

FOODEEG: An open dataset of human electroencephalographic and behavioural responses to food images

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Abstract

Investigating the neurocognitive mechanisms underlying food choices has the potential to advance our understanding of eating behaviour and inform health-targeted interventions and policy. Large, publicly available neural and behavioural datasets can enable new discoveries and targeted hypothesis tests, yet no such datasets are currently available. We present the FOODEEG dataset containing electroencephalographic (EEG) responses to a diverse array of food images, as well as behavioural measures of food cognition (food categorisation task, food go/no-go task, and food choice task responses), collected from 117 participants. We also provide normative ratings for the food image stimuli with respect to 22 food attributes, including nutritive, hedonic and taste properties, familiarity, and elicited emotions. Our dataset also includes questionnaire-based measures of participants' food motivations, dietary styles, and general motivational tendencies. In the validation analyses, we demonstrate that early food-evoked EEG responses in our dataset are consistent with observations in previous work. The FOODEEG dataset will be valuable for accelerating research into the neural substrates of visual food processing, dietary decisions, and individual differences.

Background & Summary

Food-related decisions are highly complex, often involving the integration of sensory and mnemonic information as well as the influence of social, cognitive, and contextual factors. Poor diet has been linked to a number of negative health consequences, including increased risks for cardiovascular and neurodegenerative diseases, and certain types of cancer^{1,2}; yet, maintaining a healthy diet is a challenge for many people. While recent neuroimaging work has shed light on the mechanisms underlying food-related cognition, such as the visual processing of foods^{3–7}, cognitive reappraisal^{8–13}, working memory^{14–16}, and reward processing^{17–19}, we do not yet fully understand how people make dietary decisions. Gaining insight into these neural processes could inform future interventions aimed at improving dietary choice, and subsequent health and wellbeing.

Using visual food cues (e.g., food images) to investigate the neural mechanisms of dietary choice confers several advantages. Compared to using real foods as stimuli, using food images allows for precise characterisation of stimulus features (e.g., size, colour), increases reproducibility across and within studies, and reduces cost. There is also evidence that responses to food images are associated with eating behaviour. A meta-analysis²⁰ reported that cue reactivity and craving elicited by visual food cues predicted eating behaviour and weight gain with a similar effect size as those elicited by exposure to real foods, and with a greater effect size than those elicited by olfactory food cues. Researchers can also readily access food image stimulus sets for use in experimental research, such as Food-pics^{21,22}, the FoodCast Research Image Database²³, the Cross-Cultural Food Image Database²⁴, the Open Library of Affective Foods^{25,26}, and the Macronutrient Picture System²⁷.

These image databases can be used to systematically measure neural responses to foods that vary across a wide range of choice-relevant dimensions. However, collecting a large amount of neuroimaging data using these stimulus sets can be time consuming and expensive. Recently, large-scale neuroimaging datasets collected while viewing scenes, objects, and animals have become publicly available (e.g., Natural Scenes Dataset²⁸; THINGS-EEG²⁹), allowing for a rapid expansion in research on the visual processing of objects and scenes. Providing large, high-quality

datasets containing neural responses to visual food cues can similarly help researchers to devise and test novel questions in food cognition research and nutrition science. Such datasets also enable replication analyses in conjunction with other neuroimaging datasets to test the generalisability of novel findings. Furthermore, large-scale behavioural and neural datasets can be used to train models of eating behaviour to predict health outcomes³⁰.

Food choice behaviour can also vary widely, and may reflect individual differences in eating styles^{31–33}. Such differences may also be reflected in neural activity. Examination of neural responses to food images could potentially reveal mechanisms that underpin individual differences in dietary choices. Differences in food-evoked neural responses has been reported for people who differ in their levels of restrained eating (the tendency to restrict food intake to control body size or weight)^{34–36}, external eating (tendency to eat in response to external food cues)^{37,38}, and emotional eating (tendency to eat in response to negative emotions)³⁹. Individuals can also vary in their motivations when making food choices, with different levels of consideration given to factors such as health, mood, pleasure, familiarity, habits, and hunger when choosing what to eat^{40,41}. However, few studies have examined the associations between food motivations and the neural processing of food cues. Another important facet of food cognition is inhibitory control. A recent meta-analysis⁴² revealed that food-related inhibitory control (as measured using laboratory-based behavioural tasks) plays a role in food consumption, though the strength of the association was small. However, few studies have included measures to examine similar differences in neural responses³⁸. Large datasets are particularly important when examining the links between neural processing of foods and individual differences in eating style, food motivations, and inhibitory control abilities, given the high power required for between-subject group comparisons or correlational analyses.

Here, we present a large dataset ($N = 117$) comprising electroencephalographic (EEG) and behavioural responses to a broad set of 120 food image stimuli. This dataset includes EEG data and behavioural responses recorded during a food categorisation task, and also behavioural data collected during a food go/no-go task and a paired food choice task. Questionnaire-based measures of participants' food

motivations (Food Choice Questionnaire⁴¹, Eating Motivation Survey⁴⁰), dietary styles (Dutch Eating Behaviour Questionnaire³³, Three Factor Eating Questionnaire-R21^{32,43}), general motivational tendencies (Behavioural Activation and Behavioural Inhibition Scales⁴⁴), and hedonism (Present Time Orientation Scale⁴⁵) are included. Additionally, we provide normative ratings on 22 food attributes for each of the food images, spanning nutritive properties (healthiness, calorie content, edibility, and level of transformation), hedonic properties (tastiness, willingness to eat, positive and negative valence, and arousal), familiarity (previous exposure, recognisability, and typicality), taste properties (sweetness, saltiness, sourness, bitterness, and savouriness), and emotional properties (happiness, surprise, disgust, craving, and guilt) collected from separate online samples (total $N = 624$).

Users may take advantage of the high temporal resolution afforded by EEG, which allows for the characterisation of rapid neural processes that underlie visual perception and decision-making. Previous work has shown that abstract information about objects beyond low-level visual features, such as animacy, naturalness, and elicited emotions, are reflected in EEG signals within the first several hundred milliseconds after image onset^{46–48}. This dataset is well suited for uncovering the precise timing of neural processes that occur in response to food images.

The EEG task was carefully designed to separately measure the appraisal and decision stages of food evaluation (Figure 1A). In this task, participants were presented with a food image and were asked to evaluate the food on one of three attributes: healthiness, tastiness, and willingness to eat. Each food image was presented three times (once for each attribute). We sought to separate the appraisal and decision stages of food evaluation by including a viewing phase, in which participants viewed the food image in isolation for 2 seconds, before the response phase, in which participants were presented with the prompt indicating the attribute they were to evaluate the food on. The mapping of the response options onto response keys was randomised across trials to prevent anticipatory motor preparation. We reduced priming effects by presenting healthiness, tastiness, and willingness to eat trials in a randomised order within blocks, rather than presenting each trial type in separate blocks. This ensured that the participants were not able to anticipate which of the three attributes would be judged

during food image viewing. This task design allows for investigations into the neural mechanisms underlying food appraisal and the visual processing of foods while minimising the influences of neural signals related to decision or motor preparation. Users interested in the neural correlates of food-related decision-making processes can instead examine relationships between choice behaviour (e.g., response times) and neural activity in the response phase.

The food image stimuli were carefully curated to ensure that there was substantial variance in key food attributes. This is often difficult to achieve because people tend to view most foods as appetising and positively valenced. Ninety-one images were selected from an established food image database (Food-pics²²) and consisted of mostly palatable and familiar foods. Twenty-nine images of foods that are less palatable or familiar, but still edible, were sourced from online searches and included in the stimulus set. Some existing work has compared behavioural responses towards edible versus inedible foods (e.g., rotten or decayed foods^{49,50}), but few studies have included foods that are unappetising but not harmful to eat in their stimuli.

Rating data on a wide range of food attributes indicate that our stimuli sufficiently varied on key attributes that are relevant for dietary choice. This indicates that our dataset is suitable for analyses that are sensitive to fine-grained differences in the variables of interest, such as multivariate pattern analysis (MVPA) or representational similarity analysis (RSA). For example, MVPA can be applied to investigate how fine-grained rating information about a food attribute of interest covaries with patterns of EEG data, to delineate the precise neural time-courses of these representations during food viewing (see Technical Validation for an example). Alternatively, fine-grained rating information can be used to construct models of similarity for a given food attribute and compare them to models of similarity for neural or behavioural data. Recent work has demonstrated how RSA can be applied to this dataset.³

Beyond what we provide here, additional ratings can be easily collected from new online participant samples to assess an even wider range of food attributes and how they covary with neural responses in our dataset. Users may also use these

normative ratings in combination with the food images to inform stimuli selection, to ensure a particular distribution of attributes to suit their research goals.

In sum, the FOODEEG dataset comprises of a rich array of neural and behavioural measures during food cognition tasks collected from a large sample ($N = 117$). Given the multifaceted nature of eating behaviour, we have included measures of eating styles, food motivation, and relevant personality traits which may be useful in controlling for, or investigating, individual differences. We also provide a comprehensive set of ratings on 22 food attributes that encompass nutritive, hedonic, familiarity, taste, and emotional properties of foods. The FOODEEG dataset has the potential to advance food cognition research by supporting a wide range of research projects, including investigations linking neurocognitive mechanisms to individual differences, food attribute processing, and the dynamics of choice behaviour.

Methods

Neural and behavioural data were collected in-person over two testing sessions. In Session 1, participants completed a food categorisation task (Figure 1B) while EEG was recorded. They subsequently provided healthiness, tastiness, and willingness to eat ratings for each food image. In Session 2, participants completed a food go/no-go task (Figure 1C) and a paired food choice task (Figure 1D). They also completed questionnaires relating to their dietary style, food motivations, and general motivational tendencies. A separate sample of participants completed an online food rating task, in which participants were asked to rate the food image stimuli on multiple food attributes. Figure 1A shows the overview of the data collection procedure. This study was approved by the Human Research Ethics Committee of the University of Melbourne (HREC ID 24850).

Participants

For the in-person sessions, participants ($N = 117$) were recruited through the University of Melbourne student portal and advertising via posters around the University of Melbourne. Participants indicated that they were at least 18 years old, were fluent in written and spoken English, did not have a history of any eating disorders, were not on a calorie restriction diet, and did not have any dietary

restrictions (e.g., for health or religious reasons). Participants were asked to fast for three hours prior to the start of both in-person sessions. At the end of the second session, participants were reimbursed AUD \$50. The sample had a mean age of 26.8 years ($SD = 8.50$, range 18-57 years). 84 participants identified as female, 32 as male, and one as other.

A separate sample of online participants ($N = 624$) were recruited through the University of Melbourne Research Experience Program. Participants indicated that they were at least 18 years old and fluent in written and spoken English. Participants indicated that they did not have a history of any eating disorders, were not on a calorie restriction diet, did not have any dietary restrictions, and had not previously taken part in the in-person testing sessions. The online sample had a mean age of 19.55 years ($SD = 2.92$, range 18-53 years). 482 participants identified as female, 132 as male, 7 as non-binary/other, and 3 preferred not to disclose. Participants were awarded course credit at the conclusion of the experiment.

Stimuli

The stimuli consisted of 120 colour images of food items against a white background. Ninety-one images were drawn from the Food-pics database²² and 29 images were drawn from online search. Food-pics is a database of food images containing images of a wide variety of food items, including fruit and vegetables, snack foods, meat, and sweets, designed for use in research on eating behaviour. The 91 food images were selected to include a wide range of food types (e.g., vegetables, fruits, savoury dishes, sweet dishes, meat). Food images that portrayed multiple types of foods or included cutlery were not included. An additional 29 images were taken from online image hosting websites (e.g., Flickr, Adobe Stock). We aimed to increase variance in key food attributes (e.g., tastiness, valence, familiarity) in our stimuli set. To this end, we selected foods that were generally less appetising or less familiar but were nevertheless edible. Examples of the food images are displayed in Figure 1E.

Procedure

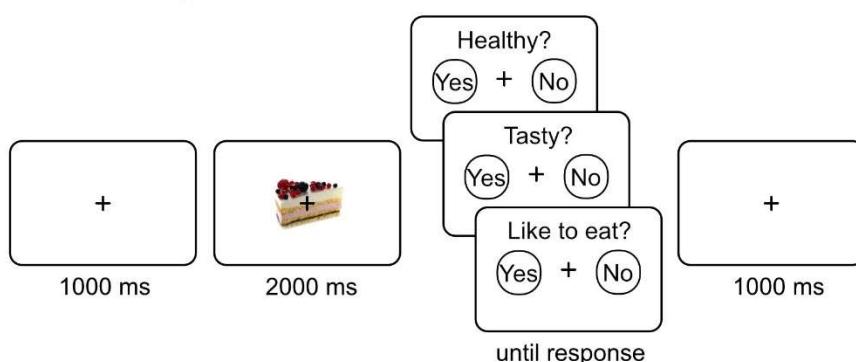
Participants attended the first and second in-person sessions on different

days, approximately 1 week apart. During the first session, participants completed a food categorisation task presented using Psychtoolbox⁵¹ interfacing Matlab R2022a. Afterwards, participants rated each food image on healthiness, tastiness and willingness to eat using Qualtrics. During the second session, participants completed two behavioural tasks: a food go/no-go task and a paired food choice task presented using PsychoPy⁵² interfacing Python (v3.11). All experimental code will be available via the Open Science Framework (<https://doi.org/10.17605/OSF.IO/Y9PMF>) at the time of publication. Participants also completed questionnaires relating to dietary styles, food motivations, and general motivational tendencies using Qualtrics.

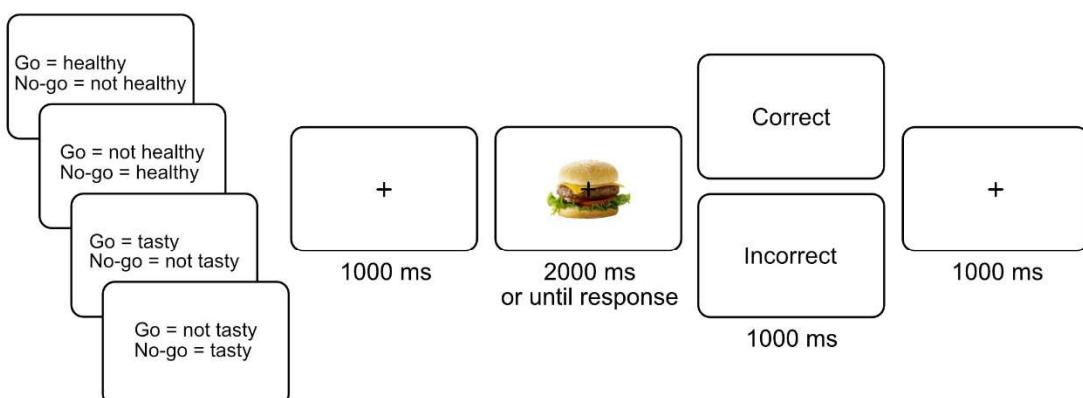
A: Data Collection Overview



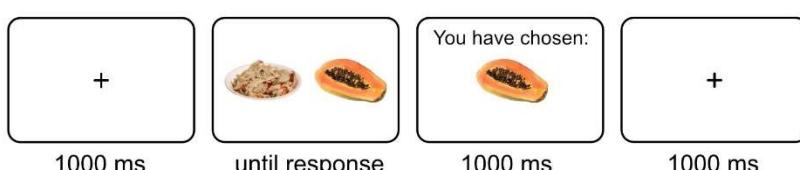
B: Food Categorisation Task



C: Food Go/No-go Task



D: Paired Food Choice Task



E: Example Food Image Stimuli



Figure 1. Overview of Data Collection and Experimental Trial Diagrams. A) Data collection overview. The data was collected over two in-person sessions and online. In the first in-person session, participants completed the food categorisation task (while EEG was recorded) and provided ratings on the food images for three attributes. In the second in-person session, participants completed two behavioural tasks (food go/no-go task and paired food choice task) and filled out survey measures of dietary styles, food motivations, and general motivational tendencies. In an online experiment using a separate sample, attribute ratings were collected for the food images for 22 choice-relevant food attributes. B) Food categorisation task. For each trial, participants viewed a food image (2 s), followed by a response screen with one of three prompts probing healthiness, tastiness, or willingness to eat and one of two keyboard press response mappings ('z' key for Yes and 'm' key for No, or the reverse). Participants were instructed to respond as quickly as possible. C) Food go/no-go task. In each trial, participants were presented with a food image and either responded as quickly as possible with a keyboard press for Go foods or withheld any responses (2 s) for the No-go foods. Feedback ('Correct' or 'Incorrect') was given after each trial. Participants completed four blocks with different Go and No-go foods. D) Paired food choice task. Participants were presented with two food images and were asked to choose the food that they would prefer to eat more, followed by a feedback screen showing the food image that they had selected. E) Example food image stimuli. Food images were selected to include a wide variety of food types and also included foods generally thought to be less appetising or familiar.

Food categorisation task with EEG recording. During the food categorisation task, participants viewed each food item three times: once each across the healthiness, tastiness, and willingness to eat trials while EEG was recorded (Figure 1A). Food images were presented centrally, subtending 6.6×3.5 degrees of visual angle on a 24.5-inch ASUS ROG Swift PG258Q monitor (1920 x 1080 pixels, 60 Hz refresh rate). Participants' heads were stabilized using a chin and forehead rest approximately 50 cm from the monitor. Each trial began with a fixation cross presented in isolation for 1 s, followed by the food image with a fixation cross for 2 s. Next, the participants were prompted with "Healthy?" in a healthiness trial, "Tasty?"

in a tastiness trial, and “Like to eat?” in a willingness to eat trial, along with a response mapping that indicated which of the left (‘z’) and the right (‘m’) keyboard keys corresponded to ‘Yes’ and ‘No’ responses. There was an equal chance of the left keyboard press indicating a ‘Yes’ response and the right keyboard press indicating a ‘No’ response, or the reverse left/right mapping, for each trial. For each participant, the order of healthiness, tastiness, and willingness to eat trials were randomised. The randomisation of the trial type order and response mapping ensured that participants could not pre-emptively prepare a keypress motor action during the food image presentation period. Participants were asked to respond as quickly and accurately as possible and to pay attention to the switching of the response mapping across trials. Participants completed five practice trials and indicated that they had understood the instructions before starting the main experiment. To reduce eye movement artifacts, participants were instructed to keep their eyes fixated on the central fixation cross, which remained on screen throughout the duration of the experimental blocks. Response time was recorded as the time taken from the presentation of the prompt to the participant’s keypress response. Participants completed six blocks of 60 trials each.

EEG was recorded using a Biosemi Active II system (Biosemi, The Netherlands) with 64 scalp electrodes at a sampling rate of 512 Hz using common mode sense and driven right leg electrodes (<http://www.biosemi.com/faq/cms&drl.htm>). We attached eight additional electrodes: two electrodes placed 1 cm from the outer canthi of each eye, four electrodes placed above and below the centre of each eye, and two electrodes placed above the left and right mastoids. Electrode offsets were kept within the range of $\pm 20 \mu\text{V}$ during recording.

Health, taste, and willingness to eat ratings. After the food categorisation task, participants rated each food image on healthiness (“How healthy is this food?”), tastiness (“How tasty is this food?”) and how willing they were to eat it (“How much would you like to eat this food?”). Participants were asked to use a computer mouse to indicate their response on a continuous rating scale ranging from ‘Not healthy at all’ to ‘Very healthy’ for the healthiness rating, ‘Not tasty at all’ to ‘Very tasty’ for the tastiness rating, and ‘Not at all’ to ‘Very much’ for the willingness to eat rating. We randomised image presentation order for each participant, while ensuring that the same food

image was not presented across two consecutive trials.

Food go/no-go task. Adapted from the standard go/no-go paradigm, the food-specific go/no-go task has been used in previous research to measure food-related inhibitory abilities^{53,54}. Stimulus set selection was carried out for each individual participant, using their attribute ratings from Session 1. Healthiness and tastiness ratings from Session 1 were used to select 60 food image stimuli across four categories: 15 Most-Healthy foods, 15 Least-Healthy foods, 15 Most-Tasty foods, and 15 Least-Tasty foods. To select the Most-Tasty foods, 15 foods with the highest tastiness ratings were selected. To select Least-Tasty foods, 15 foods with the lowest tastiness ratings were selected. From the remaining foods, Most-Healthy and Least-Healthy foods were selected in the same way. If there were more than 15 foods with the highest or lowest ratings for an attribute, 15 were randomly selected from those foods. No foods were included in multiple categories. Prior to the start of the task, participants were shown all 60 food images and the category each belonged to. It was indicated to the participants that the foods were categorised according to their own ratings from the previous session, and they were asked to let the experimenter know if they disagreed with the categorisations for any of the foods. If the participant disagreed with the categorisation of a food, the food was removed from the category and replaced with the food with the highest (for Most-Healthy and Most-Tasty categories) or the lowest (for Least-Healthy and Least-Tasty) rating on the appropriate attribute out of the unselected foods.

Participants completed four blocks of the go/no-go task. In the first block, Most-Healthy foods served as the Go stimuli and the Least-Healthy foods served as the No-go stimuli. In the second block, Most-Tasty foods served as the Go stimuli and the Least-Tasty foods served as the No-go stimuli. The Go and No-go stimuli in the first and second blocks were switched for the third and fourth blocks respectively. There were 100 trials in each block. In each block, Go foods were presented in 75% of the trials and No-go foods were presented in 25% of the trials. At the start of each block, instruction text indicated to the participants which food categories served as the Go and No-go stimuli for the block. Next, the participants were shown a series of food images and were asked to make a response via keyboard press as quickly as possible

to the Go foods and to withhold the response to the No-go foods. A fixation cross was presented for 1 s between trials. Block order was randomised for each participant.

Paired food choice task. To assess people's preferences across foods that differed in attributes (e.g., healthiness and tastiness), participants selected which food they would prefer to eat more out of pairs of foods they had previously indicated they would be willing to eat. Stimuli selection was carried out for each individual participant, using their attribute ratings from Session 1. Out of the food images that had been rated over 50 (scale: 0-100) on willingness to eat in Session 1, 25 food images were selected. If less than 25 food images had been rated over 50 on willingness to eat, the subsequent highest rated food images were included in the stimuli to bring the total number of food images to 25.

Participants completed five blocks of the paired food choice task. There were 60 trials in each block. In each trial, participants were presented with two food images on the left and right side of the screen. Participants indicated which food they would prefer to eat more with a keyboard press ('z' to select the left food and 'm' to select the right food). Every possible pairwise combination of the 25 food images was presented once, totalling 300 trials. A fixation cross was presented for 1 s between each trial.

Questionnaires. Participants' dietary styles, food motivations, general motivational tendencies, and hedonism were measured using self-report scales. Dietary styles were assessed using the Dutch Eating Behaviour Questionnaire³³ and the Three Factor Eating Questionnaire Revised 21-Item^{32,43}. Food-related motivations were assessed using the relevant subscales of the Food Choice Questionnaire (Health, Mood, Sensory Appeal, Natural Content, Weight Control, and Familiarity subscales)⁴¹ and the Eating Motivation Survey Brief (Liking, Habits, Need and Hunger, Health, Pleasure, Natural Concerns, Weight Control, Affect Regulation subscales)⁴⁰. Motivational tendencies were measured using the Behavioural Activation and Behavioural Inhibition Scales⁴⁴ and hedonism was assessed using the Present Time Orientation Scale⁴⁵. The descriptive statistics for the survey measures are presented in Table 1.

Table 1. Summary statistics for survey measures of dietary style, food motivations, general motivational tendencies, and hedonism. Mean, standard deviation, and range of individual participant's mean scores for the subscales of Food Choice Questionnaire, the Eating Motivations Survey – Brief, Three Factor Eating Questionnaire, Dutch Eating Behaviour Questionnaire, Behavioural Inhibition System/Behavioural Activation System Scale, and Present Time Orientation Scale are reported.

Scale	Subscale	Mean	SD	Min	Max
Food Choice Questionnaire	Health	3.4	0.77	1.17	5
	Mood	3.55	0.77	1.5	5
	Sensory appeal	3.81	0.67	2	5
	Natural content	2.77	1.08	1	5
	Weight control	2.56	0.99	1	5
	Familiarity	2.64	0.94	1	5
The Eating Motivations Survey - Brief	Liking	5.93	0.73	4.33	7
	Habits	4.63	1.22	1	7
	Need and hunger	5.42	0.93	3	7
	Health	4.61	1.06	2	7
	Pleasure	5.16	0.98	2.67	7
	Natural concerns	3.75	1.48	1	7
	Weight control	3.23	1.34	1	6.33
	Affect regulation	2.95	1.44	1	6.67
Three Factor Eating Questionnaire	Cognitive restraint	2.24	0.6	1	3.67
	Emotional eating	2.07	0.71	1	4
	Uncontrolled eating	2.21	0.56	1.11	3.78
Dutch Eating Behaviour Questionnaire	Restrained	2.63	0.85	1	4.75
	Emotional	2.42	0.89	1	4.83
	External	3.36	0.67	1.8	5
Behavioural Inhibition System/Behavioural Activation System Scale	Punishment				
	Sensitivity	1.84	0.49	1	3.29
	Drive	2.16	0.53	1	3.75
	Reward				
	responsiveness	1.5	0.4	1	2.6
	Fun seeking	2.01	0.61	1	3.75
Present Time Orientation Scale	Hedonism	3.4	0.62	1.67	4.67

Online food attribute ratings. A separate sample of participants were recruited for the online rating task. Participants indicated their current hunger ("How hungry are you currently?") and satiety ("How full are you currently?") level on a sliding scale from 'Not at all' to 'Very much'. They selected the time elapsed since their last food

consumption (“How long ago did you last eat food?”) from one of the following options: ‘Less than an hour ago’, ‘1-3 hours ago’, ‘3+ hours ago’, and ‘I do not know’. Each participant rated a subset of the full stimulus set (40 out of the 120 images) on 12 (out of 22) food attributes. We randomised the image presentation order for each participant as well as the order of the food attributes for each image. Table 2 displays the full list of food attributes and the corresponding prompts and additional information presented to the participants. Participants were asked to indicate their response to each prompt on a sliding scale (0-100) using a computer mouse.

Table 2. Food attribute ratings and corresponding prompts. Ratings on 22 food attributes across five categories (nutritive, hedonic, familiarity, taste, and emotional properties) were collected for the food image stimuli using the corresponding prompt for each food image. For edibility, level of transformation, and arousal, additional information was displayed alongside the prompt (in italics).

	Food attribute	Prompt
Nutritive properties	Healthiness	How healthy is the food represented in the image?
	Calorie content	How much calorie content are in the food represented in the image?
	Edibility	How much work WILL BE required to bring the food represented in the image ready to eat? <i>For example, raw chicken will require more work (i.e., more steps, long time, many ingredients) to get it ready to eat compared to an apple (i.e., less steps, short time, few ingredients).</i>
	Level of transformation	How much work WAS required to prepare the food represented in the image? <i>For example, a cake required more work (i.e., more steps, long time, many ingredients) to prepare than rice (i.e., less steps, short time, few ingredients).</i>
Hedonic properties	Tastiness	How tasty is the food represented in the image?
	Willingness to eat	How much would you like to eat the food represented in the image?
	Negative valence	How negatively do you view the food represented in the image?
	Positive valence	How positively do you view the food represented in the image?

	Arousal	How emotionally arousing is the food represented in the image? <i>Less emotionally arousing foods are calming/relaxing, and more emotionally arousing foods are stimulating/energising.</i>
Familiarity	Exposure	How often do you encounter the food represented in the image?
	Recognisability	How easy is it to identify the food represented in the image?
	Typicality	How typical is the food represented in the image?
Taste properties	Sweet	How much of the following taste is elicited by the food represented in the image?
	Salty	
	Bitter	
	Sour	
	Savoury (umami)	
Emotional properties	Happiness	How much of the following emotion is elicited by the food represented in the image?
	Disgust	
	Surprise	
	Craving	
	Guilt	

Data Records

All data and code will be publicly available at the time of publication. The raw and processed EEG recordings and behavioural data will be hosted in Brain Imaging Data Structure (BIDS) standard^{55,56} on OpenNeuro (<https://doi:10.18112/openneuro.ds006832.v1.0.0>). Custom code, food image stimuli, survey data, and normative ratings will be available from the Open Science Framework (<https://doi.org/10.17605/OSF.IO/Y9PMF>).

Technical Validation

EEG data validation

For technical validation of the current dataset, we used two complementary analyses to demonstrate that both binary categorisations and continuous ratings on food attributes covary with patterns of EEG data. We examined EEG responses following food image onset to assess whether event-related potential (ERP) waveforms differed across different food

items that categorically varied in attributes (e.g., Healthy versus Not-Healthy foods).

Furthermore, we used MVPA to demonstrate that fine-grained information about food attributes is also present in the EEG signals.

Data processing. EEG data were processed using EEGLab v2022.1⁵⁷ interfacing Matlab (R2022a). Excessively noisy channels were identified through visual inspection (mean number of bad channels = 0.26, range 0–5) and excluded from average reference calculations and independent components analysis (ICA). Excessively noisy sections were identified through visual inspection and removed. Five participants were excluded from the analyses due to noisy data or missing trials. Two additional participants were excluded due to high incongruence between their responses in the food categorisation task and their subsequent continuous food attribute ratings. The data were referenced to the average of all channels (excluding excessively noisy channels). One channel (AFz) was removed to compensate for the rank deficiency due to average referencing. EEG data were low-pass filtered at 30 Hz (EEGLab Basic Finite Impulse Response Filter New, default settings). A copy of the dataset was created for the purpose of ICA. A 0.1 Hz high-pass filter (EEGLab Basic Finite Impulse Response Filter New, default settings) was applied to the copied dataset to improve stationarity. We used RunICA extended algorithm⁵⁸ to perform the ICA. The resulting independent component information was copied to the original dataset⁵⁹. Independent components associated with eye blinks and saccades were identified and removed in line existing guidelines⁶⁰. Next, we interpolated the excessively noisy channels and AFz using spherical spline interpolation. EEG data were segmented from -100 ms to 1000 ms relative to food image onset. Segments were baseline-corrected using the data from the 100 ms time window prior to each food image presentation. Segments from trials where the participant took longer than 5 s to respond or those containing amplitudes exceeding $\pm 150 \mu\text{V}$ at any channel were excluded from the analyses. On average, 346 out of 360 trials were retained (range 283 – 360).

Event-related potentials. We demonstrate that ERP waveforms closely resemble those observed in previous work using visual food stimuli^{6,10,61–63}, and that differences in ERPs can be observed across food images that vary categorically in attributes of interest (e.g., Healthy versus Not-Healthy, Tasty versus Not-Tasty foods).

Figure 2 shows group-average ERPs measured at occipital electrodes (Oz, O2, O1, and Iz) and at parietal electrodes (Pz, P1, P2, CPz, and POz) following food image onset. For each participant, we split the trials according to the participants' responses during the food categorisation task on healthiness and tastiness trials (Healthy versus Not-Healthy, Tasty versus Not-Tasty). We then averaged EEG responses across trials to derive ERPs for each choice condition. ERPs were averaged over electrodes within each region of interest (occipital and parietal) to further improve the signal-to-noise ratio. We compared ERPs following Tasty and Not-Tasty food images, and Healthy and Not-Healthy food images, using cluster-based permutation tests to correct for multiple comparisons⁶⁴ (10,000 permutation samples, cluster forming alpha = .01, family-wise alpha = .05) using functions from the Decision Decoding Toolbox v1.1.5⁶⁵.

We observed typical patterns of visual evoked ERPs at occipital channels, including the visual P1, N1 and P2 components that occur within the first 250 ms following food image presentation^{10,61,62}. We also observed a later, sustained positivity at parietal channels, resembling the P3 or Late Positive Potential that has been measured in prior work^{6,61,62,66}. At occipital electrodes, we observed more negative-going ERP waveforms both for tasty and healthy foods between approximately 150-600 ms (Figure 2A, 2D). At parietal channels, amplitudes were more positive-going between approximately 300-550 ms following Tasty compared to Not-Tasty foods (Figure 2E) as also reported in prior work^{6,61,62}. This demonstrates that EEG responses evoked by food images do broadly differ across food attribute decision outcomes, and that it could be used to test more targeted hypotheses relating to food-evoked ERPs.

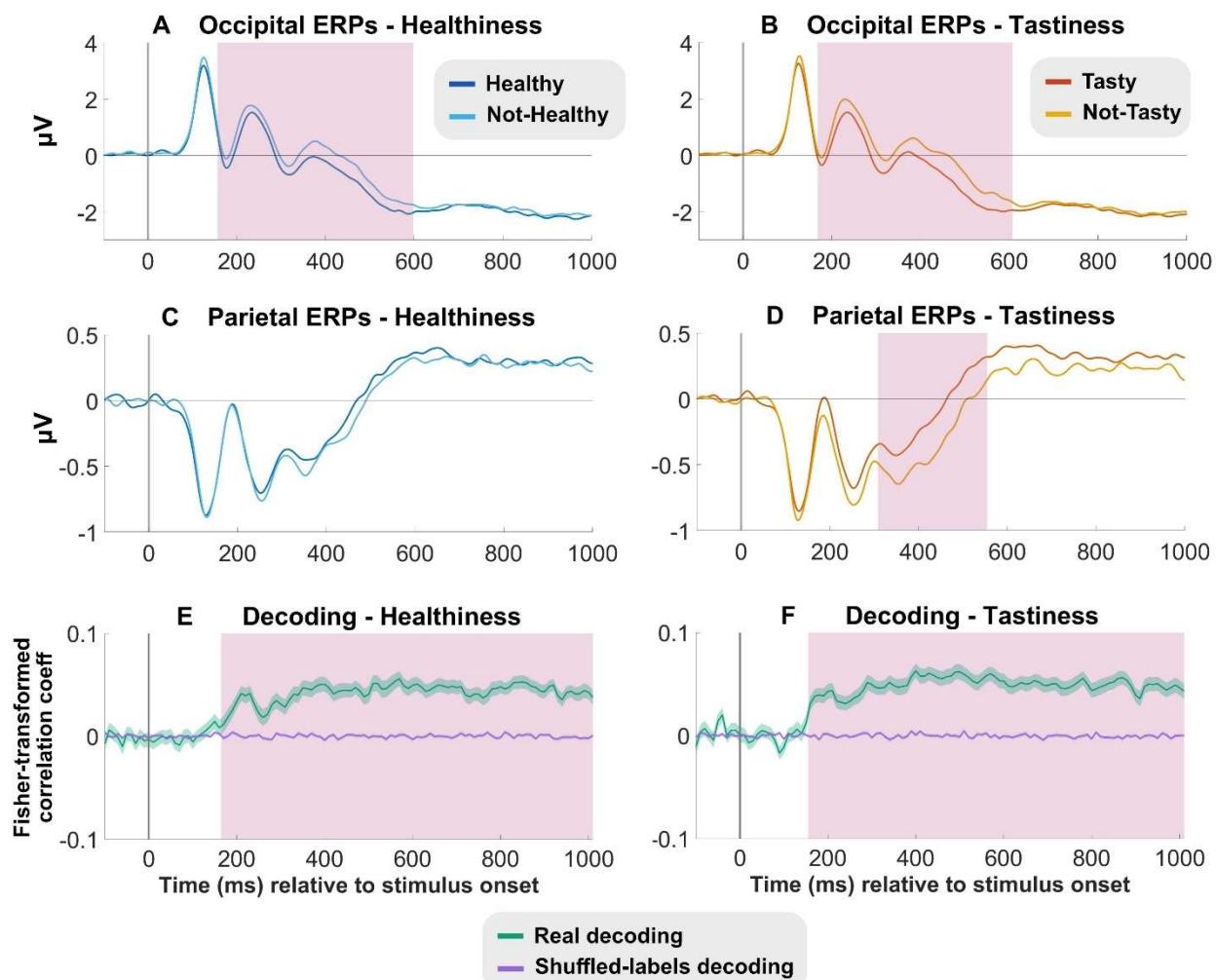


Figure 2. Group-average ERPs and decoding results following food image onset.

The top row shows ERPs evoked at occipital (Oz, O2, O1, and Iz) electrodes by A) Healthy (dark blue) and Not-Healthy (light blue) foods and by B) Tasty (red) and Not-Tasty (yellow) foods. The second row displays ERPs evoked at parietal (Pz, P1, P2, CPz, and POz) electrodes by C) Healthy and Not-Healthy foods and by D) Tasty and Not-Tasty foods. For the ERPs, pink shaded areas indicate the time windows at which the mean amplitudes between conditions significantly differed after applying multiple comparisons corrections. The third row shows the decoding results. Using support vector regressions, E) healthiness and F) tastiness ratings were predicted from the EEG data from the time relative to food image onset. Green lines indicate the real decoding results (Fisher-transformed correlation coefficients between predicted and true attribute ratings) and purple lines indicate the shuffled-labels decoding results (empirical chance distribution). Shaded areas around the green and purple lines indicate the standard error of the mean. For the decoding results, pink shaded areas

indicate the time windows at which the real decoding results significantly differed from the shuffled-labels decoding results and after applying multiple comparisons corrections.

Multivariate pattern analysis. We demonstrate using MVPA that fine-grained food attribute information beyond binary categorisations is also decodable from the current dataset. Multivariate analysis methods (e.g., MVPA, representational similarity analysis) are becoming more widely used in neuroimaging studies, due to high sensitivity that allows for the detection of distributed patterns of neural activity. Previous work has shown that such methods are useful for investigating the fine-grained neural time-courses of food attribute processing³⁻⁵. The MVPA results suggest that there are patterns in the EEG data that covaried with continuous food attribute ratings and reveal that the current dataset is suitable for multivariate analysis techniques.

Linear support vector regression (SVR) models were implemented using the Decision Decoding Toolbox v1.0.5⁶⁵ to examine whether continuous food attribute ratings were decodable from the EEG signals during food viewing. A “moving window” approach was implemented to divide each epoch of the EEG data into 10ms segments, which served as analysis time windows. These were moved through the data in steps to derive a fine-grained time-course of decoding performance. The regression model was trained to predict the continuous ratings (ranging from 0-100) for a given food attribute from the spatial brain activity patterns in each time window following the food image onset. A linear SVR model (cost parameter C=0.1, interfacing LIBSVM⁶⁷) was trained on 90% of the data, then tested on the held-out 10% of the data for each time window and for each participant. We repeated this process independently using a 10-fold cross-validation procedure so that all data sets had been used as test data once while training on all other datasets. We repeated this cross-validation process 10 times using newly selected random data to derive a conservative estimate of decoding performance for the time window, calculated as the correlation coefficient between the actual and predicted continuous ratings for the food images, averaged across the 10 x 10 iterations and Fisher-Z transformed. Results for each time window were obtained independently of any other time windows. We used cluster-based permutation tests as described above to correct for multiple tests.

Continuous healthiness and tastiness ratings were decodable from the EEG data (Figure 2C, 2F) from around 150 ms after food image onset, and remained significant until the end of the 1 s analysis window. These findings suggest that there are patterns of activity in the EEG signals that contain fine-grained information about healthiness and tastiness of foods from around 150 ms onwards. Together with the ERP analyses, they show that the FOODEEG dataset is suitable for tracking the first 1000 milliseconds of neural responses to visual food cues.

Code Availability

All custom code used to produce the technical validation analyses and figures in the manuscript will be available from the Open Science Framework (<https://doi.org/10.17605/OSF.IO/Y9PMF>).

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