Associations between olfactory pathway gene methylation marks, obesity features and dietary intakes

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Abstract

Background: Olfaction is an important sense influencing food preferences, appetite, and eating behaviors. This hypothesis-driven study aimed to assess associations between olfactory pathway gene methylation signatures, obesity features, and dietary intakes.

Methods: A nutriepigenomic analysis was conducted in 474 adults from the Methyl Epigenome Network Association (MENA) project. Anthropometric measurements, clinical data, and serum metabolic profiles of the study population were obtained from structured databases of the MENA cohorts. Habitual dietary intake was assessed using a validated semiquantitative food frequency questionnaire. DNA methylation was measured in circulating white blood cells by microarray (Infinium Human Methylation 450 K BeadChips). FDR values (p < 0.0001) were used to select those CpGs that showed the best correlation with body mass index (BMI) and waist circumference (WC). Pathway analyses involving the characterization of genes involved in the olfactory transduction system were performed using KEGG and pathDIP reference databases.

Results: Overall, 15 CpG sites at olfactory pathway genes were associated with BMI (p < 0.0001) and WC (p < 0.0001) after adjustments for potential confounding factors. Together, methylation levels at the 15 CpG sites accounted for 22% and 20% of the variability in BMI and WC (r² = 0.219, p < 0.001, and r² = 0.204, p < 0.001, respectively). These genes encompassed olfactory receptors (OR4D2, OR51A7, OR2T34, and OR2Y1) and several downstream signaling molecules (SLC8A1, ANO2, PDE2A, CALML3, GNG7, CALML6, PRKG1, and CAMK2D), which significantly regulated odor detection and signal transduction processes within the complete olfactory cascade, as revealed by pathway enrichment analyses (p = 1.94 × 10⁻10). Moreover, OR4D2 and OR2Y1 gene methylation patterns strongly correlated with daily intakes of total energy (p < 0.0001), carbohydrates (p < 0.0001), protein (p < 0.0001), and fat (p < 0.0001).

Conclusions: The results of this study suggest novel relationships between olfactory pathway gene methylation signatures, obesity indices, and dietary intakes.

Keywords: Olfactory system, Smell, Epigenetics, OR2Y1, OR4D2, Diet

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Background

Obesity epidemic represents one of the most important health challenges worldwide [1]. It has been estimated that about 1.9 billion adults present overweight or obesity based on body mass index (BMI), with an overall prevalence of 39% [2]. Epidemiological studies have identified high BMI as a risk factor for an expanding set of chronic diseases including cardiovascular disease, diabetes mellitus, non-alcoholic fatty liver disease, and many types of cancer, with relevant negative economic and social impacts [3]. At a global level, excessive weight (BMI ≥ 25) accounted for 4 million deaths (7.2% of all-cause deaths) and 120 million deaths and disability-adjusted life years (4.9% of all-cause DALYs) among adults in 2015 [4].

In general, the overconsumption of energy-dense foods, coupled with the adoption of a sedentary lifestyle, is the main environmental factors contributing to the development of obesity and associated clinical manifestations [5]. Moreover, genetic and epigenetic signatures play a role in determining individual susceptibility to fat accumulation [6, 7]. In the past, and until recently, research on obesity has focused on characterizing the biological factors that predispose people to excessive food intake and associated weight gain [8]. In this context, it has been reported that sensorial modalities, including taste and olfaction [9], may influence food preferences and eating behaviors.

In particular, olfactory ability is related to perceived food palatability [10] and may be modulated by hunger/satiety states [11, 12]. Also, exposure to food odors increases appetite for products with similar characteristics in terms of taste and energy density [13]. Indeed, a link between dietary intake and olfactory sensitivity to fat was found [14], although another study did not find changes in energy intake or food preferences in response to odor exposure in women [15]. In addition, fluctuations in the internal levels of carbohydrates, amino acids, and fats could modulate olfactory sensitivity to adjust feeding behaviors in order to maintain nutrient homeostasis [16].

Odor perception during eating depends on the interaction between olfactory and gustatory systems [17]. Besides physiological issues (i.e., age, circadian rhythmicity, endocrine secretions), sensory variability can be also driven by genetic and epigenetic marks, which in turn can impact food consumption and subsequent health outcomes [18]. Associations between taste receptors polymorphisms, dietary intakes, lipid disorders, and liver disease in Mexican subjects were reported [19–22]. Likewise, sequence variants in olfactory receptor genes appeared to contribute to the predisposition to extreme obesity [23, 24] and influenced eating behaviors and adiposity levels [25]. Additionally, DNA methylation patterns at sweet taste pathway genes were associated with BMI and carbohydrate intake in an adult population [26]. Together, these findings suggest the involvement of sensory factors, including olfaction, in appetite regulation and obesity predisposition, where epigenetics may play a crucial role. This hypothesis-driven study aimed to assess associations between olfactory pathway gene methylation signatures, obesity features, and dietary intakes.

Materials and methods

Subjects

A nutriepigenomic analysis was conducted in an adult population from the Methyl Epigenome Network Association (MENA) project (n = 474), which is constituted by previous clinical trials [27–34]. The study protocol, data confidentiality, and research procedures were in accordance with the ethical principles on human experimentation stipulated in the 2013 Declaration of Helsinki by the World Medical Association [35].

Study variables

Anthropometric measurements, clinical data, and serum metabolic profiles of the study population were obtained from structured databases of the MENA cohorts. These variables included weight, height, waist circumference (WC), systolic blood pressure (SBP), diastolic blood pressure (DBP), glucose, insulin, total cholesterol, high-density lipoprotein cholesterol (HDL-c), low-density lipoprotein cholesterol (LDL-c), and triglycerides. BMI was calculated as the ratio between weight (kg) and height (m²). Insulin resistance was estimated by the homeostatic model assessment-insulin resistance (HOMA-IR) index using the following formula: HOMA-IR = fasting insulin (mU/L) X plasma glucose (mmol/L)/22.5. Triglyceride-glucose (TyG) index was calculated as a predictor of diabetes, as described elsewhere [36].

Dietary assessment

Dietary data was available from 247 subjects of the PRE-DIMED, RESMENA, and OBEKIT cohorts, which presented similar characteristics regarding the whole sample. A validated semiquantitative food frequency questionnaire [37] was used to assess the habitual consumption of 137 food items during the previous year according to four frequency categories (daily, weekly, monthly, or never). The obtained food portions and serving sizes were further converted to daily energy (kcal) and macronutrient intakes (g) using the Spanish food composition tables [38].
DNA methylation analyses
Venous blood samples were drawn by venipuncture after a 12-h overnight fast. White blood cells (WBC) were separated from whole blood by centrifugation at 3500 rpm, at 4°C for 15 min, and immediately frozen at −80°C in buffy coat until assay, as described elsewhere [39]. Genomic DNA was isolated from WBC with the MasterPure kit (Epicentre Biotechnologies, Madison, WI, USA). Purified DNA was quantified by the PicoGreen® dsDNA Quantitation Reagent (Invitrogen, Carlsbad, CA, USA). High-quality DNA samples were modified with sodium bisulfite by using the EZ-96 DNA Methylation kit (Zymo Research Corporation, Irvine, CA, USA) according to the manufacturer’s protocol. Bisulfite-treated DNA samples were hybridized to Infinium Human Methylation 450 K BeadChips (Illumina, San Diego, CA, USA) and scanned using the Illumina HiScanSQ system. Image intensities were obtained with the GenomeStudio Methylation Software Module, v1.9 (Illumina, San Diego, CA, USA). DNA methylation data pre-processing has been recently described elsewhere [40]. Briefly, CpG methylation levels were expressed as β values ranging from 0 (unmethylated) to 1 (methylated) [41], which were corrected for type I and type II bias applying the peak-based correction. Data were normalized in R using a categorical Subset Quantile Normalization method [42]. Probes containing single-nucleotide polymorphisms, hybridizing to multiple genomic locations, or associated with X and Y chromosomes were removed from the analysis. Furthermore, DNA methylation variation explained due to different cell subtypes was corrected following the Houseman procedure [43].

Pathway analyses
Pathway mapping of genes involved in olfactory transduction (map04740) was performed using the Kyoto Encyclopedia of Genes and Genomes (KEGG) reference database (http://www.genome.jp/kegg/pathway.html). Subsequently, pathway enrichment analyses (confidence level of 99%) were further run in the Pathway Data Integration Portal (pathDIP) platform, University of Toronto, Canada (http://ophid.utoronto.ca/pathdip/).

Statistical analyses
Data normality was screened by the Kolmogorov-Smirnov test. Principal variables including BMI, WC, energy, and macronutrient intakes were normally distributed ($p > 0.05$). Continuous variables were expressed as means ± standard deviations, while categorical variables were presented as number of cases and percentages. Statistical analyses were performed in the IBM SPSS software, version 20 (IBM Inc., Armonk, NY, USA). A linear regression model (for BMI outcome) was computed in the LIMMA package for R software, which was adjusted by age, sex, study cohorts, and DNA methylation chips. Also, the Benjamini-Hochberg correction for multiple comparisons was applied. Statistically significant thresholds were based on false discovery rate (FDR) cut-offs ($p < 0.05$) and B-statistic values ($> 0$) for BMI-related analysis. Afterwards, stricter FDR ($p < 0.0001$) were used to select those CpGs that showed the best correlation with BMI. Linear regression analyses adjusted by age and sex were further performed to evaluate correlations of methylation values at olfactory transducing genes with anthropometric (BMI, WC), metabolic (glucose, insulin, lipid profile, HOMA-IR, TyG index), and dietary measurements (total energy and macronutrient intakes). Figure plots showing significant correlations were created in the GraphPad Prism® program, version 6.0C (La Jolla, CA, USA).

Results
Demographic, anthropometric, and metabolic characteristics of the whole study population are reported (Table 1). Thirty-six percent of subjects were men ($n = 171$). Eighty-two percent of the study population presented

| Variable | Average values |
|----------|----------------|
| Age (years) | 47.2 ± 14.1 |
| Men/women | 171/303 |
| Anthropometric and clinical data | |
| Weight (kg) | 81.6 ± 19.1 |
| BMI (kg/m²) | 30.1 ± 5.6 |
| WC (cm) | 95.7 ± 16.1 |
| MAP (mmHg) | 100.4 ± 16.2 |
| Metabolic profile | |
| Glucose (mmol/L) | 5.7 ± 1.7 |
| Insulin (pmol/L) | 66.7 ± 48.6 |
| HOMA-IR index | 2.44 ± 2.28 |
| Total cholesterol (mmol/L) | 5.3 ± 1.0 |
| Triglycerides (mmol/L) | 1.4 ± 0.8 |
| TyG index | 4.61 ± 0.32 |
| Dietary intake | |
| Energy (Kcal/day) | 2576 ± 759 |
| Carbohydrates (g/day) | 260.9 ± 96.8 |
| Protein (g/day) | 103.8 ± 28.3 |
| Fat (g/day) | 115.3 ± 36.4 |

Continuous variables are represented as means ± standard deviations. Men and women are number of subjects. BMI body mass index, WC waist circumference, MAP mean arterial pressure, HOMA-IR index homeostatic model assessment-insulin resistance index, TyG index triglyceride-glucose index. Dietary intake was available from 247 subjects.
excessive body weight \((n = 390)\) according to the BMI classification of the World Health Organization \((\text{BMI} \geq 25 \, \text{kg/m}^2)\). Regarding the reference values, the metabolic profile of the whole population was characterized by increased blood levels of glucose and total cholesterol.

The first screening revealed that 61 CpG sites at genes participating in the olfactory transduction pathway correlated with the BMI \((p < 0.05)\). Out of these, 35 CpGs showed best correlations with BMI based on stricter FDR values \((p < 0.0001)\). After performing linear regression tests adjusted by sex and age, 15 CpG sites still remained statistically significant. These CpG sites comprised cg19302979 \((\text{SLC8A1})\), cg02874396 \((\text{OR4D2})\), cg10610428 \((\text{ANO2})\), cg12498094 \((\text{PDE2A})\), cg17283169 \((\text{CALML3})\), cg02849894 \((\text{GNG7})\), cg15102821 \((\text{CALML6})\), cg16401207 \((\text{PRKG1})\), cg00467296 \((\text{ORS1A7})\), cg24609819 \((\text{PRKG1})\), cg13801347 \((\text{CAMK2D})\), cg15819352 \((\text{CALML6})\), cg13441213 \((\text{OR2T34})\), and cg18482656 \((\text{OR2Y1})\).

Interestingly, most of the BMI-associated CpGs were mapped to coding regions \((n = 7)\) or terminal sequences \((n = 2)\), and only six were located on gene promoters (Table 2). Together, methylation levels at these 15 CpG sites accounted for 22% of the variability in BMI \((r^2 = 0.219, p < 0.001)\). Illustrative correlations of each CpG site are reported (Fig. 1). Moreover, methylation signatures at the aforementioned 15 CpGs also correlated with WC values in a similar way (Fig. 2), explaining about 20% of variation in WC \((r^2 = 0.204, p < 0.001)\). No statistically significant relationships between methylation status at olfactory transducing genes with blood levels of metabolic markers and blood pressure measurements were found.

Pathway mapping of the BMI-associated genes within the olfactory transduction cascade is shown (Fig. 3).

### Table 2: Genomic and statistical data of CpG sites at olfactory pathway genes putatively associated with BMI

| CpG_ID | Illumina_ID | Gene name | CHR position | Genomic region | p value | FDR | B | r^2 |
|--------|-------------|-----------|--------------|----------------|---------|-----|---|-----|
| cg19302979 | Solute carrier family 8 member A1 | 1: 2,404,368,433 | Body | 4.0 × 10^{-12} | 3.2 × 10^{-09} | 15.80 | 0.12 |
| cg02874396 | Olfactory receptor family 4 subfamily D member 2 | 17: 16,524,854 | TSS1500 | 1.3 × 10^{-10} | 4.4 × 10^{-08} | 12.42 | 0.06 |
| cg10610428 | Anoctamin 2 | 12: 588,429 | Body | 1.5 × 10^{-09} | 2.7 × 10^{-07} | 10.01 | 0.07 |
| cg12498094 | Solute carrier family 8 member A1 | 1: 2,403,567,822 | Body | 1.4 × 10^{-08} | 1.4 × 10^{-08} | 7.81 | 0.05 |
| cg07736155 | Phosphodiesterase 2A | 11: 723,541,000 | TSS1500 | 1.7 × 10^{-08} | 1.6 × 10^{-08} | 7.65 | 0.07 |
| cg17283169 | Calmodulin-like 3 | 10: 55,675,24 | 3′UTR | 2.1 × 10^{-09} | 2.0 × 10^{-09} | 7.42 | 0.08 |
| cg02849894 | G protein subunit gamma 7 | 19: 260,897,13 | 5′UTR | 4.5 × 10^{-08} | 3.4 × 10^{-06} | 6.68 | 0.06 |
| cg15102821 | Calmodulin-like 6 | 1: 184,480,13 | TSS1500 | 1.4 × 10^{-07} | 7.6 × 10^{-07} | 5.61 | 0.05 |
| cg16401207 | Protein kinase, cGMP-dependent, type I | 10: 531,821,11 | Body | 1.5 × 10^{-07} | 8.0 × 10^{-06} | 5.54 | 0.05 |
| cg00467296 | Olfactory receptor family 51 subfamily A member 7 | 11: 492,876,00 | 1stExon | 1.9 × 10^{-07} | 9.7 × 10^{-06} | 5.28 | 0.04 |
| cg24609819 | Protein kinase, cGMP-dependent, type I | 10: 528,430,77 | Body | 3.8 × 10^{-07} | 1.6 × 10^{-05} | 4.62 | 0.05 |
| cg13801347 | Calcium/calmodulin dependent protein kinase II delta | 4: 11,446,005 | Body | 1.7 × 10^{-06} | 4.7 × 10^{-05} | 3.14 | 0.05 |
| cg15819352 | Calmodulin-like 6 | 1: 184,594,00 | TSS1500 | 2.1 × 10^{-09} | 5.4 × 10^{-05} | 2.96 | 0.05 |
| cg13441213 | Olfactory receptor family 2 subfamily T member 34 | 1: 248,738,75 | TSS1500 | 3.0 × 10^{-06} | 6.9 × 10^{-05} | 2.62 | 0.05 |
| cg18482656 | Olfactory receptor family 2 subfamily Y member 1 | 5: 180,167,96 | TSS1500 | 3.6 × 10^{-06} | 7.9 × 10^{-05} | 2.45 | 0.05 |

Data are sorted by FDR values

BMI: body mass index, CHR: chromosome, FDR: false discovery rate, B: LIMMA B-statistic from LIMMA

1: Studied CpG identifier
2: CpG locations were mapped using GRCh37 version of the genome from Ensembl platform
Notably, pathway enrichment analysis revealed a significant contribution ($p = 1.94 \times 10^{-10}$) of these genes to the regulation of the olfactory transduction network, which were involved in odor detection and signal processing in the nervous system (Fig. 3). These genes included the olfactory receptors \textit{OR4D2}, \textit{OR51A7}, \textit{OR2T34}, and \textit{OR2Y1} and several downstream effectors, such as \textit{SLC8A1}, \textit{ANO2}, \textit{PDE2A}, \textit{CALML3}, \textit{GNG7}, \textit{CALML6}, \textit{PRKG1}, and \textit{CAMK2D}.

Furthermore, potential associations between olfactory receptor gene methylation status and dietary intakes were screened in a subsample of the MENA cohort.
Notably, methylation at cg02874396 (OR4D2) and cg18482656 (OR2Y1) strongly correlated with daily intakes of total energy ($p < 0.0001$), carbohydrates ($p < 0.0001$), protein ($p < 0.0001$), and fat ($p < 0.0001$).

**Discussion**

Olfaction is considered an important sensorial factor influencing feeding behaviors through modulating food palatability and appetite [10, 13]. Thus, olfactory disruptions at the phenotypic and molecular levels may affect food intake and, consequently, individual predisposition to weight gain and obesity [9]. The current research shows, apparently for the first time, associations between different methylation patterns at olfactory pathway genes and adiposity indicators (BMI and waist circumference), which were independent of age and sex. These genes encompassed olfactory receptors (OR4D2, OR51A7, OR2T34, OR2Y1) and downstream signaling molecules (SLC8A1, ANO2, PDE2A, CALML3, GNG7, PRKG1, OR51A7, CAMK2D, CALML6, OR2T34, and OR2Y1). Figure taken from KEGG reference database (map04740). Pathway enrichment analyses, based on pathDIP ($p = 1.94 \times 10^{-10}$).

**Fig. 3** Pathway mapping of BMI-associated genes within the olfactory transduction network (red boxes). The following genes were computed: SLC8A1, OR4D2, ANO2, PDE2A, CALML3, GNG7, PRKG1, OR51A7, CAMK2D, CALML6, OR2T34, and OR2Y1.
obesity in humans [23]. Also, predicted damaging missense variants in olfactory receptor and protocadherin beta cluster genes were co-localized in subjects with extreme obesity [24]. Similarly, olfactory receptor gene polymorphisms showed evidence of an association with adiposity levels and some eating behaviors, including cognitive dietary restraint, susceptibility to hunger, and eating disinhibition [25].

Regarding odor-evoked transducers, PRKG1, a cGMP-dependent protein kinase, involved in foraging behavior, food acquisition, and energy balance, was located within a variably methylated region associated with BMI in a human cohort [46]. Until now, the specific roles of ANO2, CAMK2D, SLC8A1, PDE2A, CALML3, GNG7, and CALML6 genes in olfactory-related dietary patterns and obesity in humans have not been apparently

Fig. 4 Associations between methylation levels (beta values) at olfactory receptors and dietary intakes. a-d cg02874396, OR4D2 (e-h) cg18482656, OR2Y1
explored; therefore, further investigation in these research areas is warranted.

Potential relationships between eating patterns, olfactory function, and obesity have also been phenotypically studied. In this context, consumers of a Western-style diet (rich in saturated fat and added sugar) presented poorer odor identification ability, worse fat discrimination, and hedonic differences in taste and flavor perception relative to people who consumed a healthier diet [47]. Of note, an increasing BMI has been associated with a decrease in olfactory sensitivity [48]. High BMI was also related to subjective olfactory dysfunction in obese patients [49]. In addition, an impaired olfactory capacity has been reported in obese subjects compared to normal-weight controls [50]. Furthermore, an increase in visceral fat content was associated with a decrease in olfactory function [51].

To the best of our knowledge, this is a pioneer study exploring the role of epigenetics of olfaction in obesity. The strengths of this investigation include the screening of the whole olfactory transduction pathway and the relative large number of DNA samples analyzed. Moreover, methylation-related statistical analyses were adjusted for potential confounding factors such as sex, age, study cohorts, methylation chips, and cell composition variability as well as with corrections for multiple comparisons within the experiments performed in this study. Instead, the limitations of this study encompass those inherent to retrospective association studies including the inability to explain cause-effect relationships, the scientific caution concerning the reproducibility of the findings in other administrative settings having their own peculiar biases, and the difficulties in distinguishing between factors related to increased or decreased risk of developing the disease and those associated with the course of the illness [52]. Other drawbacks of this study comprise the exclusion of expression assays (RNA samples were not available) and the lack of tests evaluating olfactory function. Also, given the number of subjects analyzed and the type of study (association) type I and type II bias cannot be completely excluded despite the statistical settings. Likewise, some methylation relevant data related to obesity could have not been taken into account due to the use of rigorous FDR thresholds in the outcome-related analyses.

Another issue of this study could be the fact that DNA methylation levels were measured in circulating white blood cells instead of olfactory tissues. Interestingly, the expression of some olfactory receptors has been detected in different blood cells, including OR51A7 in leukocytes [53]. Also, our results are in accordance with a previous investigation showing associations between methylation signatures at sweet taste transducing genes, obesity, and carbohydrate intake [26]. Additionally, some studies support that methylation marks in blood cells can mirror those found in other samples, including oral mucosa [54] and subcutaneous adipose tissue [55], suggesting the possible use of the methylome in leukocytes for disease-risk prediction and therapeutic purposes.

In recent years, the implication of different epigenetic processes in the development of obesity is being extensively investigated [56]. Especially, altered DNA methylation patterns can trigger changes in gene expression associated with deregulations of energy homeostasis and weight gain predisposition [7]. In this manner, genes integrating the olfactory pathway are susceptible to covalent epigenetic modifications that might lead to odor perception dysfunction. Establishing an epigenetic basis for olfactory function may help to understand, at least in part, relationships between olfaction capacity, food consumption, and body weight regulation. In turn, this knowledge may contribute to identify epigenetic biomarkers to predict the risk of developing excessive adiposity and associated comorbidities, as well as implement epigenome-based dietary strategies for prevention, prognosis, and treatment of obesity within the era of precision nutrition [57].

Conclusions

The results of this investigation suggest novel relationships between olfactory pathway gene methylation signatures, obesity indices, and dietary intakes.

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Availability of data and materials
The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors’ contributions
JAM conceived the study, revised data, wrote, and critically revised the content of this article. ORL and JIR performed the statistical analysis and prepared the first draft of the manuscript. FIM, JLS, and MAZ wrote and critically revised the content of this article. All authors critically reviewed all drafts and approved the final manuscript.

Ethics approval and consent to participate
All procedures performed in this study were in accordance with the ethical standards of the Research Committee of the University of Navarra and with the 1964 Helsinki Declaration and its 2013 amendments. Informed consent was obtained from all individual participants before included in the study.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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References
1. Friedrich MJ. Global obesity epidemic worsening. JAMA. 2017;318:603.
2. World Health Organization: Obesity and overweight (2017). http://www.who.int/mediacentre/factsheets/fs311/en/. Accessed 03 Mar 2018.
3. Seidel JL, Halbertsd J. The global burden of obesity and the challenges of prevention. Ann Nutr Metab. 2015;66(Suppl 1):27–12.
4. GBD 2015 Obesity Collaborators. Health effects of overweight and obesity in 195 countries over 25 years. N Engl J Med. 2017;377:13–27.
5. Kang JX. Nutritional problems and solutions for the modern health epidemic. J Nutrigenet Nutrigenomics. 2014;7:188–90.
6. Goni L, Milagro FI, Cuervo M, Martínez JA. Single-nucleotide polymorphisms and DNA methylation markers associated with central obesity and regulation of body weight. Nutr Rev. 2014;72:673–90.
7. Martínez JA, Milagro FI, Claycombe KJ, Schalinske KL. Epigenetics in adipose tissue, obesity, weight loss, and diabetes. Adv Nutr. 2014;5:71–81.
8. Apovian CM, Rifkind BM. Perspectives on the global obesity epidemic. Curr Opin Endocrinol Diabetes Obes. 2017;24:307–9.
9. Rolls ET. Taste, olfactory, and food reward value processing in the brain. Prog Neurobiol. 2015;127:128–64–90.
10. Cameron JD, Goldfeld GS, Doucet E. Fasting for 24 h improves nasal chemosensory performance and food palatability in a related manner. Appetite. 2012;58:978–81.
11. Hanci D, Afton H. Hunger state affects both olfactory abilities and gustatory sensitivity. Eur Arch Otorhinolaryngol. 2016;273:1637–41.
12. Sun X, Veldhuijzen MG, Babbis AE, Sinha R, Small DM. Perceptual and brain response to odors is associated with body mass index and postprandial total ghrelin reactivity to a meal. Chem Senses. 2016;41:233–48.
13. Zoon HF, de Graaf C, Boesveldt S. Food odours direct specific appetite. Foods. 2016. https://doi.org/10.3390/foods5010012.
14. Kindleyides S, Beck KL, Walsh DCI, Henderson L, Jayasinghe SN, Golding M, Breier BH. Fat sensation: fatty acid taste and olfaction sensitivity and the link with disinhibited eating behaviour. Nutrients. 2017. https://doi.org/10.3390/nu9080879.
15. Zoon HF, He W, de Wijk RA, de Graaf C, Boesveldt S. Food preference and intake in response to ambient odours in overweight and normal-weight females. Physiol Behav. 2014;133:190–6.
16. Juilliard AK, Al Koborsy D, Fadrol DP, Palouzier-Paulignan B. Nutrient sensing: another chemosensitivity of the olfactory system. Front Physiol. 2017;8:468.
17. Green BG, Nachtigal D, Hammond S, Lim J. Enhancement of retinosomal odors by taste. Chem Senses. 2012;37:77–86.
18. Loper HB, La Sala M, Doston C, Steinele N. Taste perception, associated hormonal modulation, and nutrient intake. Nutr Rev. 2015;73:69–81.
19. Ramos-Lopez O, Roman S, Martinez-Lopez E, Gonzalez-Alcado K, Ojeda-Granados C, Sepulveda-Villegas M, Panduro A. Association of a novel TAS2R38 haplotype with alcohol intake among Mexican-mestizo population. Ann Hepatol. 2015;14:729–34.
20. Ramos-Lopez O, Panduro A, Martinez-Lopez E, Fierro NA, Ojeda-Granados C, Sepulveda-Villegas M, Roman S. Genetic variant in the CD36 gene (rs1765667) is associated with higher fat intake and high serum cholesterol among the population of West Mexico. J Nutr Food Sci. 2015;5:353.
21. Ramos-Lopez O, Panduro A, Martinez-Lopez E, Roman S. Sweet taste receptor TAS1R2 polymorphism (Val191Val) is associated with a higher carbohydrate intake and hyperglycemia/deremia among the population of West Mexico. Nutrients. 2016;8:101.
22. Ramos-Lopez O, Roman S, Martinez-Lopez E, Fierro NA, Gonzalez-Alcado K, Jose-Abrego A, Panduro A. CD36 genetic variation, fat intake and liver fibrosis in chronic hepatitis C virus infection. World J Hepatol. 2016;8:1067–74.
23. Jarick I, Vogel CI, Scherer S, Schaffer H, Hebebrand J, Hinney A, Schera A. Novel common copy number variation for early onset extreme obesity on chromosome 11q11 identified by a genome-wide analysis. Hum Mol Genet. 2011;20:840–52.
24. Mariman EC, Sokolczyk R, Bouwman FG, Aller EE, van Baak MA, Wang P. Olfactory receptor genes cooperate with protocadherin genes in human extreme obesity. Genes Nutr. 2015;10:465.
25. Choquette AC, Bouchard L, Drapeau V, Lemieux S, Tremblay A, Bouchard C, Vohl MC, Périsse L. Association between olfactory receptor genes, eating behavior traits and adiposity: results from the Quebec Family Study. Physiol Behav. 2012;105:772–6.
26. Ramos-Lopez O, Arpín A, Riezu-Boj J, Milagro FI, Mansego ML, Martinez JA, MENA project. DNA methylation patterns at sweet taste transducing genes are associated with BMI and carbohydrate intake in an adult population. Appetite. 2018;120:230–9.
27. Larsen TM, Dalskov S, van Baak M, Jebb S, Kafatos P, Pfeiffer A, Martine JL, Handjiev-Dainiksa T, Kunosov M, Holst C, Saris WH, Astrup A. The Diet, Obesity and Genes (Diogenes) Dietary Study in eight European countries - a comprehensive design for long-term intervention. Obes Rev. 2012;13:76–91.
28. Zulet MA, Bondía-Pons J, Abete I, de la Iglesia R, López-Larregui P, Forlóz L, Navas-Carretero S, Martínez JA. The reduction of the metabolic syndrome in Navarra-Spain (RESMENA-S) study: a multidisciplinary strategy based on chrononutrition and nutritional education, together with dietetic and psychological control. Nutr Hosp. 2011;26:16–26.
29. Petersen M, Taylor MA, Saris WH, Verdich C, Toubro S, Macdonald I, Rössner S, Stich V, Guy-Grand B, Langin D, Martinez JA, Pedersen OD, Holst C, Sørensen TI, Astrup A. Randomized, multi-center trial of two hypo-energetic diets in obese subjects: high- versus low-fat content. Int J Obes. 2006;30:552–60.
30. San-Cristóbal R, Navas-Carretero S, Celis-Morales C, Brennan L, Wahle M, Lovegrove JA, Daniel H, Saris WH, Traczyk S, Marinos Y, Gibney ER, Gibney MJ, Matthews JC, Martinez JA. Analysis of dietary pattern impact on weight status for personalised nutrition through on-line advice: the Food4Me Spanish cohort. Nutrients. 2015;7:5925–37.
31. Martínez-González MÁ, Toledo E, Arós F, Fiol M, Corella D, Salas-Salvadó J, Fiol M, Serra-Majem L, Pintó X, Lamuela-Raventós RM, Soriguera J, Bimbela N, Buil Costales P.
Ruiz-Gutiérrez V, Estruch R, Alonso A. PREDIMED Investigators. Extravirgin olive oil consumption reduces risk of atrial fibrillation: the PREDIMED (Prevención con Dieta Mediterránea) trial. Circulation. 2014;130:18–26.

32. Huerta AE, Navas-Carretero S, Prieto-Hontoria PL, Martínez JA, Moreno-Albaja MJ. Effects of α-lipoic acid and eicosapentaenoic acid in overweight and obese women during weight loss. Obesity (Silver Spring). 2015;23:313–31.

33. Santos JL, Yévenes J, Cataldo LR, Morales M, Galgani J, Arancibia C, Vega J, Olmos P, Flores M, Valderas JP, Pollak F. Development and assessment of the disposition index based on the oral glucose tolerance test in subjects with different glycaemic status. J Physiol Biochem. 2016;72:121–31.

34. Abete I, Gómez-Urquzá AM, Mansego ML, De Arce A, Goyenechea E, Blázquez J, Vázquez-Zabaleta MT, González-Muniesa P, López De Munain A, Martínez JA, Campón J, Milagro FJ. Epigenetic changes in the methylation patterns of KCNQ1 and WT1 after a weight loss intervention program in obese stroke patients. Curr Neurovasc Res. 2015;12:321–33.

35. World Medical Association (WMA). World Medical Association Declaration of Helsinki: ethical principles for medical research involving human subjects. JAMA. 2013;310:2191–4.

36. Navarro-González DJ, Sánchez-Silago L, Pantrana-Delgado J, Fernández-Montero A, Martínez JA. Triglyceride-glucose index (TyG index) in comparison with fasting plasma glucose improved diabetes prediction in patients with normal fasting glucose: The Vascular-Metabolic CUN cohort. Prev Med. 2016;86:99–105.

37. de la Fuente-Arrillaga C, Ruiz Z, Bes-Rastrollo M, Sampson L, Martínez-Juan C, et al. The validity and reproducibility of a food frequency questionnaire validated for the assessment of diet-related inflammatory biomarkers. J Physiol Biochem. 2013;69:485–95.

38. Moreiras O, Cabrera L, Carreño A, Arbuiña M, de la Torre R, Fernández-Real JM, Frühbeck G, Agüera Z, Langohr K, Casanueva FF, de la Torre R. A lower olfactory capacity is related to higher circulating concentrations of endocannabinoid 2-arachidonoylglycerol and higher body mass index. Sci Transl Med. 2010;2:49ra67.

39. Tangrea JA, DeOtte A, Alborzian N, Kurburski A, Allison DB, Maret W. Higher body mass index is associated with a decrease in the taste and olfactory capacity. PLoS One. 2017;12:e0171204.

40. Mansego ML, Garcia-Lacarte M, Milagro FI, Martí A, Martínez JA. Adherence to Mediterranean diet is associated with methylation changes in non-obese and obese patients. Sci Rep. 2017;7:41903.

41. Arpón A, Riezu-Boj JI, Milagro FJ, Martí A, Razquin C, Martínez-González MA, Corella D, Estruch R, Casas P, Fito M, Ros E, Salas-Salvadó J, Martínez AJ. Adherence to Mediterranean diet is associated with methylation changes in inflammation-related genes in peripheral blood cells. J Physiol Biochem. 2016;73:445–55.

42. Houseman EA, Accomando WP, Koestler DC, Christensen BC, Marsit CJ, Nelson HH, Wiencke JK, Kelsey KT. DNA methylation arrays as surrogate measures of cell mixture distribution. BMC Bioinf. 2012;13:86.

43. Tangrea JA, Alborzian N, Kurburski A, Allison DB, Maret W. Higher body mass index is associated with a decrease in the taste and olfactory capacity. PLoS One. 2017;12:e0171204.

44. Kang N, Koo J. Olfactory receptors in non-chemosensory tissues. BMB Rep. 2015;58:407–13.

45. Ihara S, Yoshikawa K, Touhara K. Chemosensory signals and their receptors in the olfactory neural system. Neuroscience. 2013;254:45–53.

46. Ramirez-Tortosa E, Ramallo-del Pozo P, Garcia-de Elía A, Castilla-López C, Pena-Villamizar G, et al. DNA methylation of miRNA coding sequences putatively members. DNA methylation of miRNA coding sequences putatively associated with childhood obesity. Pediatr Obes. 2017;12:29–72.

47. Du P, Kibbe WA, Lin SM. Illumina: a pipeline for processing Illumina microarrays. Bioinformatics. 2008;24:1547–8.

48. Touleimat N, Tost J. Complete pipeline for Infinium (®) Human Methylation 450K BeadChip data processing using subset quantile normalization for accurate DNA methylation estimation. Epigenomics. 2012;4:325–41.

49. Houseman EA, Accomando WP, Koestler DC, Christensen BC, Marsit CJ, Nelson HH, Wiencke JK, Kelsey KT. DNA methylation arrays as surrogate measures of cell mixture distribution. BMC Bioinf. 2012;13:86.

50. Arpón A, Riezu-Boj JI, Milagro FJ, Martí A, Razquin C, Martínez-González MA, Corella D, Estruch R, Casas P, Fito M, Ros E, Salas-Salvadó J, Martínez AJ. Adherence to Mediterranean diet is associated with methylation changes in inflammation-related genes in peripheral blood cells. J Physiol Biochem. 2016;73:445–55.

51. Arpón A, Riezu-Boj JI, Milagro FJ, Martí A, Razquin C, Martínez-González MA, Corella D, Estruch R, Casas P, Fito M, Ros E, Salas-Salvadó J, Martínez AJ. Adherence to Mediterranean diet is associated with methylation changes in inflammation-related genes in peripheral blood cells. J Physiol Biochem. 2016;73:445–55.

52. Mantel N, Haenszel W. Statistical aspects of the analysis of data from retrospective studies of disease. J Natl Cancer Inst. 1959;22:719–48.

53. Malik A, Fiedler J, Fricke K, Ballweg K, Pfaff M, Krautwurst D. Class I odorant receptors, TAS1R and TAS2R taste receptors, are markers for subpopulations of circulating leukocytes. J Leukoc Biol. 2015;97:533–45.

54. San-Cristobal R, Navas-Carretero S, Milagro FJ, Riezu-Boj JI, Guevara-C e, Celis-Morales C, Livingstone KM, Brennan L, Lovegrove JA, Daniel H, Satsi WH, Traczyk I, Manios Y, Gibney ER, Gibney MJ, Mathers JC, Martinez JA. Gene methylation parallelsisms between peripheral blood cells and oral mucosa samples in relation to overweight. J Physiol Biochem. 2016;73:465–74.

55. Crujeiras AB, Díaz-Lagares A, Sandoval J, Milagro FJ, Navas-Carretero S, Careira MC, Gomez A, Hervas D, Monteiro MP, Casanueva FF, Esteller M, Martinez JA. DNA methylation map in circulating leukocytes mirrors subcutaneous adipose tissue methylation patterns: a genome-wide analysis from non-obese and obese patients. Sci Rep. 2017;7:41903.

56. Milagro FJ, Mansego ML, De Miguel C, Martinez JA. Dietary factors, epigenetic modifications and obesity outcomes: progresses and perspectives. Mol Asp Med. 2013;34:782–812.

57. Ramos-Lopez O, Milagro FJ, Allayee H, Chmurzynska A, Choi MS, Curi R, De Caterina R, Ferguson LR, Góni L, Kang JX, Kohlmeier M, Marti A, Moreno LA, Pérusse L, Prasad C, Qi L, Reifen R, Rieu-Bou JI, San-Cristobal R, Santos JL, Martinez JA. Guide for current nutrigenetic, nutrigenomic, and nutriepigenetic approaches for precision nutrition involving the prevention and management of chronic diseases associated with obesity. J Nutrigenet Nutrigenomics. 2017;10:43–62.

58. Skrandies W, Zschiegisch R. Olfactory and gustatory functions and its relation to body weight. Physiol Behav. 2015;142:1–4.

59. Patel ZM, DelGaudio JM, Wise SK. Higher body mass index is associated with subjective olfactory dysfunction. Behav Neurol. 2015;2015:675635.

60. Pastor A, Fernández-Aranda F, Fito M, Jiménez-Marcuña S, Botella C, Fernández-Real JM. Frühbeck G, Tinahones FJ, Fagundo AB, Rodriguez J, Agüera Z, Langohr K, Casanueva FF, de la Torre R. A lower olfactory capacity is related to higher circulating concentrations of endocannabinoid 2-arachidonoylglycerol and higher body mass index in women. PLoS One. 2016;11:e0148734.

61. Fernández-García JC, Alcaide J, Santiago-Fernandez C, Roca-Rodriguez MM, Agüera Z, Bahías R, Botella C, de la Torre R, Fernández-Real JM, Frühbeck G, Gómez-Ambrosi J, Jiménez-Marcuña S, Menchón JM, Casanueva FF, Fernández-Aranda F, Tinahones FJ, Garrido-Sánchez L. An increase in visceral fat is associated with a decrease in the taste and olfactory capacity. PLoS One. 2017;12:e0171204.