0 min as the base of the area.

A 4 day weighed food record on four consecutive days, including one weekend day, with weighing of each food item on a digital weighing scale (that could be set to zero), was completed by the participants, starting 15±5 days after initiating the

dietary change. Nutrients were calculated using Matsedel dietary analysis software (Kost och Näringsdata AB, Bromma, Sweden). Glycaemic load was calculated by multiplying the content of available carbohydrate in the serving of each food by the food's Glycaemic Index (with glucose as the reference) as given by Foster-Powell et al. [35]. Under-reporting was checked for by comparing food records with baseline weight and achieved weight loss, and by evaluating distribution and amount of consumed food. Body composition was estimated in a subset of 15 patients by use of leg-to-leg bioelectrical impedance analysis (BIA), using a Tanita Body Fat Analyzer (Model TBF 105; Tanita Corporation of America, Arlington Heights, IL, USA).

Statistics A pre-study power calculation showed that 12 subjects would be needed in each group in order to detect, with 80% power and at a significance level of 5%, a 20% reduction in AUC Glucose₀₋₁₂₀. Group assignment was made by use of minimisation, a restricted randomisation procedure which lowers the risk of baseline differences [36], using diabetes at screening (no/yes) and BMI (below or above 27 kg/m2) as restricting variables. A two-way paired t test was used to analyse within-subject differences in absolute values, while a two-way unpaired t test and repeated-measures ANOVA were used to

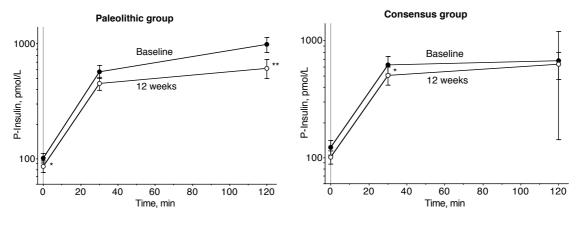


Fig 3 Plasma insulin during OGTTs at study start (baseline, closed circles) and after 12 weeks (open circles) in the Palaeolithic (a) and Consensus (b) groups. Values are means±SE. ***p<0.001.

analyse betweensubject differences in these changes. Simple and multiple linear regression was used to analyse univariate and bivariate relationships. All variables showed reasonable normal distribution in normal plots, but change in AUC Glucose_{0–120}, HOMA-IR and fruit intake showed perfect normal distribution only after ln transformation.

Results

The two groups differed at baseline only with regard to age being higher (p=0.01) and plasma triacylglycerols being lower (p=0.06) in the Palaeolithic group (Table 1). There was no relationship between age and any of the outcome variables at study start (Supplementary Tables 2, 3, 4, 5 and 6). During the 12 week dietary intervention, both groups decreased their waist circumference with a greater decrease in the Palaeolithic group (p=0.03; Table 2). Weight loss was on average 4.4 kg with no significant group difference.

In the Palaeolithic group, there was a 20% decrease in the OGTT AUC Glucose₀₋₁₂₀ during the first 6 weeks (p=0.0001), and an 8% decrease between weeks 6 and 12 (p=0.12; Figs 1 and 2, Table 2). In the Consensus group, a 10% decrease of AUC Glucose₀₋₁₂₀ was seen after the first 6 weeks (p=0.09) with no further change at 12 weeks (+4%, p=0.4), despite a further decrease of weight (p= 0.0003) and waist circumference (p=0.003). In the whole study population, there was no relationship between change in AUC Glucose₀₋₁₂₀ and changes in weight (r=-0.06, p= 0.9) or waist circumference (r=0.01, p=1.0) during the whole study period (Supplementary Table 6), which, consequently, did not explain the larger improvement of AUC Glucose₀_ ₁₂₀ in the Palaeolithic group (Supplementary Table 2).

In each group, AUC Insulin_{0–120} decreased during the first 6 weeks by 22%, but the decrease over 12 weeks was

significant only in the Palaeolithic group (Table 2, Fig. 3). After adjustment for waist loss, the tendency for a larger decrease of AUC Insulin₀₋₁₂₀ in the Palaeolithic group was no longer significant (Supplementary Table 3). Among the two groups combined, there was no association between change in AUC Glucose₀₋₁₂₀ and change in AUC Insulin₀₋₁₂₀ (r= 0.19, p=0.3), and thus the group difference in improvement of AUC Glucose₀₋₁₂₀ was independent of changes in AUC Insulin₀₋₁₂₀ (p=0.002) or ln HOMA-IR (p=0.0009; Supplementary Table 2).

Among secondary outcome variables, the most marked change was a 36% decrease in 2 h plasma glucose in the Palaeolithic group (from 8.9 to 5.6 mmol/l, p=0.0003; Table 3). In contrast, 2 h plasma glucose decreased by only 7% in the Consensus group (p=0.10), and the difference between the groups was highly significant. After 12 weeks, all 14 subjects in the Palaeolithic group had normal values, compared with 7 of 15 subjects in the Consensus group (p=0.0007 for group difference; Table 4). At 12 weeks, five subjects in the Consensus group still had diabetic values.

There was a decrease of HOMA-IR in both groups with no significant difference between the two groups (Table 4). The QUICKI index of insulin sensitivity [1/(In fasting plasma insulin+In fasting plasma glucose)] did not change more in the Palaeolithic group than in the Consensus group (p=0.23, data not shown). The early phase of postchallenge glucose and insulin responses, as represented by Incremental AUC Glucose₀₋₃₀ and Incremental AUC Insulin₀₋₃₀, did not change significantly during the trial, although a trend towards lowered Incremental AUC Insulin₀₋₃₀ was seen in both groups (Table 4).

Reported food composition differed between the two groups such that subjects in the Palaeolithic group had a much lower intake of dairy products, cereals and oil/ margarine, and a higher intake of fruits and

nuts (Table 5). The intake of vegetables, meat, meat products or fish did not differ significantly between the groups. Total fat intake was low with no difference between the groups (Table 6). Absolute protein intake was identical in the two groups while relative protein intake (as a percentage of total energy intake [E%]) was higher in the Palaeolithic group. Absolute carbohydrate intake was 43% lower in the Palaeolithic group, and 23% lower in terms of E%. Glycaemic load was 47% lower in the Palaeolithic group and correlated strongly with cereal intake (r=0.75, p<0.0001).

Energy intake was 25% lower in the Palaeolithic group (p=0.004; Table 6) despite similar quantities of consumed food (by weight; Table 5). After adjustment for energy intake, the improvement of AUC Glucose₀₋₁₂₀ was still larger in the Palaeolithic group (p=0.02; Supplementary Table 2), while the larger waist loss, and the tendency for larger decrease of AUC Insulin₀₋₁₂₀, compared with the Consensus group, disappeared (Supplementary Table 3).

In post hoc analysis among the whole study population, a positive association between intake of cereals and change in waist circumference explained 42% of waist loss among the whole study population (p=0.0003; Supplementary Table 6), and 40% in the Consensus group alone (p=0.016). In contrast, there was a negative correlation between fruit intake and change in waist circumference, which explained 21% of waist loss (p=0.01). Each of these associations remained significant after adjustment for dietary assignment, energy intake, carbohydrate intake or glycaemic load (Supplementary Table 5). Thus, waist loss increased with increasing intake of fruits and decreasing intake of cereals, associations which explained most of the group difference in waist loss. Compared with waist change, weight change was generally less clearly associated with dietary assignment and

other variables (Table 2, Supplementary Tables 2, 3, 4, 5 and 6).

Glycaemic load was positively associated with changes in waist (r=0.52, p=0.008) and AUC Glucose₀₋₁₂₀ (r= 0.50, p=0.01) but not with change in AUC Insulin₀₋₁₂₀ (r= 0.30, p=0.15). When glycaemic load and dietary assignment were entered simultaneously as explanatory variables in bivariate linear regression, neither of these was significantly associated with change in AUC Glucose_{0–120} (Supplementary Table 2). In forward stepwise linear regression with glycaemic load and dietary assignment as independent variables, only dietary assignment was associated with change in AUC Glucose₀₋₁₂₀ (data not shown).

None of the other effects, nor lack of effects, of group assignment on primary outcome variables (changes in weight, waist, AUC Glucose_{0–120} and AUC Insulin $_{0-120}$) was essentially altered after adjustment for age or baseline levels of weight, waist, glucose, insulin, AUC Glucose₀₋₁₂₀ or AUC Insulin₀₋₁₂₀, nor after adjustment for intake (g/day or E%) of carbohydrate, protein, total fat, saturated fat, monounsaturated fat, polyunsaturated fat, fibre or sodium. Repeated-measures ANOVA gave similar results for primary and secondary outcome variables, and addition of baseline values as covariates confirmed the independent effect of Palaeolithic diet on improvement of glucose tolerance (data not shown). Serum lipids changed to a similar extent in both groups, due to initiation of statin treatment in most patients, and there was no decrease in blood pressure (data not shown). The impact of medication was not analysed.

Among the 15 subjects who underwent BIA for body composition, change of fat mass did not differ between the groups (Table 7), and it explained 50% of weight change (p=0.002). In this subset of patients, change in fat mass explained <1% of change in AUC Glucose₀₋₁₂₀.

Discussion

We found marked improvement of glucose tolerance after advice to eat a Palaeolithic diet, based on lean meat, fish, fruits, vegetables, root vegetables, eggs and nuts as staple foods, while avoiding cereals, dairy products, refined fat, sugar and salt. Control subjects, who were advised to follow a Consensus (Mediterranean-like) diet based on whole grains, low-fat dairy products, fish, fruits and vegetables, did not significantly improve their glucose tolerance despite decreases in weight and waist circumference. The more pronounced improvement of glucose tolerance in the Palaeolithic group was unrelated to weight loss or decrease in waist circumference. In contrast, the insulin response changed more as a result of change in waist circumference than of dietary assignment or food choice.

The higher drop-out rate in the Palaeolithic group (three vs none) does not appear to be an important source of bias. If we assume no change in primary outcome variables in any of the drop-out subjects, had they finished the trial, we would still have found larger decreases in AUC Glucose_{0–120} (p= 0.01) and 2 h glucose (p=0.02) in the Palaeolithic group than in the Consensus group.

It is conceivable, but not very likely, that the more pronounced improvement of glucose tolerance in the Palaeolithic group was due to higher motivation (rather than different food patterns). We were meticulous in our efforts not to give the subjects in the Consensus group a feeling of belonging to a control group. Thus, we told eligible persons that we were to compare two healthy diets, not knowing which was the better one. We informed all subjects individually of the presumed benefits of their respective diet (but not of those of the other diet) during two 1 h sessions, and all subjects were provided with recipes and written dietary advice of equal length. During the 12 week trial,

waist circumference decreased more in the Palaeolithic group, but this did not explain the more pronounced decrease in fasting and post-prandial plasma glucose in these subjects.

Among the whole study population, change in AUC Glucose₀₋₁₂₀ was not related to changes in weight or waist circumference. Considering the large variation in weight loss (between -10.7 and +1.3 kg), and in light of earlier studies showing weight loss to be the major determinant of improved glucose tolerance [12], this lack of relationship is unexpected. In the Diabetes Prevention Project, weight loss was the dominant predictor of reduced diabetes incidence among glucose-intolerant subjects who were randomised to lifestyle modification [37]. However, weight change does not explain all of the improvement in glucose tolerance in such trials, and in a metaanalysis on the efficacy of lifestyle education to prevent type 2 diabetes in high-risk individuals, four out of eight trials did not find any effect on 2 h plasma glucose despite significant weight loss [12]. Furthermore, in epidemiological studies most of the variation in glucose tolerance among the general population is not explained by adiposity [38]. Therefore, an improvement in glucose tolerance that is independent of weight change is not entirely unexpected.

There was no apparent influence of dietary assignment on the HOMA-IR index of insulin sensitivity, and adjustment for changes in waist circumference or body weight eliminated the tendency towards larger decrease of AUC Insulin₀₋₁₂₀ in the Palaeolithic group. This is in contrast to a recent feeding trial in pigs, where we found markedly lower insulin response by the frequently sampled IVGTT, independent of body weight, after 15 months of a cerealfree Palaeolithic diet. compared with a cereal-based swine feed [39]. This discrepancy may be due to the use of frequently sampled IVGTT in the study in pigs, which gives a more precise

measure of insulin sensitivity than that of the present study (HOMA-IR). Since we did not perform euglycaemic insulin clamp measurements, the gold standard for assessing whole-body insulin sensitivity, we may have missed a significant effect on insulin sensitivity. If not, our findings add to the evidence that reduction of waist circumference is more important than dietary composition for the treatment of insulin resistance [19].

The very low reported energy intake in the Palaeolithic group, as calculated from 4 day weighed food records (registered early in the trial), does not necessarily imply under-reporting of food intake. An energy deficit of 4 MJ/day would be expected to cause a weight loss of 1 kg/week in the second and third months of energy restriction, and even more during the first month [40]. Thus, assuming a pre-study energy intake of at least 10 MJ/day, the reported in-trial energy intake is actually higher than expected from the observed weight loss, even in the Palaeolithic group. In addition, the similar weight loss in the two groups is not incompatible with different energy intakes. In this context, the laws of thermodynamics need to be considered thoroughly. These laws state that energy is constant and cannot be destroyed. However, they also state that energy can take various forms, including heat, and that conversion from one form of energy to another is more or less efficient [41]. Highly relevant, then, is the finding in animal experiments of decreased body temperature on low-calorie diets [42, 43]. Accordingly, identical weight loss on different energy intakes does not violate the laws of thermodynamics [41].

It is important to separate glycaemic control, as measured by HbA1c, from glucose tolerance. A habitual diet which reduces the post-prandial glucose response, such as a low glycaemic load diet, can reduce the metabolic consequences of glucose intolerance, including delaying the manifestation of diabetes, without necessarily improving glucose tolerance

itself [19, 44]. Although we cannot rule out glycaemic load as an important factor for glucose tolerance, our finding that the effect of Palaeolithic diet on glucose tolerance was independent of carbohydrate intake agrees with earlier studies which do not indicate a beneficial effect of carbohydrate restriction on glucose tolerance [20, 45–47].

The high fruit intake in the Palaeolithic group, almost sevenfold higher than the median intake among Swedish men (75 g/day) [48], and twice as high as in the Consensus group, should also be viewed against this background. Despite large variation in fruit intake (range 160–1,435 g/day in the Palaeolithic group and 53–679 g/day in the Consensus group), it was not associated with change in AUC Glucose₀-₁₂₀ (r=-0.02, p=0.9) or AUC Insulin₀₋₁₂₀ (r=-0.02, p=0.9) and did not explain the effects of group assignment on these outcome variables. Furthermore, a high fruit intake was associated with larger waist loss. Thus, our study lends no support to the notion that fruit intake should be restricted in patients with diabetes or glucose intolerance.

This is, to the best of our knowledge, the first controlled study of the effects of an ancestral human diet in patients with IGT or diabetes. In a non-controlled study of ten Australian Aborigines with diabetes and a mean BMI of 27 kg/m2, O'Dea et al. found that reversion to a hunter–gatherer lifestyle during 7 weeks led to 10% weight loss and reductions in fasting and 2 h glucose of 45 and 36% (p<0.0001 for all) [49]. Fasting insulin decreased by 48% (p<0.0001), while 2 h insulin did not change (+20%, not significant). Both diet and physical activity changed markedly, which precludes evaluation about the isolated role of diet. In contrast, in a similar study on healthy Australian Aborigines by the same authors, the insulin response to 70 g of starch from white bread (and butter) was reduced, while the glucose response was not, after 10-12

weeks of reversion to a traditional lifestyle [50].

In conclusion, we found marked improvement of glucose tolerance in IHD patients with increased blood glucose or diabetes after advice to follow a Palaeolithic diet compared with a healthy Western diet. The larger improvement of glucose tolerance in the Palaeolithic group was independent of energy intake and macronutrient composition, which suggests that avoiding Western foods is more important than counting calories, fat, carbohydrate or protein. The study adds to the notion that healthy diets based on wholegrain cereals and low-fat dairy products are only the second best choice in the prevention and treatment of type 2 diabetes

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Duality of interest The authors declare that there is no duality of interest associated with this manuscript.

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 Table 2
 Primary outcome variables

	Group		_ p value ^a
	Palaeolithic (<i>n</i> =14)	Consensus (<i>n</i> =15)	
Weight (kg)			
Baseline	91.7±11.2	96.1±12.4	0.3
6 weeks	88.0 ± 10.7	93.6±12.8	0.2
Change 0-6 weeks	-3.7 ± 2.2	-2.5 ± 2.3	0.2
95% CI	-4.9 to -2.4	-3.8 to -1.2	
p value for change within group	0.0001	0.0009	
12 weeks	86.7±11.3	92.2±12.9	0.2
Change 6-12 weeks	-1.4 ± 2.1	-1.3 ± 1.1	0.9
95% CI	−2.6 to −0.1	-1.9 to -0.7	
p value for change within group	0.03	0.0003	
Change 0-12 weeks	-5.0 ± 3.3	-3.8 ± 2.4	0.3
95% CI	-6.9 to -3.1	-5.2 to -2.5	
p value for change within group	0.0001	0.0001	
Waist circumference (cm)			
Baseline	105.8±7.6	106.6±8.0	0.8
6 weeks	102.8±7.8	105.2±8.8	0.5
Change 0-6 weeks	-3.0 ± 1.8	-1.5 ± 2.0	0.04
95% CI	−4.0 to −2.0	-2.7 to -0.2	
p value for change within group	0.0001	0.02	
12 weeks	100.2±7.7	103.6±8.6	0.11
Change 6-12 weeks	-2.6 ± 2.4	-1.5 ± 1.8	0.2
95% CI	−3.9 to −1.2	-2.7 to -0.7	
p value for change within group	0.001	0.003	
Change 0-12 weeks	-5.6 ± 2.8	-2.9 ± 3.1	0.03
95% CI	-7.2 to -3.9	-4.8 to -1.1	0.00
p value for change within group	0.0001	0.004	
AUC ^b Glucose ₀₋₁₂₀ (mmol/l × min)	0.0001	0.001	
Baseline	1,104±118	1,145±298	0.6
6 weeks	877±161	1,024±339	0.15
Change 0-6 weeks	-220 ± 206	-120±255	0.13
95% CI	-339 to -101	-262 to +21	0.5
p value for change within group	0.002	0.09	
12 weeks	807±107	1,065±250	0.001
Change 6-12 weeks	-70±156	+41±179	0.001
95% CI	-70±130 -160; +20	-59; +140	0.09
	0.12	-39, +140 0.4	
p value for change within group	-290±143	-80±168	0.001
Change 0-12 weeks			0.001
95% CI	-373 to -208	-173 to +13	
p value for change within group	0.0001	0.09	
AUC^b Insulin ₀₋₁₂₀ (nmol/l × min)	00.5.41.1	60.7.44.7	0.5
Baseline	80.5±41.1	69.7±44.7	0.5
6 weeks	63.1±30.0	54.1±37.2	0.5
Change 0-6 weeks	-17.4 ± 27.7	-15.5±16.9	0.8

95% CI	-33.4 to -1.3	−24.9 to −6.2	
p value for change within group	0.04	0.003	
12 weeks	56.1±30.1	60.4±46.4	0.8
Change 6-12 weeks	-7.0 ± 16.9	$+6.2\pm25.8$	0.12
95% CI	-16.7 to $+2.8$	-8.1 to $+20.5$	
p value for change within group	0.15	0.4	
Change 0-12 weeks	-24.3 ± 28.4	-9.3 ± 23.3	0.13
95% CI	-40.7 to -8.0	-22.2 to $+3.6$	
p value for change within group	0.007	0.14	

Values are means±SD

^aFor difference between groups ^bAUC for glucose and insulin response to a 75 g OGTT. The base of the AUC was set at 0 mmol/l for glucose and 0 nmol/l for insulin

Table 3 Glucose and insulin responses to OGTTs (secondary outcome variables) during the trial

trial			
	Group		p value ^a
	Palaeolithic (<i>n</i> =14)	Consensus (<i>n</i> =15)	_
Fasting plasma glucose (mmol/l)			_
Baseline	6.8±1.3	7.1 ± 1.8	0.6
6 weeks	5.2 ± 1.1^{b}	$5.8 \pm 1.2^{\circ}$	0.2
12 weeks	5.1±1.0	6.2 ± 1.4	0.02
Change 0-12 weeks	-1.7 ± 1.7	-0.9 ± 1.8	0.2
95% CI	-2.7 to -0.7	-1.9 to $+0.08$	
p value for change within group	0.003	0.07	
30 min plasma glucose (mmol/l)			
Baseline	10.0±1.1	10.7 ± 2.4	0.3
6 weeks	8.4 ± 1.6^{b}	9.8 ± 3.3	0.16
12 weeks	8.0±1.1	10.3 ± 2.1	0.001
Change 0-12 weeks	-2.0 ± 1.2	-0.4 ± 1.6	0.008
95% CI	-2.7 to -1.3	-1.3 to $+0.5$	
p value for change within group	0.0001	0.3	
120 min plasma glucose (mmol/l)			
Baseline	8.9±1.8	8.8 ± 3.8	1.0
6 weeks	6.6 ± 1.5^{b}	7.8 ± 4.1	0.3
12 weeks	$5.6 \pm 1.5^{\circ}$	7.9 ± 3.1	0.01
Change 0-12 weeks	-3.3 ± 1.9	-0.9 ± 2.0	0.003
95% CI	-4.4 to -2.2	-2.0 to $+0.2$	
p value for change within group	0.0001	0.10	
Fasting plasma insulin (pmol/l)			
Baseline	102±36	123±68	0.3
6 weeks	91±32	100±45	0.5
12 weeks	86±36	101±53	0.4
Change 0-12 weeks	-16 ± 27	-22 ± 54	0.7
95% CI	-32 to -0.3	-51 to +8	
p value for change within group	0.047	0.15	
30 min plasma insulin (pmol/l)			
Baseline	575±290	625±416	0.7
6 weeks	503±222	516±393	0.9
12 weeks	453±226	507±355	0.7
Change 0-12 weeks	-121 ± 230	-118 ± 202	1.0
95% CI	-254 to +12	−230 to −6	
p value for change within group	0.07	0.04	
120 min plasma insulin (pmol/l)			
Baseline	988±570	674±532	0.14
6 weeks	702±423°	482±374 ^b	0.15
12 weeks	615±443	631±633	1.0
Change 0-12 weeks	-374 ± 408	-42 ± 408	0.04
95% CI	−609 to −138	-268 to +183	
p value for change within group	0.005	0.7	

Values are means±SD

^aFor difference between groups

 $^{^{}b}p<0.01$ and $^{c}p<0.05$ by paired t test for change within group (6 week level is compared with baseline and 12 week level is compared with 6 week level)

 Table 4 Other glucometabolic variables in the two groups

	Group		p value ^a
	Palaeolithic (<i>n</i> =14)	Consensus (<i>n</i> =15)	
HbA_{lc} (%)			
Baseline	4.76 ± 0.26	4.89±0.79	0.6
6 weeks	4.61 ± 0.25^{b}	4.84 ± 0.72	0.3
12 weeks	4.64 ± 0.22	4.85±0.69	0.3
Change 0-12 weeks	-0.13 ± 0.26	-0.03 ± 0.39	0.4
95% CI	-0.28 to $+0.02$	-0.24 to $+0.17$	
p value for change within	0.09	0.7	
group			
Normal glucose levels ^c (n)			
Baseline	2	2	0.8
6 weeks	10	10	0.7
12 weeks	14	7	0.0007
Diabetic glucose levels ^d (<i>n</i>)			
Baseline	10	9	0.4
6 weeks	1	3	0.2
12 weeks	0	5	0.01
ln HOMA-IR	v		0.01
Baseline	0.62 ± 0.38	0.75 ± 0.53	0.5
6 weeks	0.47 ± 0.33^{b}	$0.55\pm0.42^{\rm b}$	0.6
12 weeks	0.39 ± 0.36	0.55±0.46	0.3
Change 0-12 weeks	-0.24±0.29	-0.19±0.36	0.7
95% CI	-0.40 to -0.07	-0.39 to +0.01	0.7
p value for change within	0.01	0.03	
group	0.01	0.03	
Insulin/Glucose ₀₋₃₀			
Baseline	172±125	145±110	0.5
6 weeks	172±125 135±61	133±144	1.0
12 weeks	139±72	112±126	0.5
Change 0-12 weeks	-33 ± 94	-33±71	1.0
95% CI	-87 to +21	-33 ± 71 -73 to +8	1.0
p value for change within	0.2	0.11	
_	0.2	0.11	
group Incremental Glucose AUC _{0.30} e			
Baseline	48±20	54.20	0.4
		54±20	0.4
6 weeks	48±19	60±40	0.3
12 weeks	44±20	62±26	0.06
Change 0-12 weeks	-4±24	+7±21	0.19
95% CI	-18 to +10	-4 to +19	
p value for change within	0.6	0.2	
group			
Incremental Insulin AUC ₀₋₃₀ ^e	5 1.40	7.5.50	0.0
Baseline	7.1±4.0	7.5±5.9	0.8
6 weeks	6.2 ± 3.0	6.2 ± 5.6	1.0

12 weeks	5.5±2.9	6.1±4.8	0.7
Change 0-12 weeks	-1.6 ± 3.1	-1.5 ± 2.7	0.9
95% CI	-3.4 to $+0.2$	-3.0 to $+0.07$	
p value for change within	0.08	0.06	
group			

Values are means±SD

^aFor difference between groups

 $^{^{}b}p$ <0.05 by paired t test for change within group (6 week level is compared with baseline and 12 week level is compared with 6 week level)

^cFasting venous plasma glucose ≤6.0 mmol/l and 2 h venous plasma glucose <7.8 mmol/l at OGTT (despite increased capillary blood glucose at screening)

^dFasting venous plasma glucose ≥7.0 mmol/l or 2 h venous plasma glucose ≥11.1 mmol/l at OGTT

 $^{^{\}rm e}$ Incremental AUC_{0.30}, incremental AUC during the first 30 min of OGTT, using levels at 0 min as the base of the area

Table 5 Diet composition (g/day) in the two groups, as estimated from 4 day weighed food records

	Group		p
	Palaeolithic (<i>n</i> =14)	Consensus (<i>n</i> =15)	_ value ^a
Fruit	493±335	252±179	0.03
Vegetables ^b	327±233	202±88	0.07
Potatoes	51±42	77±78	0.3
Nuts	11±12	2±6	0.02
Meat, fresh	143±95	97±67	0.16
Meat products	65±59	58±49	0.8
Fish	119±92	77±56	0.16
Eggs	29±23	19±18	0.21
Beans, peas	8±21	15±26	0.5
Cereals	18±52	268±96	0.0001
Milk and dairy products	45±119	287±193	0.0006
Oil, margarine ^c	1±3	16±11	0.0001
Sauce	2±6	25±31	0.02
Pastry	1±3	13±25	0.12
Jam	1±3	6±10	0.12
Total amount of food	1,311±598	1,382±222	0.7
Wine	59±63	37±51	0.3
Beer, light ^d	11±27	27±47	0.3
Sweet beverages (excluding juice)	18±46	53±90	0.2
Juice	38±75	88±141	0.3

Values are means±SD

^aFor difference between groups

^bIncluding root vegetables (but excluding potatoes and beans with pods)

^cButter was not reported to be consumed by anyone

^dStronger beer or liquor was not consumed, as reported

Table 6 Daily intake of macronutrients, dietary fibre, cholesterol, sodium, potassium, magnesium and calcium in the two groups, as estimated from 4 day weighed food records.

	Group			
	Palaeolithic (<i>n</i> =14)	Consensus (<i>n</i> =15)	_	
Energy				
MJ	5.6 ± 2.2	7.5 ± 1.3		
kcal	1,344±521	1,795±306	0.01	
Protein				
Weight (g)	90±41	89±20	0.9	
g/kg body weight	0.98 ± 0.4	0.95 ± 0.2	0.8	
E%	27.9 ± 6.8	20.5±3.6	0.002	
Total fat				
Weight (g)	42±20	50±13	0.2	
g/kg body weight	0.44 ± 0.2	0.55 ± 0.2	0.12	
E%	26.9 ± 6.4	24.7±4.3	0.3	
Fatty acids				
Saturated (g)	11.5±4.8	16.8 ± 4.2	0.005	
E%	7.7 ± 2.4	8.3±1.7	0.4	
Monounsaturated (g)	16.3 ± 7.4	19.0±5.0	0.3	
E%	10.7 ± 2.6	9.4±1.9	0.2	
Polyunsaturated (g)	9.6±7.5	9.0 ± 3.0	0.8	
E%	5.8 ± 2.5	4.4±1.1	0.06	
Carbohydrate				
Weight (g)	134±56	231±48	0.0001	
g/kg body weight	1.4 ± 0.6	2.5 ± 0.6	0.0001	
E%	40.2 ± 8.3	51.7±5.3	0.0002	
Glycaemic load ^b	65±30	122±28	0.0001	
Alcohol (E%)	3.9 ± 4.4	2.3 ± 3.0	0.3	
Fibre (g)	21.4±13.2	26.8 ± 7.4	0.2	
Cholesterol (mg)	397±192	295±122	0.11	
Salt (g)				
Sodium	1.9±0.6	2.9 ± 0.7	0.0006	
Sodium chloride	4.7±1.6	7.2 ± 1.7	0.0006	

Values are means±SD

^aFor difference between groups

^bThe glycaemic index (with glucose as the reference food) multiplied by the amount of carbohydrate

Table 7 Leg-to-leg BIA in a subset of patients (n=15)

	Group		p
	D-11'41.'- (0)	C(7)	_ value ^a
	Palaeolithic (<i>n</i> =8)	Consensus (<i>n</i> =7)	
Fat mass (kg)	20.7.5.4	22.0.0.6	0.0
Baseline	28.7±5.4	33.0±8.6	0.3
6 weeks	$26.5 \pm 4.5^{\text{b}}$	31.7±8.5 ^b	0.16
12 weeks	$24.9 \pm 4.5^{\text{b}}$	30.8±8.7	0.12
Change 0-12 weeks	-3.9 ± 2.9	-2.3 ± 1.0	0.18
95% CI	-6.3 to -1.5	-3.2 to -1.4	
p value for change within	0.007	0.0009	
group			
Fat mass (% of body mass)			
Baseline	30.0 ± 3.0	32.6±5.7	0.3
6 weeks	28.9 ± 2.7	31.8 ± 5.6^{b}	0.2
12 weeks	$27.4 \pm 2.7^{\circ}$	31.0±5.9	0.14
Change 0-12 weeks	-2.6 ± 2.3	-1.6 ± 0.6	0.3
95% CI	-4.5 to -0.7	-2.2 to -1.1	
p value for change within	0.02	0.0004	
group			
Fat free mass (kg)			
Baseline	66.6±6.3	66.7±4.8	1.0
6 weeks	64.8±6.1	66.6±4.9	0.5
12 weeks	65.6 ± 6.6^{b}	66.9±4.9	0.7
Change 0-12 weeks	-1.0 ± 2.7	$+0.2\pm0.9$	0.3
95% CI	-3.3 to $+1.3$	-0.7 to $+1.0$	
p value for change within	0.3	0.6	
group			
Total body water (kg)			
Baseline	48.7±4.6	48.8±3.5	1.0
6 weeks	47.4±4.5	48.7±3.6	0.6
12 weeks	48.0±4.8°	49.0±3.6	0.7
Change 0-12 weeks	-0.7±2.0	$+0.2\pm0.7$	0.7
95% CI	-2.4 to +0.9	-0.5 to +0.8	0.5
p value for change within	0.3	0.5	
1	0.5	0.5	
Volves are many SD			

Values are means±SD

^aFor difference between groups

 $^{^{}b}p$ <0.05 and ^{c}p <0.01 by paired t test for change within group (6 week level is compared with baseline and 12 week level is compared with 6 week level)

ESM 2. Adjusting the effect of dietary assignment on change^a in AUC^b Glucose₀₋₁₂₀: standardized regression coefficients in bivariate (multiple linear) regression, using change in AUC Glucose₀₋₁₂₀ as the dependent variable.

Independent variables	Standardized regression coefficients (B)	P	Cumulative adjusted R ²
Dietary assignment ^c	0.60	0.001	0.29
Weight change	-0.13	0.4	
Dietary assignment	0.68	0.0008	0.33
Waist change	-0.28	0.13	
Dietary assignment	0.57	0.002	0.27
Change in AUC Insulin ₀₋₁₂₀	0.02	0.9	
Dietary assignment	0.59	0.0008	0.34
Change in fasting plasma insulin	0.25	0.12	
Dietary assignment	0.59	0.002	0.28
Change in 120-min plasma insulin	-0.04	0.8	
Dietary assignment	0.48	0.001	0.54
Change in log HOMA	0.51	0.0006	
Dietary assignment	0.61	0.003	0.28
Age	0.08	0.7	
Dietary assignment Baseline AUC Glucose ₀₋₁₂₀	0.62 -0.45	0.0001 0.003	0.49
Dietary assignment	0.77	0.02	0.24
Cereals	-0.28	0.4	
Dietary assignment Dairy products	0.48 0.08	0.04 0.7	0.22
Dietary assignment	0.46	0.02	0.26
Pastry	0.21	0.2	
Dietary assignment	0.47	0.01	0.27
Dietary fiber	0.21	0.2	
Dietary assignment	0.48	0.02	0.23
Energy intake	0.11	0.6	
Dietary assignment	0.58	0.02	0.22
Carbohydrate intake, g	-0.07	0.8	
Dietary assignment	0.29	0.2	0.28
Glycaemic load	0.34	0.2	

^achange betweeen 0 and 12 weeks;

^bAUC, area under curve during oral glucose tolerance test;

^cPaleolithic diet = 1, Consensus diet = 2.

ESM 3. Bivariate (multiple linear) regression coefficients for change^a in AUC^b Insulin₀₋₁₂₀ (dependent variable).

Independent variables	Standardized regression coefficients (B)	P	Cumulative adjusted R ²
Weight change	0.47	0.01	0.24
Dietary assignment ^c	0.19	0.3	
Waist change	0.62	0.001	0.36
Dietary assignment	0.03	0.9	
Waist change	0.68	0.003	0.35
Carbohydrate intake, g/d	-0.08	0.7	
Waist change	0.66	0.003	0.35
Glycaemic load	-0.05	0.8	
Waist change	0.78	0.001	0.38
Cereals	-0.23	0.3	
Dietary assignment	0.28	0.2	0.01
Age	-0.01	1.0	
Dietary assignment	0.23	0.2	0.20
Baseline AUC Insulin ₀₋₁₂₀	-0.42	0.02	
Dietary assignment	0.18	0.4	0.00
Nuts	-0.14	0.5	
Dietary assignment	0.04	0.9	0.00
Cereals	0.24	0.5	
Dietary assignment	0.29	0.12	0.18
Alcohol, g/d	0.43	0.02	
Dietary assignment	0.34	0.08	0.18
Alcohol, E%	0.44	0.02	
Dietary assignment	0.10	0.6	0.05
Energy intake	0.28	0.2	
Dietary assignment	0.04	0.9	0.03
Carbohydrate intake, g/d	0.29	0.3	
Dietary assignment Glycaemic load	0.08 0.23	0.8 0.4	0.01

^achange betweeen 0 and 12 weeks;

^bAUC, area under curve during oral glucose tolerance test;

^cPaleolithic diet = 1, Consensus diet = 2.

ESM 4. Bivariate (multiple linear) regression coefficients for weight change dependent variable).

Independent variables	Standardized regression coefficients (B)	P	Cumulative adjusted R ²
Dietary assignment ^b	0.18	0.4	0.05
Age	0.02	0.9	
Dietary assignment	0.17	0.4	0.05
Baseline weight	0.02	0.9	
Dietary assignment	-0.02	0.9	0.11
Fruit	-0.43	0.05	
Dietary assignment	0.22	0.2	0.17
Meat products	0.45	0.02	
Dietary assignment	0.18	0.4	0.04
Meat, fresh	-0.04	0.9	
Dietary assignment	-0.16	0.5	0.10
Carbohydrate intake, g/d	0.51	0.06	
Dietary assignment	-0.14	0.6	0.12
Carbohydrate intake, E%	0.51	0.05	
Dietary assignment	-0.15	0.6	0.08
Glycaemic load	0.48	0.09	
Dietary assignment	-0.16	0.5	0.17
Sodium intake	0.57	0.02	
Dietary assignment	-0.27	0.5	0.05
Cereals	0.54	0.15	
Dietary assignment	0.05	0.8	0.02
Energy intake	0.29	0.2	
Cereals	0.07	0.8	0.09
Carbohydrate intake, g/d	0.34	0.2	
Cereals	0.21	0.3	0.06
Energy intake	0.20	0.4	

^achange betweeen 0 and 12 weeks;

^bPaleolithic diet = 1, Consensus diet = 2.

ESM 5. Bivariate (multiple linear) regression coefficients for change^a in waist circumference (dependent variable).

Independent variables	Standardized regression coefficients (B)	P	Cumulative adjusted R ²
Dietary assignment ^b	0.30	0.025	0.60
Weight change	0.68	0.0001	
Dietary assignment	0.49	0.027	0.13
Age	0.15	0.5	
Dietary assignment	0.44	0.027	0.13
Baseline waist	-0.07	0.7	
Dietary assignment	0.25	0.2	0.23
Fruit	-0.38	0.07	
Dietary assignment	0.32	0.14	0.16
Nuts	-0.25	0.2	
Dietary assignment	-0.40	0.2	0.41
Cereals	0.98	0.003	
Dietary assignment	0.20	0.3	0.32
Sauce	0.50	0.02	
Dietary assignment	0.21	0.3	0.25
Energy intake	0.42	0.05	
Dietary assignment	0.24	0.3	0.20
Saturated fat intake, g	0.34	0.13	
Dietary assignment	0.02	0.9	0.28
Carbohydrate intake, g/d	0.57	0.03	
Dietary assignment	0.21	0.4	0.17
Carbohydrate intake, E%	0.33	0.2	
Dietary assignment	0.12	0.6	0.21
Glycaemic load	0.44	0.10	
Dietary assignment	0.08	0.7	0.36
Sodium intake	0.59	0.007	
Cereals	0.51	0.01	0.42
Energy intake	0.26	0.16	
Cereals	0.46	0.05	0.40
Carbohydrate intake, g/d	0.25	0.3	
Cereals	0.59	0.01	0.37
Carbohydrate intake, E%	0.09	0.7	
Cereals	0.58	0.027	0.37
Glycaemic load	0.09	0.7	

Fruit	-0.48	0.004	0.46
Energy intake	0.51	0.003	
Fruit	-0.38	0.03	0.43
Carbohydrate intake, g/d	0.49	0.006	
Fruit	-0.36	0.08	0.26
Carbohydrate intake, E%	0.30	0.14	
Fruit	-0.39	0.03	0.36
Glycaemic load	0.42	0.02	
Fruit	-0.12	0.6	0.37
Cereals	0.57	0.02	

^achange betweeen 0 and 12 weeks;

^bPaleolithic diet = 1, Consensus diet = 2.

т М	1, , , , , , ,	10	1 40 50)
	I eat porridge, breakfast cereals	OI	+ # S. 32	² I eat fish
A.	a. rarely or not at all		K.	a. no more than once per month
Λ.	b. a few times per week		IX.	b. a few times per month
	c. almost every day			c. once per week
	b. low fat margarine			d. at least twice per week
	c. no fat			
J.	For lunch or dinner I eat meat, pork, minced		T.	As a rule I cut away visible fat from my meat
	meat or sausage (meat from wild animals and			when I eat
	chicken not included)			a. yes
	a. almost every day			b. no
	b. a few times per week			
	c. once per week			
	d. less often or never			
	margarine			
	d. no fat			
C.	On my bread I usually use		M.	I eat chips or cheese doodles
	a. fat cheese (>28%), sausage, liver paste			a. almost every day
	b. hamburger meat, mackerel, low-fat cheese,			b. a few times per week
	soft cheese, whey-cheese			c. once per week
	c. alternative a about as often as alternative b			d. less often or never
	d. marmalade, fruits, vegetables			d. less often of fiever
	e. plain bread with nothing except for butter or margarine			
	of margarine			
D	If I get broad with aboug I wought take		N.	I eat chocolate bar or chocolate creams
D.	If I eat bread with cheese I usually take		IN.	
	a. at least three slices of cheese (comparable			a. almost every day
	with three thick portion packed slices of			b. a few times per week
	cheese)			c. once per week
	b. two slices of cheese			d. less often or never
	c. one slice of cheese			
E.	Of bread I eat every day		O.	I eat buns, Danish pastry, cakes, tarts or ice-
	a. no more than two slices			cream
	b. 2-5 slices			a. almost every day
	c. 6-9 slices			b. a few times per week
	d. 10 slices or more			c. once per week
				d. less often or never
F.	Most often I eat		P.	I eat vegetables (tomatoes, cucumbers or green
	a. white bread			salad not included)
	b. rye meal bread			a. a few times per week or less often
	c. coarse rye-bread			b. once per day
	d. whole-meal bread, crisp bread			c. more often than once per day
G.	As far as milk and yoghurt are concerned,		Q.	I eat potatoes or other roots
٠.	I drink/eat every day		ζ.	a. a few times per week or less often
	a. at least one litre			b. once per day
	b. 0,3 - 1,0 litre			c. more often than once per day
	c. no more than 0,3 litre			c. more often man once per day
	d. almost no milk at all			
	u. aiiiiost iio iiiik at ali			
ŢŢ	As fan as will and no shout and a		D	Last aline on fried notato
H.	As far as milk and yoghurt are concerned,		R.	I eat chips or fried potatoes
	I drink/eat every day			a. a few times per week
	a. standard milk (3% of fat)			b. once per week
	b. milk with 1,5% fat			c. some times per month
	c. low fat milk (no>0,5% fat)			d. less often
I.	I eat porridge, breakfast cereals		S.	I eat fish
	a. rarely or not at all			a. no more than once per month