

# A low-fat diet up-regulates expression of fatty acid taste receptor gene *FFAR4* in fungiform papillae in humans: a co-twin randomised controlled trial

Andrew Costanzo<sup>1†</sup>, Dongli Liu<sup>1†</sup>, Caryl Nowson<sup>2</sup>, Konsta Duesing<sup>3</sup>, Nicholas Archer<sup>4</sup>, Steve Bowe<sup>5</sup> and Russell Keast<sup>1\*</sup>

<sup>1</sup>*School of Exercise and Nutrition Sciences, Centre for Advanced Sensory Science, Deakin University, Geelong, VIC 3220, Australia*

<sup>2</sup>*School of Exercise and Nutrition Sciences, Institute for Physical Activity and Nutrition, Deakin University, Geelong, VIC 3220, Australia*

<sup>3</sup>*Health & Biosecurity, Commonwealth Scientific and Industrial Research Organisation (CSIRO), North Ryde, NSW 2113, Australia*

<sup>4</sup>*Agriculture & Food, CSIRO, North Ryde, NSW 2113, Australia*

<sup>5</sup>*Biostatistics Unit, Faculty of Health, Deakin University, Geelong, VIC 3220, Australia*

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## Abstract

Fatty acid taste (FAT) perception is involved in the regulation of dietary fat intake, where impaired FAT is associated with increased fatty food intake. There are a number of FAT receptors identified on human taste cells that are potentially responsible for FAT perception. Manipulating dietary fat intake, and in turn FAT perception, would elucidate the receptors that are associated with long-term regulation of FAT perception. The present study aimed to assess associations between diet-mediated changes to FAT receptors and FAT perception in humans. A co-twin randomised controlled trial was conducted, where each matching twin within a pair were randomly allocated to either an 8-week low-fat (LF; <20 % energy fat) or an 8-week high-fat (HF; >35 % energy fat) diet. At baseline and week 8, fungiform papillae were biopsied in the fasted state and FAT receptor gene expressions (cluster of differentiation 36 (*CD36*), free fatty acid receptor 2 (*FFAR2*), *FFAR4*, G protein-coupled receptor 84 (*GPR84*) and a delayed rectifying K<sup>+</sup> channel (K<sup>+</sup> voltage-gated channel subfamily A member 2; *KCNA2*)) were measured using RT-PCR; and FAT threshold (FATT) was assessed using three-alternate forced choice methodology. Linear mixed models were fitted, adjusting for correlation between co-twins. Intake was compliant with the study design, with the LF and HF groups consuming 14.8 and 39.9 % energy from fat, respectively. Expression of *FFAR4* increased by 38 % in the LF group ( $P = 0.023$ ; time–diet interaction  $P = 0.063$ ).  $\Delta FFAR4$  ( $\Delta$ , week 8–baseline) was associated with  $\Delta$ fat intake (g) ( $\beta = -159.4$ ;  $P < 0.001$ ) and  $\Delta FATT$  ( $\beta = -8.8$ ;  $P = 0.016$ ). In summary, *FFAR4* is involved in long-term diet-mediated changes to FAT perception. Manipulating dietary fat intake, and therefore *FFAR4* expression, might aid in reducing taste-mediated passive overconsumption of fatty foods.

**Key words:** Fat intake: Fatty acid taste: Randomised controlled trials: Free fatty acid receptor 4: G protein-coupled receptor 120

The chemoreception of fatty acids (FA) in the oral cavity, also known as fatty acid taste (FAT), is involved in the regulation of dietary fat intake<sup>(1–3)</sup>. Individuals with impaired FAT sensitivity are more likely to consume greater amounts of dietary fat due to compromised cephalic phase and post-ingestive satiety hormone responses following oral fat exposure<sup>(4–9)</sup>. This is reflected throughout the alimentary canal, as individuals who are less sensitive to FAT in the oral cavity also have reduced hormonal response

following FA stimulation in the gastrointestinal tract (GIT)<sup>(8)</sup>. FAT sensitivity is attenuated by oral fat exposure and conversely can be increased by long-term reduced dietary fat intake<sup>(10–12)</sup>, which may be due to the regulation of some or all FAT receptors in the oral cavity.

Three types of lingual papillae – fungiform, foliate and circumvallate – house taste bud cells (TBC) which express FAT receptors. A recent study of our group localised several

**Abbreviations:** CD36, cluster of differentiation 36; FAT, fatty acid taste; FATT, FAT threshold; FFAR, free fatty acid receptor; GIT, gastrointestinal tract; GPR, G protein-coupled receptor; HF, high fat; KCNA2, delayed rectifying K<sup>+</sup> channel (K<sup>+</sup> voltage-gated channel subfamily A member 2); LF, low fat; TBC, taste bud cells; TRA, Twins Research Australia.

\* **Corresponding author:** Russell Keast, email [russell.keast@deakin.edu.au](mailto:russell.keast@deakin.edu.au)

† These authors contributed equally to this work and are co-first authors.

candidate FAT receptor genes – cluster of differentiation 36 (*CD36*), free fatty acid receptor 2 (*FFAR2*), *FFAR4*, G protein-coupled receptor 84 (*GPR84*) and a delayed rectifying K<sup>+</sup> channel (K<sup>+</sup> voltage-gated channel subfamily A member 2; *KCNK2*) – in TBC of human fungiform papillae<sup>(13)</sup>. Each of these receptors has specificities with regard to the type of FA they bind<sup>(14)</sup>; however, the exact receptors that might be responsible for oral chemoreception of FA in humans remain unresolved. Previously, we showed that increased *CD36* was associated with short-term fat intake, particularly saturated fat, and acute dislike of fatty foods<sup>(13)</sup>. However, no association was found between any FAT receptors and FAT perception. Since FAT sensitivity can be modified by dietary fat intake<sup>(10–12)</sup>, it may be possible to use a dietary fat intervention to elucidate the receptor that might be responsible for long-term changes in FAT perception.

While no human studies have investigated changes in FAT receptors in response to changes in the dietary fat, animal studies have been conducted albeit with conflicting results. An 8-week high-fat (HF) (40 % energy from fat) diet induced significant down-regulation of *CD36* in rat circumvallate TBC<sup>(3)</sup>. However, this down-regulation was not observed in fasted mice fed an HF (34.2 %) diet for 4 weeks<sup>(15)</sup>. Another study found that protein levels of *CD36*, rather than mRNA, decreased, while *FFAR4* protein increased in raft membranes of human and mouse fungiform TBC following a 2-month HF (40 % energy from fat) diet<sup>(16)</sup>. As for short-term, an acute HF (30 % energy from fat) oral exposure triggered an immediate decrease in *CD36* protein in mice circumvallate TBC, which returned to preexposure levels following 11 h of fasting, with no obvious effect seen for *FFAR4* protein<sup>(17)</sup>. The discrepancies between the studies may be attributed to the fat content, the different area of the tongue used for analysis and/or fasting status. Diet-mediated regulation of FAT receptors has not been investigated in humans. Manipulating dietary fat intake, and in turn FAT taste perception, would elucidate the key receptors that are associated with long-term regulation of oral chemoreception of FA.

The present study aimed to determine the FAT receptors that are responsible for diet-mediated changes to FAT perception in humans. The aim was accomplished by assessing the association between changes to key FAT receptor genes expressed in fungiform papillae (as determined by Liu *et al.*<sup>(13)</sup>) and FAT perception following an 8-week dietary fat intervention. A secondary aim was to assess the association between changes in macronutrient intake and FAT receptor gene expressions.

## Subjects and methods

The study presented here is an 8-week follow-up analysis of a previous study where the baseline data have been published<sup>(13)</sup>.

### Participants

A co-twin design was chosen as it controls for age, common environmental and partial genetic factors shared by co-twins in each experimental group. Twins Research Australia (TRA) invited via mail 1881 twin pairs (3762 individuals) from the Melbourne metropolitan area to participate in a larger study on the effect of diet and genes on FAT<sup>(10)</sup>. Twins were eligible to participate in

the study if they were aged between 18 and 69 years, were able to attend three laboratory sessions in Burwood, Victoria and were willing to alter their diet for a period of 8 weeks. Both monozygotic and dizygotic twin pairs were invited to participate. Subjects were excluded from recruitment if they had any dairy allergies and intolerances, if they had any illnesses preventing them from eating foods included in the study, or if they were pregnant or lactating. Due to the nature of the twin study design, if one individual from a twin pair was excluded or withdrawn from the study, their co-twin was also excluded. In sixty-six pairs, both twins expressed interest in participating and were then screened for eligibility. Forty-six twin pairs (ninety-two individuals) aged between 18 and 68 years were recruited into the larger study; however, only thirteen pairs (twenty-six individuals; ten monozygotic pairs, three dizygotic pairs) consented to the additional testing described in this paper (Fig. 1). Co-twins from each pair were randomised into either a low-fat (LF) or an HF diet, where one twin from each pair was allocated to the LF diet and the other twin allocated to the HF diet. Prior to recruitment, a block randomised sequence was generated with blocks of size two. TRA was responsible for recruitment and therefore characteristics of the participants were blinded to the researchers. Participants were allocated to the randomised sequence based on their TRA twin number; therefore, allocation of participants to diet group was concealed. Due to the nature of the intervention, blinding of participants was not feasible.

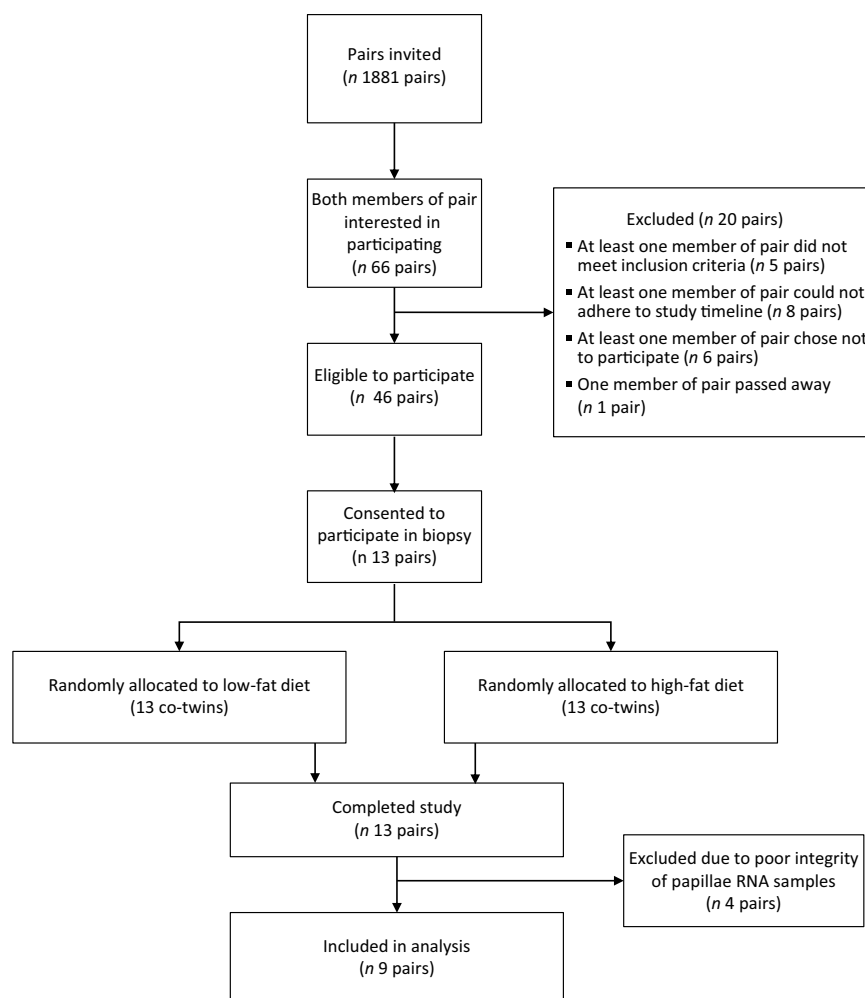
### Ethics

The present study was conducted according to the guidelines laid down in the Declaration of Helsinki, and all procedures involving human subjects/patients were approved by the Deakin University Human Research Ethics Committee (ID: 2013-163). Written informed consent was obtained from all subjects. The present study is registered as a clinical trial with the Australian New Zealand Clinical Trials Registry (ID: ACTRN12613000466741; <https://www.anzctr.org.au/>).

### Study outline

Participants attended two tasting sessions and two biopsy sessions at the Centre for Advanced Sensory Science at Deakin University, Burwood, Victoria. Recruitment and data collection occurred between July 2014 and May 2016. The first tasting session occurred on the day prior to beginning the dietary intervention, and the second tasting session occurred 8 weeks later on the last day of the intervention. Tasting sessions were conducted in a temperature- and sound-controlled environment with a 15-min break in the middle of their session to prevent fatigue. Participants were asked to avoid eating or drinking anything but water and to avoid brushing their teeth or using mouthwash up to an hour prior to each tasting session. Tasting sessions measured FAT threshold (FAT<sub>T</sub>) for oleic acid (C18:1) and anthropometric measurements. A 24-h food recall was collected by a nutritionist during the first session. Between tasting sessions, participants recorded six 24-h diet records, two weekdays and one weekend during the first 4 weeks, and two weekdays and one weekend during the final 4 weeks of the trial.

Collection of fungiform papillae (biopsy) occurred the day after each tasting session. Participants fasted for at least 10 h prior to each fungiform papillae collection. Fungiform papillae tissue



**Fig. 1.** Study CONSORT flow chart diagram.

and serum blood samples were collected. After the biopsy session, participants were provided with an LF or HF breakfast snack depending on the diet that they were allocated.

### Dietary intervention

The LF diet was defined as <20 % of energy from fats and the HF diet was defined as >35 % of energy from fats. These values were chosen as they fall outside the acceptable macronutrient distribution range for fat intake (20–35 %)<sup>(18)</sup>. Participants on the HF diet were encouraged to choose foods higher in monounsaturated and polyunsaturated fats rather than saturated fats in order to maintain a healthy diet. A diet booklet for each diet was created with the aid of an accredited practicing dietitian, which described the parameters of each diet, a list of foods that should be and should not be eaten and some example recipes that adhere to the diet. Participants were given the HF or LF booklet, their assigned diet was explained and they were taught how to interpret a nutrition information panel in order to identify the food that were acceptable for their diet.

Participants were requested to start the assigned diet the day after the baseline measurement. As food were not provided in the present

study, food choice was up to the participants. To maximise adherence to the diets, participants were contacted via phone fortnightly and questioned on their dietary habits. If the researcher felt that participants were not following the diet adequately, they were provided with suggestions and encouragement to aid in diet adherence. Participants were also asked a series of questions to ensure that they did not experience any negative effects from the diet. These questions included ‘Do you feel like you have less energy since starting the diet?’ ‘Do you feel like your weight has changed significantly since starting the diet?’ ‘Is the diet affecting your day-to-day activities?’ If the researcher felt that participants were suffering from major negative effects due to the diet (e.g. nausea, inability to work), they would be asked to stop the diet and were dropped out of the trial.

The first three completed diet diaries were inspected and reviewed at week 4 to assess compliance to the assigned diet.

### Dietary assessment

A single three-pass 24-h dietary recall<sup>(19)</sup> of the day prior to the first tasting session was conducted by a trained nutritionist. For the ongoing diet records throughout the trial, participants were asked to avoid filling out diet records on a non-standard day (e.g. if they attended a wedding reception). They were

taught to quantify foods in standard serving sizes (cups, teaspoons, tablespoons, etc.) using a food model booklet and asked to weigh their food and drinks wherever possible. Details such as brand, cooking method and foods additives (e.g. sugar added to coffee) were included in the diet records.

Food recall and records were analysed for carbohydrate, protein, fat and fibre intake (g and % of energy) using computer software FoodWorks (version 8; Xyris).

### Anthropometry

Body weight was measured after removal of shoes, heavy clothing and any items in their pockets using electronic scales (OHAUS NV4101), and height was measured using a free-standing stadiometer (SECA). BMI was calculated as weight (kg)/height (m)<sup>2</sup>.

### Fatty acid taste receptor expression

Fungiform papillae biopsy was conducted without anaesthetic by a registered doctor following the procedure described previously<sup>(20,21)</sup>. Fungiform papillae were chosen as they were the least invasive biopsy target and provide the best result compared with other oral sampling methods<sup>(22)</sup>. For each participant, up to eight fungiform papillae were collected; four papillae chosen from the left side of the tongue and four from the right side of the tongue, which were chosen at random sites within the fungiform region of the tongue by the doctor. The eight papillae were pooled as an individual sample to reduce the impact of variation from different sites of collection. The gene expression of FAT receptors found within fungiform papillae tissue<sup>(13)</sup> including *CD36*, *FFAR2*, *FFAR4*, *GPR84* and *KCNA2* was analysed with real-time RT-PCR. RNA was extracted from the pooled fungiform papillae samples using TRIzol reagent (Life Technologies) following the manufacture's protocol. The purified RNA pellet was dissolved in 20 µl RNase-free water treated with RNase-free DNase set (Qiagen) and quantified with NanoDrop ND-1000 spectrophotometer. The RNA integrity was measured with Bioanalyser 2100 (Agilent Technologies) and the cut-off value was set at 5. For the RT-PCR, 1 µg of total RNA was used to synthesise complementary DNA (cDNA) using the High-Capacity cDNA Reverse Transcription kit (Applied Biosystems). Each cDNA sample was diluted 1:5 at first. Standards were then prepared with a serial dilution of 1:5 from the top standard (an aliquot of all the 1:5 dilution cDNA samples). For the standard curve, one gene copy was assigned to the lowest concentrated standard, which had a Cp value. Gene concentration of other standards were set accordingly based on the dilution value. The notional concentration of each sample was calculated based on the standard curve (log concentration against Cp). The expression of the genes of interest were analysed with the Taqman gene expression assays (Life Technologies) (Table 1). For each gene analysis, a negative control of the sample that had not been reversely transcribed was included. Housekeeping genes glyceraldehyde 3-phosphate dehydrogenase (*GAPDH*) and 60S acidic ribosomal protein P0 (*RPLP0*) were included for normalising the transcript numbers.

**Table 1.** Taqman gene expression assays used for the real-time RT-PCR analysis

Gene	Assay ID	Description
<i>CD36</i>	Hs01567185_m1	Probe spans exons
<i>FFAR4</i>	Hs00699184_m1	Probe spans exons
<i>FFAR2</i>	Hs00271142_s1	Probes are within single exon
<i>GPR84</i>	Hs01874713_s1	Probes are within single exon
<i>KCNA2</i>	Hs04187587_g1	Probe spans exons
<i>RPLP0</i>	Hs99999902_m1	Probe spans exons
<i>GAPDH</i>	Hs02758991_m1	Probe spans exons

*CD36*, cluster of differentiation 36; *FFAR4*, free fatty acid receptor 4; *FFAR2*, free fatty acid receptor 2; *GPR84*, G protein-coupled receptor 84; *KCNA2*, delayed rectifying K<sup>+</sup> channel (K<sup>+</sup> voltage-gated channel subfamily A member 2); *RPLP0*, 60S acidic ribosomal protein P0; *GAPDH*, glyceraldehyde 3-phosphate dehydrogenase.

### Fatty acid taste threshold

Fat-free milk (Devondale) solutions containing C18 : 1 (Sigma Aldrich) at varying concentrations (0.02, 0.06, 1.00, 1.40, 2.00, 2.80, 3.80, 5.00, 6.40, 8.00, 9.80, 12.00 and 20.00 mM) were prepared using established methods<sup>(22)</sup>. Control samples were prepared in the same way but without added C18 : 1.

FATT for C18 : 1 were determined using an ascending series three-alternate forced choice methodology<sup>(10)</sup>. To prevent confounding non-taste sensory inputs, participants wore nose clips and all tests were conducted under red light. FATT was transformed to an ordinal variable ranging from 0 to 12, with a higher score implying lower FAT sensitivity<sup>(2)</sup>.

### Statistical analysis

Analyses were conducted using computer software SPSS (v24.0; IBM). Null hypotheses were rejected at *P* < 0.05. The present study is a secondary analysis of participants from a larger study<sup>(10)</sup>, who provided consent to additional testing; therefore, sample size of the present study was determined by feasibility of recruitment.

The effect of the diet on FAT receptor gene expressions was assessed using linear mixed models including diet (LF and HF), time (baseline and week 8) and the time–diet interaction as fixed effects; with twin pair as a random effect and co-twin as the subject with repeated measures to account for the correlation between co-twins. A *post hoc* Sidak's test was used to compare means between each fixed effect and to correct for multiple comparisons. Time–diet interaction, *post hoc* Sidak's test and CI are reported. The above model was also used to assess differences between diet groups at baseline and to determine compliance to the prescribed diets.

To explore the strength of the association ( $\hat{\beta}$ ) between change in FATT and change in FAT receptor gene expression over the 8-week period, the same model as above was conducted with  $\Delta$  FATT ( $\Delta$ , week 8 – baseline) as the outcome and  $\Delta$  FAT gene expression as a fixed effect and twin pair as a random effect. Pearson's correlations (*r*) for each association are also reported for descriptive purposes. The same analysis was repeated for the association between  $\Delta$  macronutrient intake (total fat, saturated fat, monounsaturated fat, polyunsaturated fat, carbohydrate, protein and dietary fibre) (g) and  $\Delta$  FAT gene expression.





**Table 2.** Baseline characteristics of participants (Mean values and standard deviations)

	All participants (n 18)		LF diet (n 9)		HF diet (n 9)	
	Mean	SD	Mean	SD	Mean	SD
Age (years)	41.6	16.5	—	—	—	—
Weight (kg)	72.5	17.2	71.9	17.5	73.0	18.0
BMI (kg/m <sup>2</sup> )	26.9	6.3	26.7	6.1	27.2	6.9
Female (n)	16		8		8	

LF, low fat; HF, high fat.

## Results

Thirteen twin pairs were recruited in the present study. However, four twin pairs were not included in the RT-PCR analysis of gene expression due to low concentration or poor integrity of the collected RNA samples according to the Nanodrop and Bioanalyser analyses. As a result, we present here nine pairs (seven monozygotic and two dizygotic; eight female pairs and one male pair) with valid samples that underwent RT-PCR analysis (Fig. 1). No harmful or unintended effects were observed in either group. None of the participants reported any significant changes to their weight, energy levels or ability to perform day-to-day tasks throughout the dietary intervention. Baseline characteristics of the participants are described in Table 2. No significant differences in baseline characteristics were observed between diet groups.

### Dietary compliance

Intake of energy from fat throughout the trial was within the aim of the study, with the LF group consuming 14.8 (95 % CI 10.0, 19.7) % energy from fat and the HF group consuming 39.9 (95 % CI 35.0, 44.7) % energy from fat (online Supplementary Table S1). Intake of energy from fat at baseline was already considered high in the HF diet group at 36.4 (95 % CI 31.6, 41.3) %, so there was no significant change in fat intake across the trial in this group. A significant decrease in the intake of energy was observed from fat in the LF diet group (−20.3 (95 % CI −26.2, −14.4) %,  $P < 0.001$ ). No significant change in total energy intake was observed in either diet group, though a between-group difference was observed at week 8 (2.3 (95 % CI 0.5, 4.2) MJ,  $P = 0.021$ ).

### Effect of dietary fat intake on fatty acid taste threshold

No difference in FATT was observed between the LF and HF diet groups at baseline (Table 3). Over the 8 weeks, FATT decreased by 76 % in the LF diet group ( $P < 0.001$ ) and increased by 23 % in the HF diet group ( $P = 0.049$ ). At week 8, FATT in the HF diet group was 3.8 times higher than the LF diet group ( $P < 0.001$ ), and a significant time–diet interaction was observed for FATT ( $P < 0.001$ ) (Table 3).

### Effect of dietary fat intake on fatty acid taste receptor gene expressions

There was evidence for a time–diet interaction for *FFAR4* expression ( $P = 0.063$ ), as expression increased by 38 % in the LF diet

group ( $P = 0.023$ ) (Table 3). No significant time–diet interactions were observed for *CD36*, *GPR84*, *FFAR2* and *KCNA2* expression, though *GPR84* expression was 61 % greater in the LF diet group compared with the HF diet group at baseline.

### Associations between fatty acid taste threshold and fatty acid taste receptor gene expressions

The relationship between FAT sensitivity and FAT receptor expression was assessed by comparing change ( $\Delta$ ) in FATT with change in receptor expression from baseline to week 8 (Table 4) (Fig. 2). A significant negative association was observed between  $\Delta$  FATT and  $\Delta$  *FFAR4*, indicating that as *FFAR4* expression increased, FATT also decreased concurrently (i.e. sensitivity to FAT increased). Conversely, the positive association between  $\Delta$  FATT and  $\Delta$  *GPR84* indicates that as *GPR84* expression increased, FATT also increased (i.e. sensitivity to FAT decreased). There were no significant associations between  $\Delta$  FATT and  $\Delta$  *CD36*,  $\Delta$  *FFAR2* or  $\Delta$  *KCNA2*.

### Associations between dietary macronutrient intake and fatty acid taste receptor gene expressions

Significant negative associations were observed between  $\Delta$  *FFAR4* and  $\Delta$  fat intake (g) ( $\hat{\beta} = -159.4$ ;  $r = -0.744$ ;  $P < 0.001$ ),  $\Delta$  saturated fat intake (g) ( $\hat{\beta} = -79.4$ ;  $r = -0.759$ ;  $P < 0.001$ ),  $\Delta$  mono-unsaturated fat intake (g) ( $\hat{\beta} = -53.4$ ;  $r = -0.711$ ;  $P = 0.001$ ) and  $\Delta$  polyunsaturated fat intake (g) ( $\hat{\beta} = -14.8$ ;  $r = -0.533$ ;  $P = 0.023$ ). There was a statistical trend for a negative association between  $\Delta$  *KCNA2* and  $\Delta$  polyunsaturated fat intake (g) ( $\hat{\beta} = -72.7$ ;  $r = -0.459$ ;  $P = 0.056$ ). Finally, there was a significant positive association between  $\Delta$  *FFAR2* and  $\Delta$  dietary fibre intake (g) ( $\hat{\beta} = 22.5$ ;  $r = 0.560$ ;  $P = 0.016$ ). No significant associations were observed for  $\Delta$  *CD36* or  $\Delta$  *GPR84*, though a significant association was observed between baseline *GPR84* expression and intake of energy from dietary fibre (%) ( $\hat{\beta} = 8.8$ ;  $r = 0.399$ ;  $P = 0.023$ ).

## Discussion

The present study assessed changes in fasting expression of FAT receptor genes following 8 weeks of LF or HF dietary intake, for the first time in humans. It is well established that FAT sensitivity is modulated by dietary fat intake<sup>(10–12)</sup>. It was hypothesised that expression of FAT receptors would be modulated due to dietary fat intake, similar to rodent models<sup>(3,17)</sup>. However, only *FFAR4* was affected by the dietary intervention, with expression of *FFAR4* increasing by 38 % in the LF diet group from baseline to week 8. While the magnitude of reduction in *FFAR4* expression in the HF group (3 %) was much lower than the increase seen in the LF group, this is likely because in the HF group, fat (g) increased by only 34 %; whereas in the LF group, there was a 72 % reduction in fat consumed. Further, change in *FFAR4* expression was associated with change in FATT, or in essence, as *FFAR4* increased, FAT sensitivity also increased concurrently. This is supported by the association between *FFAR4* and intake of fat, saturated fat, monounsaturated fat and polyunsaturated fat, indicating that dietary fat with any level of FA saturation down-regulates *FFAR4*. These results indicate that *FFAR4* expressed in fungiform papillae may be responsible

**Table 3.** Means and between-group differences in fatty acid taste threshold (FATT) and fatty acid taste receptor gene expression levels (relative gene copy number) in fungiform papillae taste bud cells over the 8-week trial† (Mean values and 95 % confidence intervals)

	LF diet		HF diet		Between-group difference		Time–diet interaction ( <i>P</i> )
	Mean	95 % CI	Mean	95 % CI	Mean	95 % CI	
<b>FATT</b>							<0.001
Baseline	7.8	5.8, 9.7	7.4	5.4, 9.3	−0.4	−3.1, 2.4	
Week 8	1.9***	0.4, 3.4	9.1*	7.6, 10.6	7.2†††	5.1, 9.3	
<b>CD36</b>							0.248
Baseline	1.20	0.65, 1.75	1.26	0.71, 1.81	0.06	−0.56, 0.68	
Week 8	1.39	0.87, 1.92	1.10	0.58, 1.63	−0.29	−0.88, 0.29	
<b>FFAR4</b>							0.063
Baseline	0.84	0.57, 1.12	0.93	0.66, 1.21	0.09	−0.30, 0.48	
Week 8	1.16*	0.89, 1.43	0.90	0.62, 1.17	−0.26	−0.65, −0.12	
<b>FFAR2</b>							0.409
Baseline	0.95	0.53, 1.36	0.60	0.18, 1.01	−0.35	−0.93, 0.23	
Week 8	0.93	0.48, 1.38	0.75	0.30, 1.20	−0.18	0.86, 0.50	
<b>GPR84</b>							0.214
Baseline	0.057	0.020, 0.095	0.022	−0.015, 0.060	−0.035†	−0.069, −0.001	
Week 8	0.041	0.009, 0.074	0.029	−0.003, 0.062	−0.012	−0.025, 0.000	
<b>KCNA2</b>							0.460
Baseline	0.17	0.12, 0.22	0.18	0.14, 0.23	0.01	−0.04, 0.07	
Week 8	0.20	0.12, 0.28	0.19	0.11, 0.26	−0.01	−0.12, 0.09	

LF, low fat; HF, high fat; *CD36*, cluster of differentiation 36; *FFAR4*, free fatty acid receptor 4; *FFAR2*, free fatty acid receptor 2; *GPR84*, G protein-coupled receptor 84; *KCNA2*, delayed rectifying K<sup>+</sup> channel (K<sup>+</sup> voltage-gated channel subfamily A member 2). Significantly different from baseline: \* *P* < 0.05, \*\*\* *P* < 0.001. Significant between-group difference: † *P* < 0.05, ††† *P* < 0.001. ‡ Between-group difference calculated as HF diet – LF diet. Estimated means, CI and *P* values obtained under a mixed model including twin pair as a random effect. *Post hoc* Sidak's test, CI and time–diet interaction are reported.

**Table 4.** Associations between Δ fatty acid taste threshold (FATT) and Δ fatty acid taste receptor gene expressions\*

	Δ FATT		
	<i>r</i>	$\hat{\beta}$	<i>P</i>
Δ <i>CD36</i>	−0.061	−0.8	0.822
Δ <i>FFAR4</i>	−0.590	−8.8	0.016
Δ <i>FFAR2</i>	0.270	3.4	0.311
Δ <i>GPR84</i>	0.517	68.7	0.040
Δ <i>KCNA2</i>	−0.302	−21.3	0.256

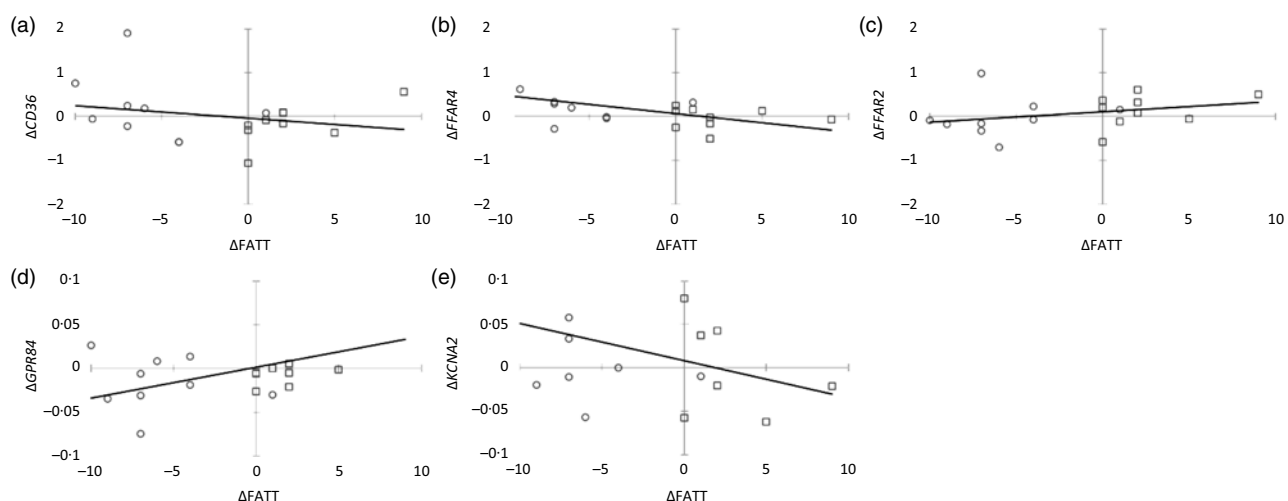
Δ, Week 8 – baseline; *CD36*, cluster of differentiation 36; *FFAR4*, free fatty acid receptor 4; *FFAR2*, free fatty acid receptor 2; *GPR84*, G protein-coupled receptor 84; *KCNA2*, delayed rectifying K<sup>+</sup> channel (K<sup>+</sup> voltage-gated channel subfamily A member 2). \*  $\hat{\beta}$  and *P* values obtained under a mixed model including twin pair as a random effect and time as a repeated effect; *r* obtained using Pearson's correlation for descriptive purposes only.

for long-term changes in FAT perception. To that point, increasing expression of *FFAR4* might lead to an increased secretion of intestinal peptides following chemoreception of FA in the oral cavity or GIT<sup>(8)</sup>, reducing subsequent desire to eat and therefore energy intake. Also, since systemic *FFAR4* is involved in the facilitation of energy homeostasis<sup>(23)</sup>, brown fat activation<sup>(24)</sup> and inflammation<sup>(25)</sup>, the ability to up-regulate *FFAR4* via dietary intervention may have the potential to manage obesity and metabolic disease.

Interestingly, there was no effect of diet on *CD36* in fungiform papillae in the present study, which contradicts the finding in our previous research<sup>(13)</sup>. In the present study, there was a small trend in the same direction as *FFAR4*, in that *CD36* increased by 16 % in the LF diet and decreased by 13 % in the HF diet after

8 weeks; however, no significant time–diet interaction was observed. This is likely because the baseline dietary data were based on a single 24-h recall, whereas the intervention dietary data were based on six diet records over the course of 8 weeks. If participants consumed an HF meal on the night before testing, there might have been some residual effect on gene expression. Short-term oral exposure to dietary fat in mice decreased *CD36* levels by 2-fold within 1 h of re-feeding<sup>(17)</sup>. However, 11 h of fasting returned *CD36* levels to preprandial levels. Therefore, it is possible that TBC up- and down-regulate *CD36* relatively quickly following acute oral exposure to fat compared with *FFAR4*. We speculate that the role of *CD36* may be to mediate short-term response to dietary fat and is down-regulated throughout an eating event, then up-regulated to premeal expression following a prolonged fast; whereas *FFAR4* is involved in regulating long-term response to dietary fat between meals with greater gradual change in expression over longer periods of time. Analysis of *CD36* in human papillae immediately before and after consumption of dietary fat is necessary to confirm this. Some studies have also shown association between *CD36* and hedonic preference for fatty foods in humans<sup>(13)</sup> and rodents<sup>(26–29)</sup>, which strengthens the argument that the role of *CD36* is more specific to mediating fat intake within a given meal. There is also evidence of interaction between *FFAR4* and *CD36* in TBC<sup>(30)</sup>, though the significance of this interaction remains unclear.

While the dietary intervention was not designed to control for polyunsaturated fat intake as participants were able to choose to consume any food sources of fat, intake of polyunsaturated fat increased on the LF diet and decreased on the HF diet. Despite this, time–diet interaction for *KCNA2*, which is the



**Fig. 2.** Scatterplots of  $\Delta$  fatty acid taste threshold (FATT) and  $\Delta$  fatty acid taste receptor gene expressions. Circle markers (○) indicate participants on the low-fat diet ( $n=9$ ) and square markers (□) indicate participants in the high-fat group ( $n=9$ ). (a)  $\Delta$  FATT v.  $\Delta$  cluster of differentiation 36 (*CD36*); (b)  $\Delta$  FATT v.  $\Delta$  free fatty acid receptor 4 (*FFAR4*); (c)  $\Delta$  FATT v.  $\Delta$  free fatty acid receptor 2 (*FFAR2*); (d)  $\Delta$  FATT v.  $\Delta$  G protein-coupled receptor 84 (*GPR84*); (e)  $\Delta$  FATT v.  $\Delta$  K<sup>+</sup> voltage-gated channel subfamily A member 2 (*KCNA2*).

receptor gene specific to PUFA chemoreception, was not significant. An 18% increase in *KCNA2* was observed in the LF diet group, which was in the hypothesised direction, and no change in *KCNA2* in the HF diet group was likely due to the relatively minor increase in fat intake over 8 weeks. There was a statistical trend for a negative association between  $\Delta$  *KCNA2* and  $\Delta$  polyunsaturated fat intake ( $P=0.056$ ), which indicates some effect of polyunsaturated fat intake on *KCNA2*. Therefore, it could be suggested that *KCNA2* might be involved in the regulation of PUFA intake, where an individual who is not meeting the PUFA intake requirement might express greater *KCNA2* levels on TBC to promote greater subsequent PUFA intake. However, a dietary intervention that is well-controlled for PUFA intake is necessary to confirm this.

Another interesting outcome of the present study was that there was no effect of the diet on *FFAR2*, despite there being strong evidence for an association between *FFAR2* and fat intake, particularly saturated fat, in the baseline data of the present study<sup>(13)</sup>. The previous finding was unexpected as *FFAR2* is mainly responsible for SCFA chemoreception<sup>(14)</sup>. Instead *FFAR2* might not be regulated by fat intake but rather by total energy, as high saturated fat intake (g and %) is highly correlated with increased energy and reduced dietary fibre intake in this sample (unreported). Further to this, changes in *FFAR2* was associated with changes to dietary fibre intake which is likely due to incidental changes to fibre intake as an artifact of the diets. This result is noteworthy, as dietary fibre intake leads to production of SCFA by biota in the GIT<sup>(31)</sup>. Since regulation of receptors is presumably analogous throughout the alimentary canal<sup>(8)</sup>, it is possible that increased exposure to SCFA in the GIT may cause increased *FFAR2* in TBC. Similarly, although no time–diet interaction was found for *GPR84* expression, an association was found between *GPR84* and energy from dietary fibre (%). As *GPR84* is mainly involved in the chemoreception of SCFA, this too may be due to increased exposure to SCFA in the GIT.

There was a strong time–diet interaction for FATT ( $P<0.001$ ), suggesting a physiological change to taste mechanisms was causing the change in FATT. While this is likely due to changes in FAT receptor gene expressions, namely *FFAR4*, there are many complexities to the taste system that should be discussed. First, small changes in gene expression may result in large physiological effects due to potent effects of a protein, differences in the amount of protein products and the half-life of the mRNA. To that point, we only measured gene expression in fungiform papillae as it was the least invasive tissue<sup>(21)</sup>, whereas there may have been larger changes in gene expression in foliate and circumvallate papillae<sup>(32)</sup>. Second, coordination and interaction of receptors may confound the results. For example, up-regulation of either *FFAR4* or *CD36* independently may have an insignificant effect on FATT; but when these up-regulated together, it resulted in a larger attenuation of FATT due to intracellular signal transduction<sup>(30)</sup>. Finally, there may be salivary factors that were not measured in the present study that are additionally causal to the FATT change<sup>(33,34)</sup>.

This randomised controlled trial has some limitations that should be noted when interpreting these results. First, the analysis of FAT receptor gene expressions was only conducted in fungiform papillae, as collection of foliate and circumvallate papillae in living humans is fairly invasive. The relative sensitivities and expressions of these genes within and between human taste papillae are not clear<sup>(35,36)</sup>, so the results from the present study should be interpreted only as indicative of the entire oral cavity. Second, the sample size was small and there was only one male twin pair in the sample, so there may be limited power to detect small changes in gene expression. Third, while we explored associations between nutrient intake and FAT receptor gene expressions, the present study was designed as an intervention to dietary fat intake. Changes in other nutrient intake were incidental, and therefore the observed association need to be confirmed in trials that are designed around those nutrients

specifically. Lastly, due to the small sample of twin pairs, quantitative genetic effects could not be evaluated in the present study.

In summary, *FFAR4* is the only FAT receptor on fungiform papillae associated with FAT perception in the fasted state, and both are mediated by diet concurrently. The role of *FFAR4* appears to regulate long-term diet-mediated changes in FAT perception. Increasing the expression of *FFAR4* in papillae, via diet or otherwise, could aid in reducing taste-mediated passive overconsumption of dietary fat. In addition, there is some evidence to suggest that *CD36* is involved in regulating short-term within-meal responses to oral FA exposure, and *KCNA2* may also have some role in regulating PUFA intake, though the future research should be conducted to confirm these.

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There are no conflicts of interest.

### Supplementary material

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### References

- Cartoni C, Yasumatsu K, Ohkuri T, *et al.* (2010) Taste preference for fatty acids is mediated by GPR40 and GPR120. *J Neurosci* **30**, 8376–8382.
- Costanzo A, Orellana L, Nowson C, *et al.* (2017) Fat taste sensitivity is associated with short-term and habitual fat intake. *Nutrients* **9**, 781.
- Zhang XJ, Zhou LH, Ban X, *et al.* (2010) Decreased expression of CD36 in circumvallate taste buds of high-fat diet induced obese rats. *Acta Histochem* **113**, 663–667.
- Keast RS, Azzopardi KM, Newman LP, *et al.* (2014) Impaired oral fatty acid chemoreception is associated with acute excess energy consumption. *Appetite* **80**, 1–6.
- Little TJ & Feinle-Bisset C (2011) Effects of dietary fat on appetite and energy intake in health and obesity—oral and gastrointestinal sensory contributions. *Physiol Behav* **104**, 613–620.
- Newman L, Haryono R & Keast R (2013) Functionality of fatty acid chemoreception: a potential factor in the development of obesity? *Nutrients* **5**, 1287–1300.
- Smeets AJ & Westerterp-Plantenga MS (2006) Satiety and substrate mobilization after oral fat stimulation. *Br J Nutr* **95**, 795–801.
- Stewart JE, Seimon RV, Otto B, *et al.* (2011) Marked differences in gustatory and gastrointestinal sensitivity to oleic acid between lean and obese men. *Am J Clin Nutr* **93**, 703–711.
- Wisén O, Björvell H, Cantor P, *et al.* (1992) Plasma concentrations of regulatory peptides in obesity following modified sham feeding (MSF) and a liquid test meal. *Regul Peptides* **39**, 43–54.
- Costanzo A, Nowson C, Orellana L, *et al.* (2018) Effect of dietary fat intake and genetics on fat taste sensitivity: a co-twin randomized controlled trial. *Am J Clin Nutr* **107**, 683–694.
- Newman LP, Bolhuis DP, Torres SJ, *et al.* (2016) Dietary fat restriction increases fat taste sensitivity in people with obesity. *Obesity* **24**, 328–334.
- Stewart JE & Keast RS (2012) Recent fat intake modulates fat taste sensitivity in lean and overweight subjects. *Int J Obes* **36**, 834–842.
- Liu D, Costanzo A, Evans MD, *et al.* (2018) Expression of the candidate fat taste receptors in human fungiform papillae and the association with fat taste function. *Br J Nutr* **120**, 64–73.
- Liu D, Archer N, Duesing K, *et al.* (2016) Mechanism of fat taste perception: association with diet and obesity. *Prog Lipid Res* **63**, 41–49.
- Chevrot M, Bernard A, Ancel D, *et al.* (2013) Obesity alters the gustatory perception of lipids in the mouse: plausible involvement of lingual CD36. *J Lipid Res* **54**, 2485–2494.
- Ozdener MH, Subramaniam S, Sundaresan S, *et al.* (2014) CD36-and GPR120-mediated Ca<sup>2+</sup> signaling in human taste bud cells mediates differential responses to fatty acids and is altered in obese mice. *Gastroenterology* **146**, 995–1005.
- Martin C, Passilly-Degrace P, Gaillard D, *et al.* (2011) The lipid-sensor candidates CD36 and GPR120 are differentially regulated by dietary lipids in mouse taste buds: impact on spontaneous fat preference. *PLoS ONE* **6**, e24014.
- National Health and Medical Research Council (NHMRC) (2006) *Nutrient Reference Values for Australia and New Zealand Including Recommended Dietary Intakes*. Canberra, Australia: NHMRC.
- Department of Health and Ageing (2007) *User Guide 2007 Australian National Children's Nutrition and Physical Activity Survey*. Canberra, Australia: Commonwealth Government.
- Spielman AI, Pepino MY, Feldman R, *et al.* (2010) Technique to collect fungiform (taste) papillae from human tongue. *JoVE J Vis Exp* **42**.
- Archer NS, Liu D, Shaw J, *et al.* (2016) A comparison of collection techniques for gene expression analysis of human oral taste tissue. *PLOS ONE* **11**, e0152157.
- Haryono RY, Sprajcer MA & Keast RS (2014) Measuring oral fatty acid thresholds, fat perception, fatty food liking, and papillae density in humans. *JoVE J Vis Exp* **88**.
- Ichimura A, Hirasawa A, Poulain-Godefroy O, *et al.* (2012) Dysfunction of lipid sensor GPR120 leads to obesity in both mouse and human. *Nature* **483**, 350.
- Quesada-López T, Cereijo R, Turatsinze JV, *et al.* (2016) The lipid sensor GPR120 promotes brown fat activation and FGF21 release from adipocytes. *Nat Commun* **7**, 13479.
- Talukdar S, Bae EJ, Imamura T, *et al.* (2010) GPR120 is an omega-3 fatty acid receptor mediating potent anti-inflammatory and insulin-sensitizing effects. *Cell* **142**, 687–698.





26. Camandola S & Mattson MP (2017) Toll-like receptor 4 mediates fat, sugar, and umami taste preference and food intake and body weight regulation. *Obesity* **25**, 1237–1245.
27. Laugerette F, Passilly-Degrace P, Patris B, *et al.* (2005) CD36 involvement in orosensory detection of dietary lipids, spontaneous fat preference, and digestive secretions. *J Clin Invest* **115**, 3177–3184.
28. Sclafani A, Ackroff K & Abumrad NA (2007) CD36 gene deletion reduces fat preference and intake but not post-oral fat conditioning in mice. *Am J Physiol Regul Integr Comp Physiol* **293**, R1823–R1832.
29. Braymer HD, Zachary H, Schreiber AL, *et al.* (2017) Lingual CD36 and nutritional status differentially regulate fat preference in obesity-prone and obesity-resistant rats. *Physiol Behav* **174**, 120–127.
30. Abdoul-Azize S, Selvakumar S, Sadou H, *et al.* (2014) Ca<sup>2+</sup> signaling in taste bud cells and spontaneous preference for fat: unresolved roles of CD36 and GPR120. *Biochimie* **96**, 8–13.
31. Kles KA & Chang EB (2006) Short-chain fatty acids impact on intestinal adaptation, inflammation, carcinoma, and failure. *Gastroenterology* **130**, S100–S105.
32. Gilbertson TA, Liu L, Kim I, *et al.* (2005) Fatty acid responses in taste cells from obesity-prone and-resistant rats. *Physiol Behav* **86**, 681–690.
33. Mounayar R, Septier C, Chabanet C, *et al.* (2013) Oral fat sensitivity in humans: links to saliva composition before and after stimulation by oleic acid. *Chemosens Percept* **6**, 118–126.
34. Poette J, Mekoué J, Neyraud E, *et al.* (2014) Fat sensitivity in humans: oleic acid detection threshold is linked to saliva composition and oral volume. *Flavour Frag J* **29**, 39–49.
35. Mattes RD (2009) Oral thresholds and suprathreshold intensity ratings for free fatty acids on 3 tongue sites in humans: implications for transduction mechanisms. *Chem Senses* **34**, 415–423.
36. Costanzo A (2018) The influence of diet and genes in fat taste. PhD Thesis, Deakin University.