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A multicentre weight loss study using a low-calorie diet over 8 weeks: regional differences in efficacy across eight European cities

Angeliki Papadaki^a, Manolis Linardakis^a, Maria Plada^a, Thomas M. Larsen^b, Marleen A. van Baak^c, Anna Karin Lindroos^d, Andreas F.H. Pfeiffer^e, J Alfredo Martinez^f, Teodora Handjieva-Darlenska^g, Marie Kunešová^h, Claus Holstⁱ, Wim H.M. Saris^e, Arne Astrup^b, Anthony Kafatos^a on behalf of DiOGenes^j

^a Department of Social Medicine, Preventive Medicine & Nutrition Clinic, University of Crete, Heraklion, Crete, Greece

^b Department of Human Nutrition, Faculty of Life Sciences, University of Copenhagen, Denmark

^c Department of Human Biology, Nutrition and Toxicology Research Institute Maastricht, University of Maastricht, The Netherlands

^d MRC Human Nutrition Research, Elsie Widdowson Laboratory, Cambridge, United Kingdom

^e Department of Clinical Nutrition, German Institute of Human Nutrition Potsdam-Rehbruecke, Nuthetal, Germany and Charité Universitätsmedizin Berlin, Department of Endocrinology, Diabetes and Nutrition, Berlin, Germany

^f Department of Physiology and Nutrition, University of Navarra, Pamplona, Spain

^g Department of Human Nutrition, Dietetics and Metabolic Diseases, National Multiprofile Transport Hospital, Sofia, Bulgaria

^h Obesity Management Centre, Institute of Endocrinology, Prague, Czech Republic

ⁱ Institute of Preventive Medicine, Copenhagen University Hospitals, Copenhagen, Denmark

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Abbreviations:

DiOGenes Diet, Obesity and Genes

BMI body mass index

LCD low calorie diet

WC waist circumference

HC hip circumference

DXA dual-energy X-ray absorption

BIA bioelectric impedance analysis

FFM fat-free mass

FM fat mass

SD standard deviation

Summary

PRINCIPLES: The efficacy of low-calorie diets (LCDs) has not been investigated in large-scale studies or among people from different regions, who are perhaps unaccustomed to such methods of losing weight. The aim of the

present study was to investigate changes in obesity measures among overweight/obese adults from eight European cities (from Northern, Central and Southern Europe) during the 8-week LCD phase of the DiOGenes study (2006–2007), a family-based, randomised, controlled dietary intervention.

METHODS: 938 overweight/obese adults completed baseline examinations and underwent an 8-week LCD, providing 3.3–4.2 MJ/day to replace all meals. Anthropometric measurements and body composition were assessed at baseline and post-LCD.

RESULTS: 773 (82.4%) adults (mean age, 43.1 y) completed the LCD successfully. The highest drop-out rate was observed in Southern (24.9%) and the lowest in Northern (13.3%) European cities. Overall, the LCD induced favourable changes in all outcomes, including an approximate 11.0% reduction in body weight and body fat percentage. Changes in outcomes differed significantly between re-

gions, with North- and Central-European cities generally achieving higher percentage reductions in most anthropometric measurements assessed. Nonetheless, participants in Southern Europe reduced their body fat percentage significantly more than participants in Northern Europe (-11.8 vs. -9.5% , $P = 0.017$).

CONCLUSIONS: The LCD significantly improved anthropometric and body composition measurements in all cities participating in DiOGenes.

TRIAL REGISTRATION: ClinicalTrials.gov NCT00390637.

Key words: *DiOGenes; obesity; low-calorie diets; LCD; meal replacements; weight loss*

Introduction

Obesity has, in recent years, evolved into a worldwide epidemic [1]. Despite the investigation of meal patterns with varying macronutrient composition, it seems that the most effective diet leading to successful weight loss and maintenance has yet to be established [2]. Low-calorie diets (LCDs), in the form of energy-controlled, micronutrient-fortified liquid meals consumed as meal replacements, have often been prescribed for obese subjects or when rapid weight loss is a medical necessity [3]. Although not intended to act as a substitute for lifestyle modifications, LCDs appear popular, as they generally result in greater weight loss [4] and long-term weight maintenance [5, 6], compared to conventional diets. The success of LCDs is attributed to the production of greater satiety than alternative foods of equal or greater energy content [7]. LCDs, especially when combined with behavioural therapy and active follow-up with nutritional education and physical activity, are likely to lead to a larger long-term weight maintenance [5, 8–12], probably the result of the greater initial weight loss induced by such diets [9].

The efficacy of LCDs has not been investigated in large-scale studies or among people from different regions. DiOGenes (Diet, Obesity and Genes) is an integrated, family-based, randomised controlled dietary intervention study performed in eight European cities, examining dietary means of preventing weight regain after an initial 8-week run-in LCD weight-loss period [13–16]. To date, the majority of published research on LCDs has been undertaken in North America and North or Central European countries [17], including countries participating in DiOGenes (i.e., the Netherlands [18, 19], Denmark [20, 21], UK [22, 23], Germany [24], the Czech Republic [25, 26] and Spain [27, 28]. The aim of the current paper was to investigate the degree of weight loss, as well as changes in anthropometric and body composition measurements, among overweight/obese adults from eight European cities (from Northern, Central and Southern Europe) during the 8-week LCD phase of the DiOGenes study.

Materials and methods

Participants

Volunteer families were recruited from eight cities across eight European countries: Maastricht (the Netherlands),

Copenhagen (Denmark), Cambridge (United Kingdom), Heraklion (Greece), Potsdam (Germany), Pamplona (Spain), Sofia (Bulgaria) and Prague (Czech Republic). Recruitment commenced in March 2006 and was completed in April 2007. Families attended a screening examination to determine eligibility. Participating adults had to be healthy, overweight/ obese ($27 \leq$ body mass index (BMI, kg/m^2) < 45) and aged < 65 y. A detailed description of recruitment strategies, adult exclusion criteria and screening examinations is provided elsewhere [14]. Informed consent was obtained from all participants and the study was approved by the local Medical Ethical Committees in the respective research centers, in accordance with the Helsinki Declaration.

Study design and procedures

Eligible adults underwent a clinical examination (representing ‘baseline’) following their screening, when anthropometric measures and body composition were assessed. Subsequently participants entered an 8-week LCD period, with the aim of obtaining a minimum weight loss of 8%. A large weight loss in a short time period was chosen, to motivate participants to adhere to the main phase of the study [14] and to select the participants who were likely to show compliance during the randomised phase of the DiOGenes intervention [14]. Participants met the research dietitians approximately every two weeks (a total of approximately 6 visits) during the LCD period for weighing, compliance assessment and dietary instructions. Participants who reported difficulties with adhering to or non-compliance to the LCD were excluded from the study. At the end of the 8-week LCD period, those participants who achieved the target weight loss underwent a second clinical examination (representing ‘post-LCD’) and were subsequently randomised to one of five energy *ad libitum* diets for 6–12 months [13, 14, 29]. The present paper reports on these participants. In exceptional circumstances, e.g. when participants were unavailable to attend the post-LCD examination at exactly 8 weeks, less or more time on the LCD was allowed, provided the diet’s duration was between 7–9 weeks.

LCD intervention

The energy-restricted LCD (MODIFAST[®], Nutrition et Santé, France) consisted of 3.3 MJ/d (800 kcal/d), with a macronutrient composition of 15–20% of total energy from fat, 35–40% from protein and 45–50% from carbohydrates. The MODIFAST[®] products were available in 55 g sachets and included a range of products in a variety of flavours, namely powder drinks, crèmes and soups. Participants were required to consume a total of four sachets daily, eaten at intervals distributed across the day to replace breakfast, lunch, dinner and one snack-meal. This provided them with a daily intake of 54 g protein, ~5 g essential fatty acids and the daily requirement for vitamins and minerals. In exceptional cases, when lack of satiety was reported, a fifth sachet could be consumed.

Participants were free to add spices, herbs and low-calorie/calorie-free flavourings (i.e., calorie-free juice, coffee or instant coffee) to increase variety and taste of the products. In addition to the four sachets, participants were also per-

mitted without limitation to: drink coffee and tea (if necessary adding a little skimmed milk); drink sufficient quantities of water; drink calorie-free soft drinks; and chew sugar-free chewing gum/pastilles. Further, it was permitted to eat 200 g tomatoes, 125 g cucumber and 50 g lettuce on a daily basis. Thus, participants were provided with 3.3–4.2 MJ/d (800–1000 kcal/d).

Anthropometric measurements and body composition

Standard operation procedures were produced for all investigations undertaken to ensure standardisation across the cities and the same measurement devices and methods were used in each research centre on every occasion a measurement was provided [14]. Subjects had been fasting for twelve hours before the clinical examinations in the early morning and they were measured in their underwear, with an empty bladder.

Weight was measured on calibrated digital scales to the nearest 0.1 kg on all examinations and dietary counselling sessions, and height was measured at baseline, to the nearest mm with a wall-mounted stadiometer. BMI was calculated as weight divided by height squared (kg/m^2). Waist circumference (WC), measured midway between the lower rib and iliac crest) and hip circumference (HC), measured at the widest point between the hips and buttocks) were measured twice, to the nearest 0.5 cm, with a tape in a vertical plane and with the subject standing and gently breathing out. Waist-to-hip ratio was then calculated. Sagittal abdominal diameter was measured using an abdominal caliper to the nearest mm, at the highest point of the abdomen during expiration, with subjects supine [14]. Fat mass (kg), body fat percentage and fat-free mass (kg, defined as the sum of lean body mass and bone mineral content) were determined by dual-energy X-ray absorption (DXA) or by bioelectric impedance analysis (BIA). Adults who participated in the determination of body composition were fasting (no intake of foods or liquids for at least 4 hours prior to the examinations, except for a water intake of 350–500 mL).

Statistical analyses

All analyses were performed using the Statistical Package for the Social Sciences (IBM SPSS for Windows, release 19, 2010, SPSS, Chicago, Illinois). Drop-out from the study during the 8-week LCD phase (non-completers) was determined by assessing the number of participants who entered the LCD phase and the number who successfully completed the LCD and were randomised to the main DiOGenes intervention (completers). For both LCD completers and non-completers the distribution of participants according to gender was compared between the different cities using the Pearson's chi-square test. Differences in gender distribution and age between completers and non-completers were assessed using Pearson's chi-square test and analysis of variance respectively. Age of participants and LCD duration (in weeks) were compared between the cities using the Kruskal Wallis rank test. Comparisons of overall changes in anthropometric measurements and body composition during the LCD period between Northern (Maastricht, Copenhagen, Cambridge), Central (Prague, Potsdam) and Southern (Heraklion, Pamplona, Sofia)

European regions were performed using analysis of covariance, with age, gender and LCD duration as covariates and pair-wise comparisons, with age, gender, baseline body fat percentage and LCD duration as covariates.

Results

A total of 1 209 adults were registered to attend screening and 938 (77.6%) attended the baseline examination and entered the LCD period. Of these, 773 (82.4%) adults (273 males, 500 females, mean age, 43.1 y) successfully completed the LCD and were included in the present analyses. Overall, 165 adults (17.6% of those who entered the LCD) dropped out or were excluded during the LCD. The distribution of participants across different cities and regions is shown in figure 1. The highest drop-out was observed in Heraklion, Pamplona and Cambridge, and the lowest in Copenhagen. The combined analyses of cities according to region showed that mean drop-out was higher in Southern (24.9%) than in Central (16.2%) or Northern European (13.3%) cities (fig. 1). Among non-completers of the LCD

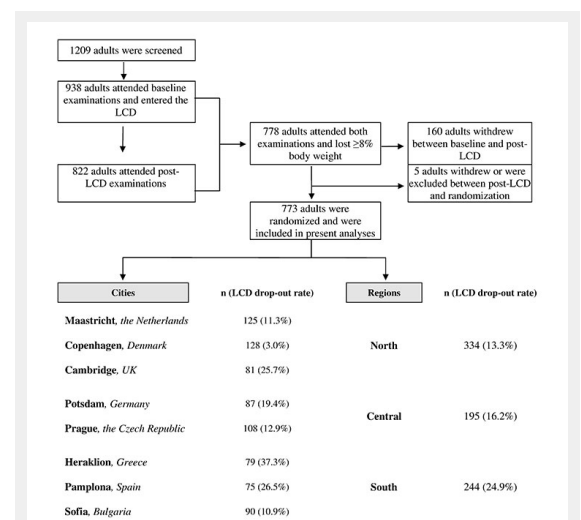


Figure 1

Flow diagram showing progress of the DiOGenes participants from screening to post-LCD, 2006–2007.

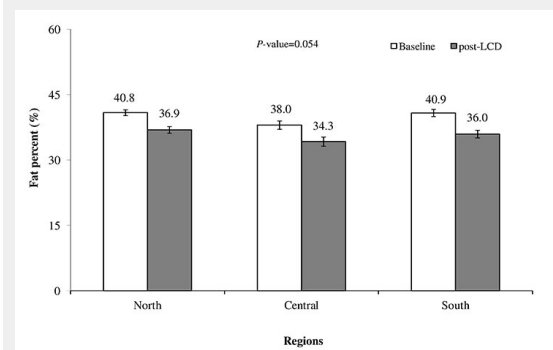


Figure 2

Body fat percentage over the LCD period in the DiOGenes participating regions, 2006–2007. LCD = low-calorie diet; bars (I) indicate 95% confidence intervals; comparisons in body fat percentage changes between regions were performed using analysis of covariance, with age, gender, baseline body fat percentage and LCD duration (weeks) as covariates.

period, there was no difference between the cities in the distribution of genders ($P = 0.869$, data not shown). Non-completers were younger (41.2 vs. 43.1y, $P < 0.001$) than LCD completers.

Among cities, gender distribution differed significantly ($P = 0.033$). Participants in Heraklion were significantly younger ($P < 0.001$) and followed the LCD for a longer period ($P < 0.001$), compared to other cities (table 1).

The mean and percental changes in anthropometric and body composition measurements during the LCD period in the three different European regions and the total sample are shown in table 2. Overall, the LCD induced favourable changes in all measured outcomes and an 11.1% (11.1 kg) and 11.0% reduction in body weight and fat percentage, respectively. Changes in all outcomes, except body weight and BMI, differed significantly between the regions. In general, participants in the Northern and Central European regions achieved higher reductions in most anthropometric measurements assessed, compared to the Southern European region (table 2). However, participants from the Southern European cities reduced their fat mass significantly more (-21.5% , $P < 0.01$) and their fat-free mass significantly less (-3.6% , $P < 0.01$), compared to participants from the Northern (-19.3% and -5.7% , respectively) and Central (-19.6% and -5.7% , respectively) European regions. Body fat percentage changes during the LCD period tended to differ between the groups of cities ($P = 0.054$) (fig. 2). Pair-wise comparisons revealed that participants in the Southern region reduced their body fat significantly more than participants in the Northern European region ($P = 0.017$) (fig. 2).

Discussion

LCDs have frequently been scientifically evaluated in the last decades, as they result in rapid weight loss, which renders this method popular among obese subjects who consider it acceptable [6, 30] and are likely to help achieve long-term weight loss maintenance [6, 9, 24], compared to conventional diets. In the present study, overweight/obese adult volunteers from 8 European cities participating in the

DiOGenes study followed an LCD for a mean of 8.2 weeks and lost an average of 11.1% body weight. Such a reduction in body weight has been associated with significant improvements in obesity-related disorders and complications [11, 24]. In addition, the LCD produced favorable changes in a variety of anthropometric and body composition measurements across different European regions, indicating its efficacy in a large-scale study where a large weight loss in a short time period was required.

Participants in the present study lost an average of 2.8 kg (4.7%) of fat-free mass, corresponding to 25.2% of the weight lost, which is comparable to earlier research and considered acceptable in weight loss efforts [18, 31–34]. Although earlier studies have been inconsistent with regard to the role of initial BMI and body composition in the loss of fat-free mass and body weight respectively [18, 33], we accounted for any potential effect of baseline values by examining absolute and percentage changes in these parameters. Nevertheless, weight reduction in the present study was mainly attributed to reduction in body fat. It should be noted however, that comparisons with earlier studies are hindered, due to differences in diet type and energy content (LCD vs. very-low-calorie diets), LCD definition and duration and baseline characteristics of participants, such as initial BMI. DiOGenes is, to the best of our knowledge, the first study to provide an opportunity for a comparison of LCD efficacy between different European regions. All eight research centres successfully implemented the LCD, although changes in the majority of outcome measures differed significantly between regions, despite standardisation of protocols and procedures. In general, participants in the Northern and Central European regions achieved higher percentage reductions in most anthropometric measurements assessed, compared to participants from Southern Europe. Nonetheless, participants in the Southern European region generally achieved significantly higher reductions in body fat percentage.

Drop-out in the present study was slightly higher than withdrawal rates reported in some earlier studies [35, 36], but lower compared to the only other multi-centre study we could identify, which comprised a national intervention in the USA [37], and to attrition reported in the review by Tsai & Wadden [2006]. In DiOGenes, most subjects who did not complete the LCD period withdrew during the first half of the LCD period. No severe adverse effects were reported and participants were excluded if they reported non-

Table 1: Distribution of gender, participant age and LCD duration in the DiOGenes participating cities ($n = 773$), 2006–2007.

	n	Gender		Pa	Age (y)	Pb	LCD duration (weeks)		Pb
		Males	Females				Mean (SD)	Pb	
Maastricht	125	39.2	60.8	0.033	43.8 (5.9)	<0.001	8.2 (1.4)	<0.001	
Copenhagen	128	40.6	59.4		43.5 (5.4)		8.3 (1.4)		
Cambridge	81	25.9	74.1		43.9 (7.2)		8.0 (1.2)		
Heraklion	79	36.7	63.3		40.1 (5.8)		9.0 (1.2)		
Potsdam	87	39.1	60.9		42.6 (6.1)		8.7 (1.1)		
Pamplona	75	45.3	54.7		44.3 (5.6)		7.8 (1.5)		
Sofia	90	28.9	71.1		44.0 (7.5)		7.9 (0.9)		
Prague	108	25.9	74.1		42.3 (6.6)		7.9 (1.3)		
Total	773	35.3	64.7		43.1 (6.3)		8.2 (1.3)		

LCD = low-calorie diet; SD = standard deviation; a the distribution of gender at the different centres is compared using Pearson chi-square test; b age and LCD duration at the different centres are compared using Kruskal Wallis rank test.

compliance with the treatment or voluntarily withdrew. It is usual in weight loss studies that patients who do not achieve the intended outcomes are more likely to drop out [11] and it has recently been suggested that early weight loss is a predictor of final weight loss during an 8-week LCD [38]. In the present study, many of the participants who withdrew reported that, although initially driven to participate due to the rapid weight loss to be induced by the LCD, they realized that this treatment method did not agree with their lifestyle, such as outdoor eating obligations.

The marked differences in drop-out between different research centres, e.g. 3% in Copenhagen and ~37% in Heraklion, and regions, require further investigation. However, participants in Heraklion who completed the LCD achieved higher percentage changes in anthropometric measurements, compared to the other Southern European cities, and the biggest changes in body composition measurements among all participating cities (data not shown). One might argue that these changes are the result of the longer LCD period in this research centre, thus allowing more time to achieve favourable results. However, it has been suggested that increases in LCD duration of more than 8 weeks result in adherence difficulties and reduced compliance, due to lack of dietary variation [34]. In addition, LCD duration was taken into account in the present analyses. As participants in Heraklion also had one of the highest percentage body weight reductions among all participating cities (data not shown), it may be that the weight loss

achieved encouraged higher degrees of compliance among participants not withdrawing from the LCD period early on in the procedure. Issues of palatability/familiarity with meal replacement products should also be considered when attempting to clarify the present drop-out rates. For example, soups provided in the current study were asparagus and potato and leek soups, which are not part of the usual Greek diet and might have discouraged some participants from Heraklion from continuing on the LCD phase. Therefore, future studies utilizing meal replacement products to promote weight loss should ideally provide products with flavours similar to local cuisines. Nevertheless, future studies should include objective measures of palatability/acceptability of LCDs, instead of relying on self-reports of non-compliance with the treatment, in order to better understand differences in compliance, drop-out and effects among different regions.

In the present study, no severe adverse effects of the LCD were reported. Nonetheless, it should be noted that such diets are not intended to substitute long-term behavioural modification for weight loss/control. Indeed, the superiority of LCDs to conventional diets has been questioned in the past [9, 11], whereas the combined treatment of LCDs and behavioural therapy appears to be more effective, compared to LCD alone, especially with regard to long-term weight maintenance [5, 11].

To the best of our knowledge, this is the first study to investigate the effect of LCDs on anthropometric measure-

Table 2: Mean (SD) and percental changes in anthropometric and body composition measures over the LCD period in the DiOGenes participating regions (n 773), 2006–2007.

Measurement		Regions			Total
		North	Central	South	
Weight (kg) (n = 773)	Baseline	99.1 (16.0)	100.0 (16.0)	101.5 (16.0)	100.0 (16.1)
	Mean change	-11.3 (3.3)	-11.1 (3.3)	-10.8 (3.3)	-11.1 (3.3)
	% change	-11.4	-11.1	-10.6	-11.1
BMI (kg/m ²) (n = 771)	Baseline	33.7 (4.9)	33.7 (4.9)	36.1 (4.8)	34.6 (5.0)
	Mean change	-3.8 (1.1)	-3.7 (1.1)	-3.6 (1.1)	-3.8 (1.1)
	% change	-11.3	-11.0	-10.0	-11.0
WC (cm) b (n = 756)	Baseline	107.4 (11.9)	105.7 (11.9)	109.8 (11.9)	107.9 (11.9)
	Mean change	-10.2 (4.7)	-10.2 (4.7)	-8.9 (4.7)	-9.7 (4.8)
	% change	-9.5	-9.6	-8.1	-9.0
HC (cm) b (n = 756)	Baseline	114.8 (10.3)	116.6 (10.3)	119.3 (10.3)	117.1 (10.3)
	Mean change	-6.9 (4.1)	-7.9 (4.0)	-7.1 (4.0)	-7.2 (3.9)
	% change	-6.0	-6.8	-6.0	-6.1
Waist-to-hip ratio b (n = 756)	Baseline	0.937 (0.073)	0.908 (0.069)	0.921 (0.061)	0.923 (0.055)
	Mean change	-0.036 (0.055)	-0.029 (0.042)	-0.022 (0.045)	-0.029 (0.055)
	% change	-3.8	-3.2	-2.4	-3.1
Sagittal height (cm) a (n = 727)	Baseline	24.4 (3.6)	26.2 (3.5)	25.8 (3.5)	25.4 (3.4)
	Mean change	-3.6 (1.7)	-3.4 (1.7)	-2.7 (1.7)	-3.2 (1.7)
	% change	-14.8	-13.0	-10.5	-12.6
FM (kg) b (n = 563)	Baseline	40.9 (10.9)	37.7 (10.9)	42.4 (10.9)	40.3 (10.9)
	Mean change	-7.9 (4.4)	-7.4 (4.4)	-9.1 (4.4)	-8.3 (4.4)
	% change	-19.3	-19.6	-21.5	-20.6
FFM (kg) b (n = 563)	Baseline	59.2 (8.4)	61.6 (8.4)	60.9 (8.4)	60.1 (8.1)
	Mean change	-3.4 (4.0)	-3.5 (4.0)	-2.2 (4.0)	-2.8 (4.0)
	% change	-5.7	-5.7	-3.6	-4.7
Fat percent (%) b (n = 563)	Baseline	40.8 (5.6)	38.0 (5.6)	40.9 (5.6)	40.0 (5.5)
	Mean change	-3.9 (3.9)	-3.7 (3.9)	-4.8 (3.9)	-4.4 (3.9)
	% change	-9.5	-9.7	-11.8	-11.0

LCD = low-calorie diet; SD = standard deviation; BMI = body mass index; WC = waist circumference; HC = hip circumference; FM = fat mass; FFM = fat-free mass. Comparisons of overall mean changes between regions were performed using analysis of covariance, with age, gender and LCD duration (weeks) as covariates. a *P* < 0.001, b *P* < 0.01.

ments and body composition in such a large and diverse sample of overweight/obese individuals. The current findings might therefore prove helpful to scientists intending to use this treatment method at a national level in the future. However, it should be noted that our sample was not representative of the general population in either the respective cities or countries, and thus our findings cannot be generalised. A limitation of the current study is that physical activity levels during the LCD were not assessed. Thus, we cannot rule out that the observed favourable changes in obesity measures were not the combined result of the LCD and increased physical activity. However, and also for reasons of standardisation assurance across the research centres, subjects were particularly requested not to change their usual lifestyle habits, including physical activity, during the LCD period. Nevertheless, it is noteworthy that studies specifically designed to examine the effect of physical exercise revealed no additional weight loss when physical exercise was added to the LCD treatment [39]. In addition, our database did not provide information on more specific acceptability measures, such as satiety produced by the diet or the taste of the LCD products. Such measurements should, if possible, be assessed during treatment to provide an additional estimate of product acceptability and explain potential drop-out differences. Furthermore, the LCD in this study was not compared to any other treatment, so recommendations regarding its efficacy might seem inappropriate. Nevertheless, this study provides additional evidence on the effect of LCDs in a free-living situation and shows this treatment's potential for use in different regions.

In conclusion, the LCD significantly improved anthropometric and body composition measurements in participants in all cities participating in the DiOGenes study. Future studies should examine objective acceptability measures of this treatment in order to further explain regional differences in the effect of this diet on obesity.

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Correspondence: Angeliki Papadaki, PhD, Centre for Exercise, Nutrition and Health Sciences, School for Policy Studies, University of Bristol, 8 Priory Road, UK-Bristol,

BS8 1TZ, United Kingdom,
[angeliki.papadaki\[at\]bristol.ac.uk](mailto:angeliki.papadaki[at]bristol.ac.uk)

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Figures (large format)

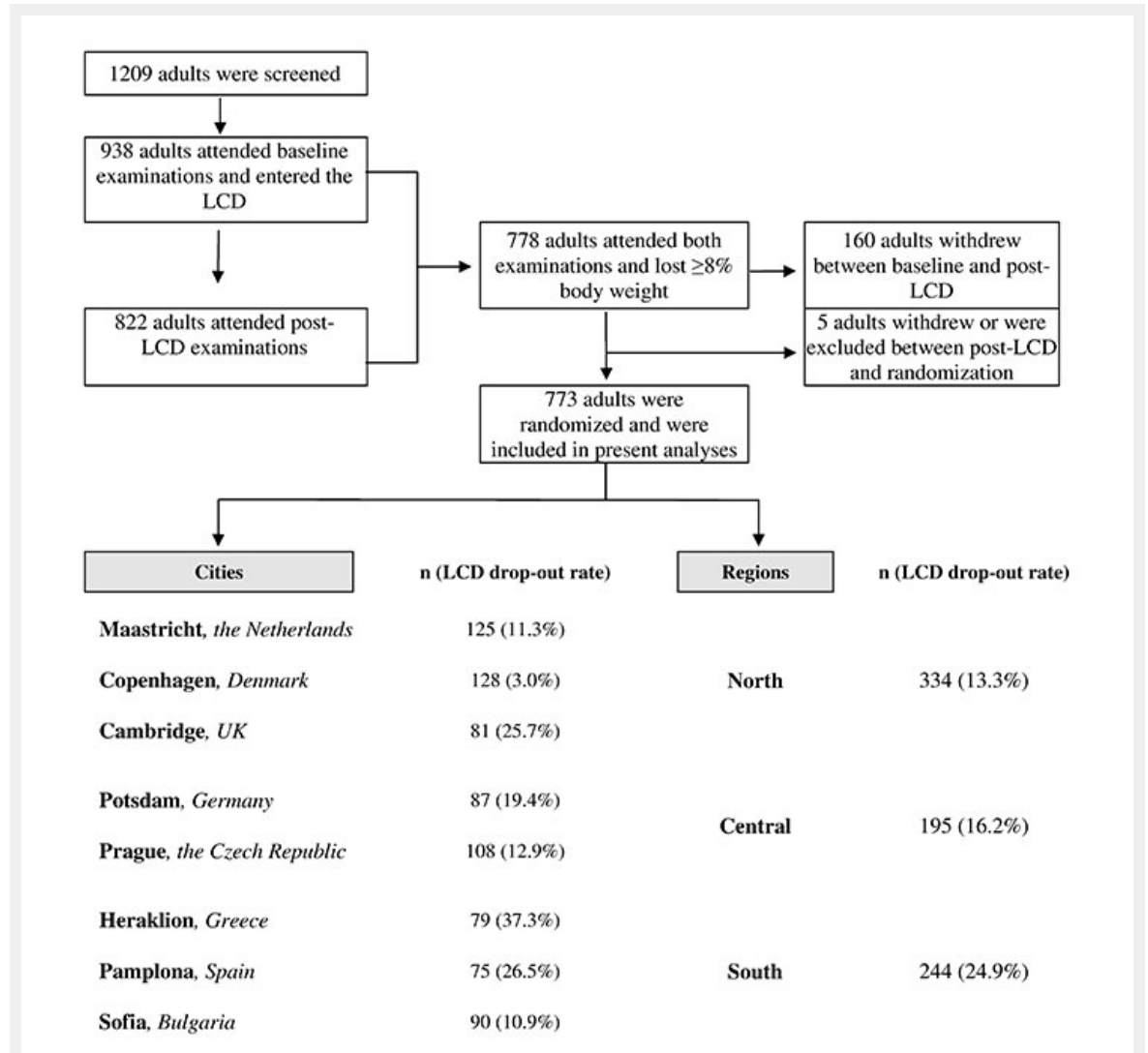


Figure 1
Flow diagram showing progress of the DiOGenes participants from screening to post-LCD, 2006–2007.

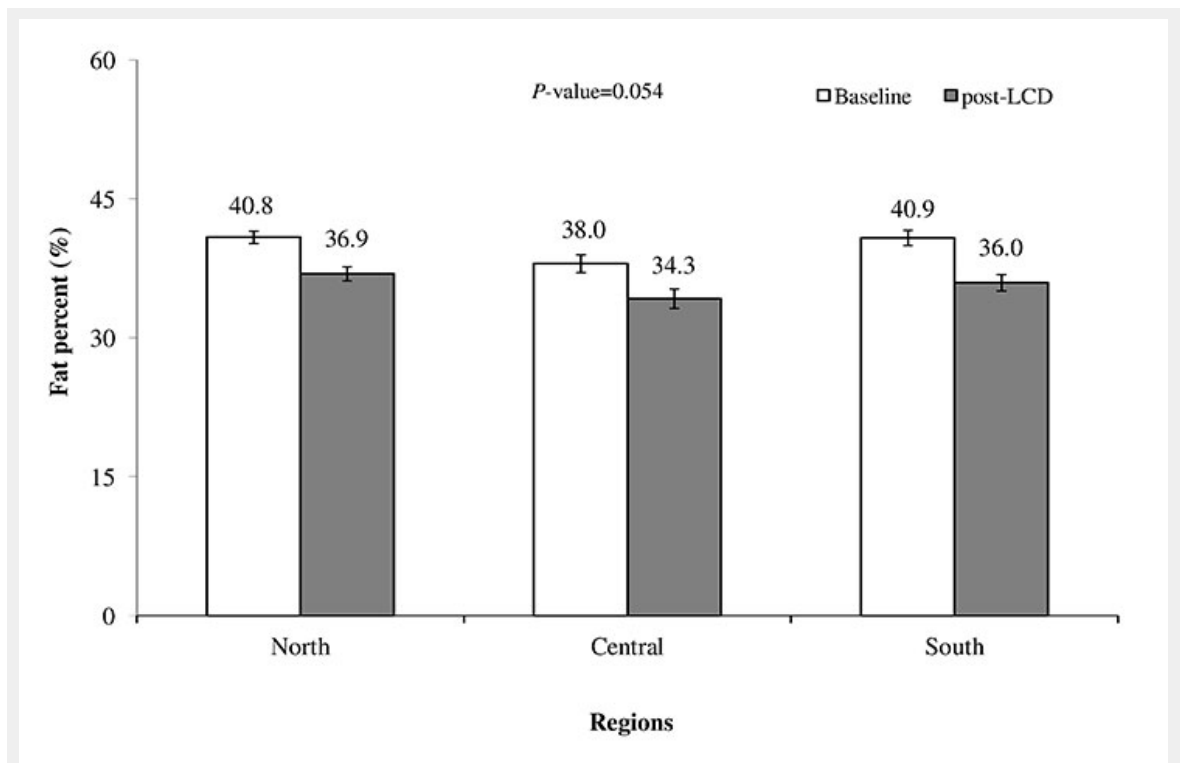


Figure 2

Body fat percentage over the LCD period in the DiOGenes participating regions, 2006–2007. LCD = low-calorie diet; bars (I) indicate 95% confidence intervals; comparisons in body fat percentage changes between regions were performed using analysis of covariance, with age, gender, baseline body fat percentage and LCD duration (weeks) as covariates.