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Nicolas Coquery, Sophie Menneson, Paul Meurice, Regis Janvier, Pierre Etienne, et al.. fMRI-Based Brain Responses to Olfactory Stimulation with Two Putatively Orexigenic Functional Food Ingredients at Two Different Concentrations in the Pig Model. *Journal of Food Science*, 2019, 84 (9), pp.2666-2673. 10.1111/1750-3841.14772 . hal-02281404

HAL Id: hal-02281404

<https://univ-rennes.hal.science/hal-02281404>

Submitted on 26 Nov 2019

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fMRI-based brain responses to olfactory stimulation with two putatively orexigenic functional food ingredients at two different concentrations in the pig model

Nicolas Coquery¹, Sophie Mennesson^{1,2}, Paul Meurice¹, Régis Janvier¹, Pierre Etienne², Virginie Noirots², David Val-Laillet¹

¹ INRA, INSERM, Univ Rennes, Nutrition Metabolisms and Cancer, NuMeCan, Rennes, France

² Phodé, Terssac, France

Corresponding author:

Nicolas Coquery, INRA UMR1341 NuMeCan, 35590 St Gilles, France

nicolas.coquery@inra.fr

Running title: fMRI responses to olfactory stimulation in pig

Abstract

Natural plant extracts are increasingly used as functional feed ingredients in animal husbandry and food ingredients in human alternative medicine to improve welfare and health. We investigated in 20 growing pigs *via* fMRI the brain Blood Oxygen Level-Dependent (BOLD) responses to olfactory stimulation with two sensory functional feed ingredients, A and B, at two different concentrations. Functional ingredient A contained extracts from *Citrus sinensis* (60-80%), and ingredient B contained a mixture of extracts *Oreganum vulgare* (40-55%) and *Cymbopogon flexuosus* (20-25%). Increased concentration of ingredients induced a higher activation in reward and cognitive areas compared to lower concentrations. Moreover, considering both ingredients at the highest concentration, the ingredient A elicited higher brain responses in brain areas involved in hedonism/pleasantness compared to ingredient B, and more specifically in the caudate nucleus and orbitofrontal cortex. Our findings shed new light in the scope of emotion regulation through olfactory modulation *via* sensory functional ingredients, which opens the way to further preclinical studies in animal models and translational research in the context of nutrition, welfare, and health.

Key words

Olfaction; Functional food ingredients; Pig; Brain; fMRI; Hedonism.

Practical Application

Functional food/feed ingredients are gaining interest for improving health and welfare in humans and animals. Besides representing an alternative to antibiotics for example, food ingredients and their sensory characteristics might have a positive impact on emotions and consequently on well-being. Functional brain imaging in large animals such as in the pig model is a promising approach to investigate the central and behavioural effects of food

ingredients, and determine the most effective blends and concentrations to modulate internal and emotional states.

Accepted manuscript

1 Introduction

The use of natural plant extracts as functional food ingredients is gaining interest due to both their impacts on welfare and health security (Piątkowska & Rusiecka-Ziółkowska, 2016), *e.g.* as an alternative to antibiotics in animal production, and their potential to prevent chronic diseases in humans (van Breda & de Kok, 2018). Besides an obvious post-ingestive impact on microbiota (Cairo et al., 2018; Dudek-Wicher, Junka, & Bartoszewicz, 2018), intestinal physiology (Patra, Amasheh, & Aschenbach, 2018), hormonal regulation (Bower, Real Hernandez, Berhow, & de Mejia, 2014), and immunity (Williams et al., 2017), food ingredients from natural plant extracts might also act directly on exteroception, and notably olfaction. Olfactory stimulations can elicit specific brain responses related to emotions and hedonic valuation (Billot et al., 2017; Sorokowska et al., 2017; Soudry, Lemogne, Malinvaud, Consoli, & Bonfils, 2011).

In a previous series of studies using the pig model, we investigated 8 different sensory functional feed ingredients, which induced different outcomes in terms of feed preferences and eating behavior (Clouard & Val-Laillet, 2014). Based on these behavioral data, we further investigated the impact of one of these ingredients (mainly composed of *Citrus sinensis* extracts) on brain glucose metabolism *via* ^{18}F -fluoro-deoxy-glucose (FDG) positron emission tomography (PET), and in response to olfactogustatory response with or without prior familiarization to the ingredient (Val-Laillet, Meurice, & Clouard, 2016). Because the FDG-PET imaging modality can test only one condition per imaging session, we developed a functional magnetic resonance imaging (fMRI) approach in the pig model to discriminate brain responses to different gustatory stimulations (Coquery et al., 2018). Here, we adapted this set-up to olfactory stimulation to evaluate brain responses to the odor of two feed ingredients at different concentrations within the same imaging session.

Among natural product-based extracts, fruit extracts, such as *Citrus sinensis*, present some promising properties for food intake regulation and mood modulation (Dosoky & Setzer, 2018; Igarashi, Ikei, Song, & Miyazaki, 2014; Lampion et al., 2016). Herbal extracts also present some interesting features. *Oreganum vulgare* and its principal compounds, carvacrol, as well as *Cymbopogon flexuosus*, *i.e.* lemongrass, can have positive effects on behavior and emotions (Amiresmaeili, Roohollahi, Mostafavi, & Askari, 2018; Blanco, Costa, Freire, Santos, & Costa, 2009; Costa et al., 2011; Jugl-Chizzola et al., 2006; Mehan et al., 2011; Pires et al., 2013; Sharifi-Rad et al., 2018; Zotti et al., 2013). In this study, we compared two food ingredients with putative orexigenic and hedonic outcomes: ingredient A, derived from *Citrus sinensis* extracts, and ingredient B, a combination of *Oreganum vulgare* and *Cymbopogon flexuosus* extracts. First, given that the hedonic value of an odor can be related to the concentration used (Clouard, Jouhannau, Meunier-Salaün, Malbert, & Val-Laillet, 2012), we analyzed the impacts of two different concentrations of each ingredient on brain responses to identify the best concentration for comparing ingredients. We also aimed at identifying the ingredient that promoted the highest outcomes on the reward circuit. Second, we characterized the specific neuronal networks modulated by each ingredient and the associated neurocognitive functions involved based on the scientific literature.

2 Materials and Methods

2.1 Animals

Experiments were conducted in accordance with the current ethical standards of the European Community (Directive 2010/63/EU), Agreement No. C35-275-32 and Authorization No. 35-88. The Regional Ethics Committee in Animal Experiment of Brittany has validated and approved the entire procedure described in this paper (project N° 2017070518585877). A total of twenty eighteen-week-old Piétrain x (Large White/Landrace) female pigs were used in

this study investigating the brain response to food ingredients. Before imaging, these pigs were used initially in a research project investigating the gut-brain adaptations to a six-week psychosocial stress to describe a new relevant preclinical model, and were separated into two groups. These two groups (10 animals per group) differed only in their housing conditions: either housed alone in individual pens ($150 \times 60 \times 80 \text{ cm}^3$) without environmental enrichment, or by two animals in twice larger pens ($150 \times 120 \times 80 \text{ cm}^3$) with environmental enrichments (a balloon and a suspended chain). Animals housed in individual pens were also exposed to unpredictable environmental visual and auditory stimuli (noise and rotating red light warning during day, every ten minutes, and night, every sixty minutes). For the statistical analysis, group attribution was included as a covariable. For all animals, the room was maintained at $\sim 24^\circ\text{C}$ with a natural light–dark cycle (from April to June in French Brittany).

2.2 Anesthesia

For brain imaging, pre-anesthesia was performed with an intramuscular injection of ketamine (5 mg/kg – Imalgene 1000, Merial, Lyon, France) in overnight-fasted animals, *i.e.* imaging being performed 24 to 30 hours after the last meal distribution. Isoflurane inhalation (Aerane 100 ml, Baxter SAS, Guyancourt, France) was used to suppress the pharyngotracheal reflex and then establish a surgical level of anesthesia, 3-5 % and 2-3 % respectively. After intubation, anesthesia was maintained with 2.5-3 % isoflurane and mechanical ventilation allowed adjustment of respiratory frequency at 17 breaths/minute with a tidal volume of 650 ml. Cotton wool with an additional headset was used to conceal the animal's ears and tape was used to keep the eyes closed.

2.3 Olfactory stimulation

We used an improved custom-made stimulation apparatus quite similar to that used in previous studies (Clouard, Jouhanneau, Meunier-Salaün, Malbert, & Val-Laillet, 2012), located outside the magnet-shielded room (5-m distance) to deliver olfactory stimulations. Briefly, animals were equipped with a tube inserted in the right nostril, allowing air circulation in the entire nasal cavity.

Two functional ingredients formulated by Phodé (Terssac, France) were used in this study: the *ingredient A* was mainly made of a natural extract of orange, *Citrus sinensis* (60-80%), and the *ingredient B* was mainly made of a natural extract of oregano, *Origanum vulgare* (40-55%) and lemongrass, *Cymbopogon flexuosus* (20-25%). Vehicle for both solutions was composed of distilled water (60-80%) and glyceryl polyethylene glycol ricinoleate (20-40%). For olfactory stimulation, solutions A and B were diluted in distilled water (1 L) at two concentrations (A1: 0.0105 %; A10: 0.105 %; B1: 0.02 %; B10: 0.2 %). The lowest concentration of ingredient A was comparable to the concentration used in a previous pig study (Val-Laillet, Meurice & Clouard, 2016). The lowest concentration of ingredient B was selected in order to get the same odor intensity (assessed by a human expert panel). The control solution consisted of the vehicle diluted in distilled water at the higher concentration used for ingredients (Control: 0.2 %). In order to obtain the highest brain responses for each animal, one block of stimulation/acquisition was performed for each ingredient and concentration: ingredient stimulation (30 sec, 4 L/min), and control stimulation (30 sec, 4 L/min) repeated 8 times. The order of ingredients was: A then B, from the lowest concentration to the highest for each ingredient in order to reduce odor saturation, resulting to the following order: A1, A10, B1, B10. The stimulation protocol duration per ingredient and per concentration was about 4 min, *i.e.* 16 min in total for both ingredients at two different concentrations.

2.4 MRI image acquisition

Imaging was performed during five days, with four animals being imaged per day. Image acquisition was performed as previously described (Coquery et al., 2018) on a 1.5-T magnet (Siemens Avanto, Erlangen, Germany) at the Rennes Platform for Multimodal Imaging and Spectroscopy (PRISM). Acquisitions were performed using a combination of coils (Body and Spine matrix coils) for optimized signal to noise ratio acquisition. *T1 weighted anatomical image acquisition*: a MP-RAGE sequence was adapted for adult minipig anatomy ($1.2 \times 1.2 \times 1.2 \text{ mm}^3$, NA=2, TR=2400 ms, TE=3.62 ms, TI=854 ms, FA=8°, acquisition duration 15 min). *BOLD (Blood-Oxygen-Level Dependent) signal acquisition*: an echo planar imaging sequence was adapted for pig head geometry (TR/TE: 2500/40 ms, FA: 90°, voxel size: $2.5 \times 2.5 \times 2.5 \text{ mm}^3$). The first four acquired volumes were excluded for the data analysis, meaning that no stimulation was performed during this period. For some animals, we detected a loss of MR signal in some part of the frontal lobes due to the anatomical presence of an air cavity anterior to the brain. The part of the frontal lobe that was not covered in all animals was thus excluded from the data analysis and is depicted as a dark grey area in the global brain activation maps (Figures 1-3).

2.5 Data analysis and statistical image analysis

Data analysis was performed with SPM12 (version 6906, Wellcome Department of Cognitive Neurology, London, UK). After slice timing correction, realignment and spatial normalization on a pig brain atlas (Saikali et al., 2010), images were smoothed with a Gaussian kernel of 4 mm. Due to limitations related to the size of the pig brain and the effect of anesthesia on brain activity, we used a non-standard statistical analysis with regards to human statistical standards. Further details regarding the validity and limitations of the statistical approach used in this model and paradigm are described in Coquery et al. (2018). *Voxel-based statistic*: first-

level (within-individual contrast) and second-level (within-group contrast) statistics were assessed with a threshold set at $p < 0.05$ to produce the brain maps of activation. *SVC-based statistics (Small Volume Correction)*: anatomical regions of interest (ROIs) from the Saikali pig brain atlas (Saikali et al., 2010) were used for SVC-based statistics with a p -value corrected for multiple ROIs comparison with a Bonferroni correction and at a threshold of 0.05 (peak level). For the analysis of the brain responses for each ingredient at different concentrations, sixteen ROIs corresponding to eight bilateral brain structures were used based on *a priori* hypothesis from a previous study performed with similar food ingredients (Val-Laillet, Meurice, & Clouard, 2016), (i) olfactory and gustatory sensory brain regions: prepyriform area and insular cortex; (ii) motivational and reward brain regions: caudate nucleus and putamen; (iii) associative learning brain regions: amygdala and parahippocampal cortex; and (iv) hedonic and executive brain regions: anterior prefrontal cortex and dorsal-lateral prefrontal cortex. The related uncorrected p -value threshold after Bonferroni correction was 0.0031 corresponding to a corrected p -value of 0.05. Uncorrected p -values between 0.0031 and 0.005 were considered as a statistical trend. For the comparison between ingredients, the brain regions related to olfaction and gustation were not included, resulting in a number of six analyzed brain regions corresponding to twelve ROIs bilaterally. The related uncorrected p -value after Bonferroni correction was 0.0042 corresponding to a corrected p -value of 0.05. Uncorrected p -values between 0.0042 and 0.005 were considered as a statistical trend.

For voxel-based statistics and SVC-based statistics, no suprathreshold voxels were detected with FDR (False Discovery Rate) correction at $p < 0.05$. Due to technical problem during functional acquisition, *i.e.* removal of the stimulation tube or animal heads movement, only sixteen animals were used for analysis.

3 Results

3.1 Brain responses to the ingredient A

Brain activation maps (Figure 1A). Independent of the concentration used, ingredient A promoted brain responses in sensory brain regions including the prepyriform area (PP), an olfactory relay/center, and the insular cortex (IC), a main gustatory center, in reward and motivational brain regions including the caudate nucleus (Cd), the putamen (Put) and the nucleus accumbens (Ac), and in associative learning brain regions, including a decreased activity in the amygdala (AMY). A ten-fold increase of ingredient A concentration (A10 vs. A1) promoted modulations of brain activation in sensory brain regions including the PP, in cingulate cortex areas (Ventral-Posterior: VP-CC, Dorsal-Posterior: DP-CC and Dorsal-Anterior: DA-CC), in the associative learning brain regions including a decreased activation in the AMY and the parahippocampal cortex (PHC). We could also detect an increased activation in reward and motivational brain regions, and in prefrontal brain regions, including the anterior prefrontal cortex (A-PFC) and the orbitofrontal cortex (OFC).

Corrected SVC-based statistic (Figure 1B). We detected an impact of ingredient A1 in reward and motivational brain regions: increased activation in the PUT and decreased activation in the Cd whereas A10 promoted a modulation in brain responses in the IC and a tendency toward a reduced activation in the A-PFC. A ten-fold increased of the ingredient A concentration (A10 vs A1) promoted a modulation in the reward and motivational brain regions, including a tendency toward increased activation in the PUT, a decreased activation in the IC and the PHC, and a tendency toward increased activation in the A-PFC. Of interest, is the fact that opposing brain responses were seen in the Cd between the left vs right hemispheres, with A10 (increased right Cd) vs A1 (increased left Cd) exposure.

3.2 Brain responses to the ingredient B

Brain activation maps (Figure 2A). Independent of the concentration used and similar to ingredient A, ingredient B promoted brain responses in sensory brain regions related to olfaction and gustation, and in reward and motivational brain regions except for the Ac. A ten-fold increase of ingredient B concentration (B10 vs. B1) promoted an increased activation in the PP, in the anterior cingulate brain regions (Ventral-Anterior: VA-CC and DA-CC) and in prefrontal brain regions (Dorsal-Lateral Prefrontal Cortex: DL-PFC and A-PFC), except for the OFC. We could also detect a brain activity modulation in reward and motivational brain regions as well as in the associative learning brain regions except for the AMY.

Corrected SVC-based statistic (Figure 2B). We detected an impact of ingredient B1 toward an increased activation in the IC, the PHC, and toward a decreased activation in the Cd and the DL-PFC. A10 promoted similar modulations with additional features, such as an increased activation in the A-PFC, a tendency toward increased activation in the AMY and no modulation in the DL-PFC. A ten-fold increased of the ingredient B concentration (B10 vs B1) promoted a decreased activation in the IC, and a tendency toward increased activation in the A-PFC and the PUT.

3.3 Different brain responses to the ingredient A and B

Brain activation maps (Figure 3A). A10 and B10 were able to promote different modulations of brain activity. A10 promoted higher brain responses in the Cd, a reward and motivational brain region, in the frontal lobes, including OFC but not A-PFC, whereas B10 promoted higher brain activation in the A-PFC, and in cingulate brain regions (DP-CC, VA-CC, DA-CC), and in associative learning brain regions, including the AMY and the PHC.

Corrected SVC-based statistic (Figure 3B). With respect to the brain activation maps, A10 was able to promote a higher activation in the reward and motivational brain region (Cd) whereas B10 promoted a higher activation in the A-PFC.

4 Discussion

For the first time, we described in the pig model the fMRI BOLD brain responses to contrasted and novel olfactory stimulations. We were able to detect the brain responses elicited by two plant-based sensory functional feed ingredients at two different concentrations. We also provided a comparison of the brain responses to these two ingredients at the higher concentration, for which higher activation of the reward and cognitive areas were detected compared to the lower concentrations tested.

A previous study from our lab had investigated, with fluoro-deoxy-glucose (FDG) positron emission tomography (PET), the brain responses under dual gustatory and olfactory stimulation from a single food ingredient with or without previous exposure, *i.e.* familiarization from weaning to testing a month later (Val-Laillet, Meurice, & Clouard, 2016). The ingredient used for this previous brain imaging study (similar to ingredient A in the present study) was chosen on the basis of the behavioral responses elicited by different feed ingredients at different concentrations (Clouard & Val-Laillet, 2014), because it led to increased palatability and acceptance of an unfamiliar diet the day of feed transition. Here, we implemented the experimental stimulation set-up to fMRI in order to investigate, within the same imaging session and without prior familiarization to the ingredients, the brain responses under olfactory stimulation with two different feed ingredients at two different concentrations. We have previously shown the potential of fMRI in the pig model for the screening of brain responses to pleasant/unpleasant gustatory stimulations (Coquery et al., 2018). In the present study, our paradigm was successful in screening the brain responses to each ingredient and allowed for direct comparison between them.

The main limitation of our paradigm is that the ingredient B was always tested after the ingredient A, which implies a possible order bias in brain activation. Interpretation of the

comparison between ingredients A and B must be done with cautious then, since brain responses might have been different had the order of exposure to the two ingredients been reversed. It is reassuring though to note that no general increase or decrease in brain activity was observed along time during the fMRI session, since the comparison between ingredients A and B led to increased responses in some brain areas and decreased responses in others. Be that as it may, for future experiments, the presentation order should be reversed in half of the subjects to compensate for this hypothetical order bias.

The choice of odor concentration is a major issue considering that, depending on the concentration used, a given olfactory compound might yield contrasted or even opposite responses, from pleasantness to aversion (Clouard, Jouhannau, Meunier-Salaün, Malbert, & Val-Laillet, 2012). Here, a ten-fold increase of ingredients A and B promoted an increased BOLD signal in the PP, a primary olfactory center/relay, and in frontal brain areas, suggesting a potential increased perception and integration along with increased concentration. Interestingly, a decreased BOLD signal was detected in the IC (gustatory and associative cortex), which usually also responds to olfactory stimulation, especially for taste-like aspects of food odors (Veldhuizen, Nachtigal, Teulings, Gitelman, & Small, 2010). Here, we used the orthonasal route for stimulation, which has been shown to promote distinct brain responses compared to the retronasal route (Small, Gerber, Mak, & Hummel, 2005). Additionally, the animals were naïve to both ingredients, preventing any association between these ingredients and the feed. This might account for a surprisingly decreased response in the IC with increased ingredient concentration. A ten-fold increase of ingredient A concentration (A10 vs. A1) tended to induce higher activation in the reward and motivational areas, *i.e.* the Cd and the PUT, but also, as seen with the global activation maps, to promote increased activation in the Ac which is involved in reward, and in the OFC, a brain region involved in olfactory/gustatory food valence (Anderson et al., 2003) and pleasantness/hedonism

(Grabenhorst, Rolls, Margot, Silva, & Velazco, 2007; Zou, van Harteveld, Kringelbach, Cheung, & Chan, 2016)11/26/2019 9:39:00 AM. Overall these observations suggested an increased pleasantness specific to the increase in ingredient A concentration.

In our study, both sensory ingredients A and B promoted brain responses in brain structures responding during early stages of odor processing such as the primary olfactory cortex (PP), hippocampus (HPC), PHC, AMY, and OFC (Stadlbauer et al., 2016). However, compared to each other, and at the higher concentration used, the two ingredients promoted contrasting brain responses. Ingredient A is mainly composed of *Citrus sinensis*, a food extract from sweet orange that has been reported to promote a broad spectrum of effects (Dosoky & Setzer, 2018) and was already tested in the pig using PET neuroimaging (Val-Laillet, Meurice, & Clouard, 2016), whereas ingredient B contained *Origanum vulgare*, of which one of the main active compounds is carvacrol, a molecule promoting positive effects in human health (Sharifi-Rad et al., 2018), but also modulating behavior in lambs (Simitzis, Deligeorgis, Bizelis, & Fegeros, 2008) and depression-like behavior in rat (Amiresmaeili et al., 2018). Ingredient B also contained *Cymbopogon flexuosus*, the so-called lemongrass, which has been reported in mice to elicit positive effects on mood regulation (Blanco et al., 2009), and notably reduced anxiety (Costa et al., 2011). Previous studies in humans either showed a reduced activation in the prefrontal cortex in association with “comfort” feeling after orange oil inhalation (Igarashi et al., 2014), or an increased cerebral blood flow in the prefrontal cortex associated with increased cognitive functions in response to flavone-rich citrus juice (Lamport et al., 2016). In our study, exposure to ingredient A promoted a lower activation in the A-PFC compared to exposure to ingredient B, but a higher activation in the DL-PFC, as seen in the global activation maps. The anterior (memory and perception) and dorsolateral (executive functions and attention) parts of the prefrontal cortex are linked to different cognitive processes, which probably explains the different outcomes observed in our study.

We showed here that the two ingredients used modulated differently both these areas *via* olfaction only. It would be very interesting to assess their effects in a pig model of chronic psychosocial stress and/or anxio-depressive symptoms for example, as modulations of the aforementioned brain areas were recently described in this model (Menneson et al. 2019).

Ingredient A promoted a higher activation in the reward/motivation brain regions, especially the Cd, which is in line with the activation in dopaminergic brain areas with food-related odors in humans (Sorokowska et al., 2017). It is also interesting to notice that the higher concentration of ingredient A activated the right Cd compared to its lower concentration, and that a reversed effect was observed in the left Cd. Despite that carvacrol has also been reported to promote similar effects on the dopaminergic system in rat (Zotti et al., 2013), and to enhance well-being in humans (Mechan et al., 2011), the activation in the related-brain regions, *i.e.* dorsal striatum (Cd and PUT) and OFC, was lower with ingredient B. This suggested that, at the concentrations used, ingredient A promoted higher pleasantness than ingredient B, which needs to be further confirmed by behavioral tests. Additional brain exploratory methods, such as *in vivo* nuclear brain imaging (Gautier et al., 2018) or post-mortem brain immunohistochemistry, might also be used to confirm that the dopaminergic system is specifically involved in the brain response to feed ingredients, with a focus on the striatum and prefrontal cortex, which are well-known dopaminergic areas that significantly responded to the olfactory stimulations in our study.

Exposure to ingredient B promoted higher brain responses in the AMY compared to exposure to ingredient A, as seen with the global activation maps. In humans, AMY is involved in olfactory hedonic processing-related tasks (Zou et al., 2016), and the higher activation with ingredient B could suggest a higher odor emotional salience (Winston, Gottfried, Kilner, & Dolan, 2005). However, given that carvacrol promotes in humans a perception of “warmth” through trigeminal stimulation (Klein et al., 2014), and that the AMY has been shown in rats

to receive trigeminal inputs (Hummel & Livermore, 2002), the higher activation in the AMY obtained with ingredient B compared to A might also be a consequence of the trigeminal stimulation. This is in line with a recent human study (Fournel, Ferdenzi, Sezille, Rouby, & Bensafi, 2016).

Finally, the olfactory stimulations used in our imaging paradigm did not permit to detect any modulation of brain activity in homeostatic brain areas, such as the hypothalamus. First, this might be due to limitations of the imaging method because the hypothalamus is a small brain structure composed of several specific nuclei, which complicates acquisition of images with a good resolution. The hypothalamus is also endowed with fenestrated capillaries probably enhancing interface imaging artifacts or noise. Second, we might also hypothesize that olfactory stimulation only was not sufficient in our study to modulate homeostatic signals and the hunger/satiety internal state, and that the observed effects were restricted to the hedonic dimension of eating control. Physiological measurements (*e.g.* gut hormones) concomitant to the olfactory stimulation might help in the future to disentangle this question.

5 Conclusion

We provided here a global comparison of the brain responses to two sensory functional feed ingredients at two different concentrations with fMRI in the pig model, and showed that increased concentration of ingredients promoted different brain responses, which were also dependent on the ingredient composition. The ingredient A, mainly composed of *Citrus sinensis* extracts was able to elicit higher brain responses in brain areas involved in pleasantness, compared to ingredient B that was principally composed of *Origanum vulgare* and *Cymbopogon flexuosus* extracts. Our findings might find relevance in the scope of the growing interest in the understanding of emotion regulation through olfaction modulation (Billot et al., 2017; Soudry et al., 2011).

Acknowledgements

This study and the PhD grant of Sophie Menneson were funded by Phodé, the ANRT (Agence Nationale Recherche Technologie, French Association for Research and Technology), and INRA. We thank the staff from the animal facility UEPR (Unité Expérimentale Porcs de Rennes, St Gilles, France) of INRA for their technical support: Serge Dubois, Bruno Fontaine, Renan Delaunay, Alain Chauvin. We acknowledge the PRISM (Plateforme de Recherche en Imagerie et Spectroscopie Multimodales, Rennes, France) core facility for its technical support, and especially Stéphane Quellec.

Conflict of interest statement

The functional feed/food ingredients used in this study were designed by Phodé that also co-funded, together with the ANRT (French Association for Research and Technology) and INRA, the research project and PhD grant of Sophie MENNESON. Authors Sophie MENNESON, Virginie NOIROT, and Pierre ETIENNE were employed by company Phodé. All other authors declare no competing interests.

Authors' contribution

Experimental design: NC, SM, PE, VN, and DVL. Technical development: NC, RJ, and PM. Performing the experiments: NC, SM, PM, and RJ. Data analysis: NC. Manuscript writing: NC and DVL. Manuscript revising: all co-authors.

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Figure

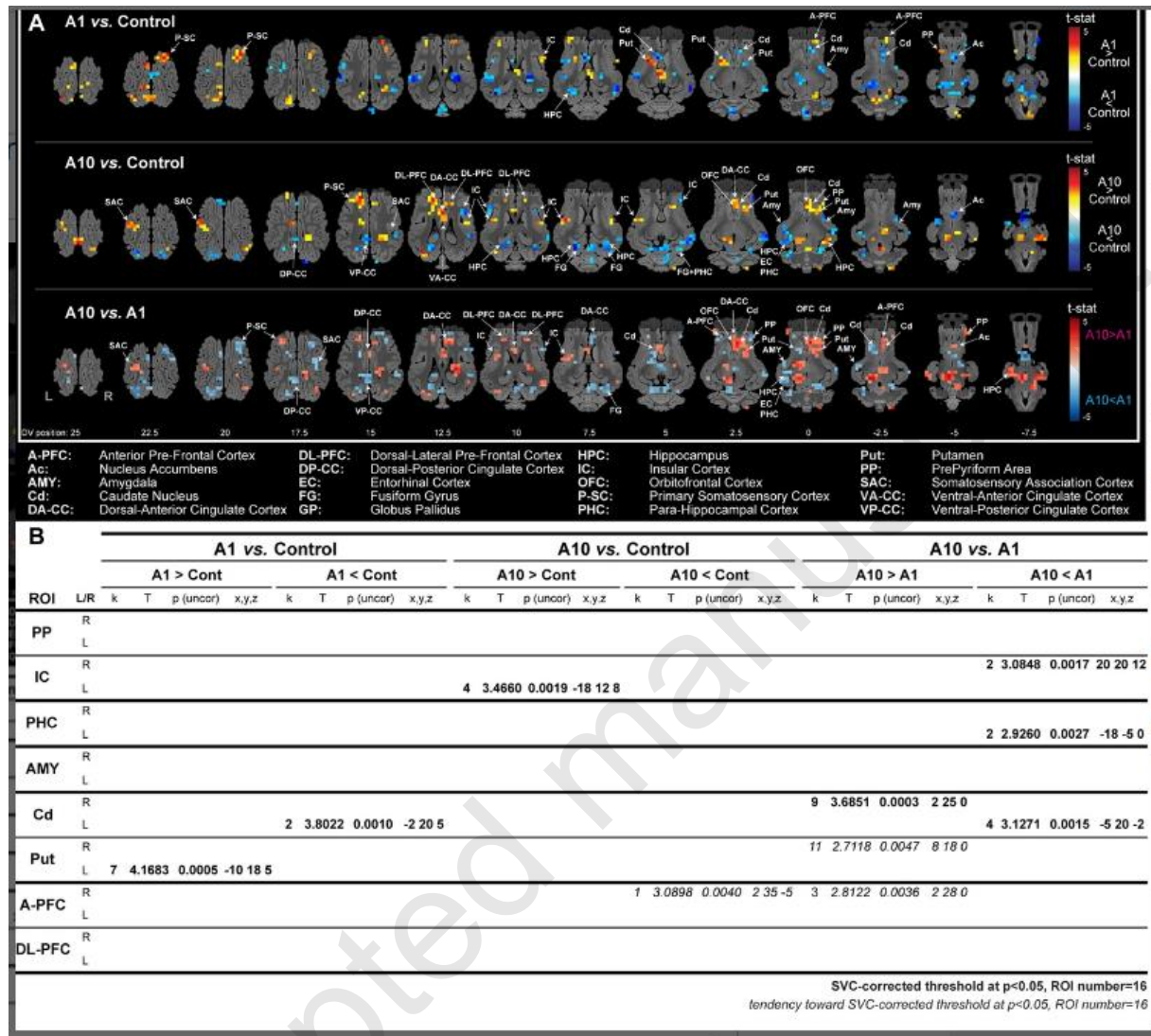


Figure 1-(A) Horizontal maps of global brain BOLD responses (contrast between average brain responses) to the functional ingredient A (at two concentrations: A1 and A10, which was ten-fold increased compared to A1) compared to control stimulations, and between the two concentrations of the ingredient A (A10 vs. A1). P-value threshold=0.05, $k > 4$, DV: dorsal-ventral position in mm related to the posterior commissure. The part of the brain that was not covered with the average BOLD-based statistical maps is superimposed in dark grey on the anatomical maps (frontal cortices). (B) SVC-based statistics: related ROIs with uncorrected p-value that reach the criteria of $p < 0.05$ after Bonferroni correction for multiple

ROIs comparison for sixteen regions of interest (ROI) from Saikali's atlas (Saikali et al., 2010) chosen upon *a priori* hypothesis. Statistical differences are depicted in bold, and tendency toward statistical differences are depicted in italic. L: ROI in the left hemisphere, and R: ROI in the right hemisphere. ROI abbreviations are detailed at the bottom of the panel A.

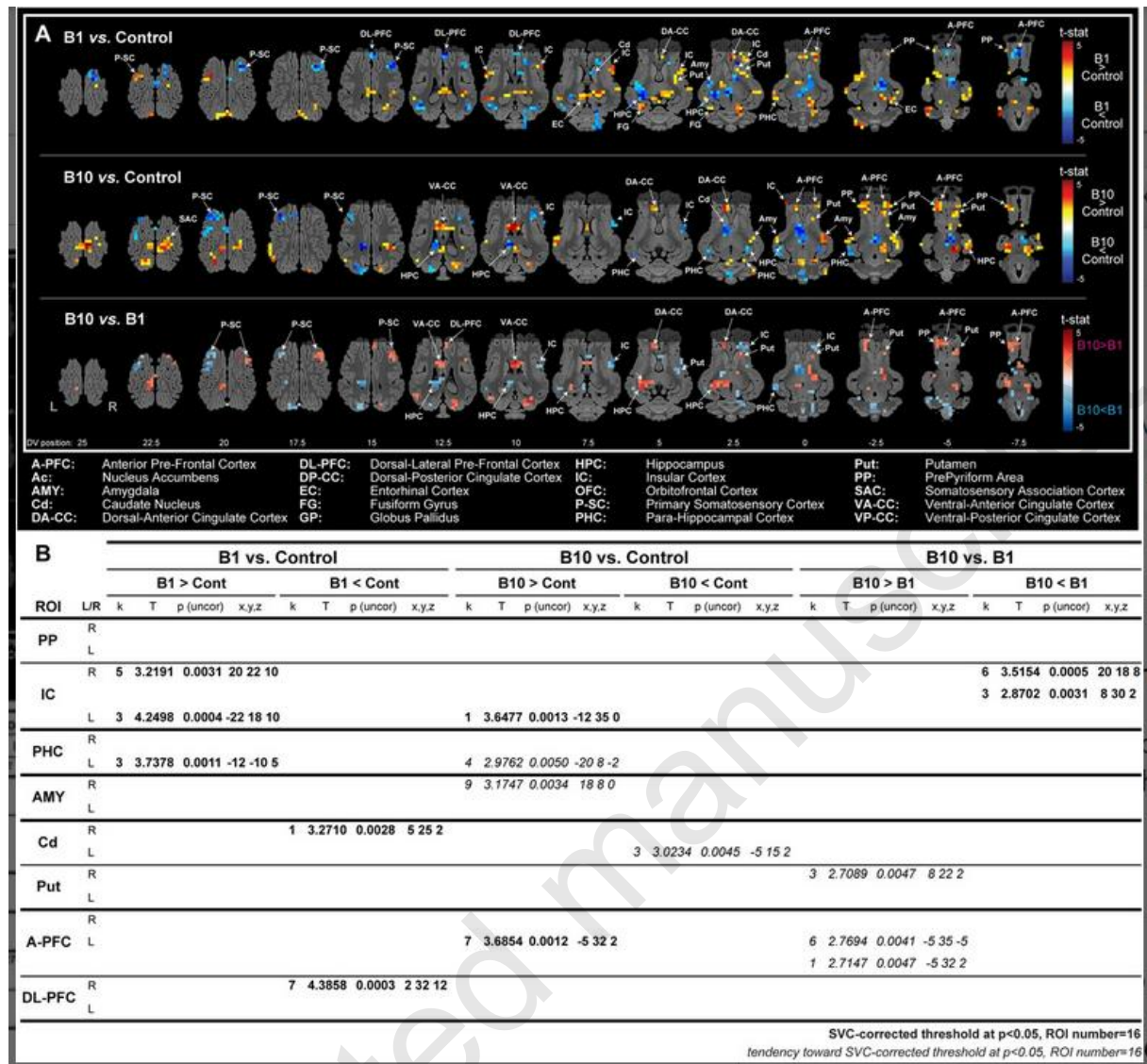


Figure 2-(A) Horizontal maps of global brain BOLD responses (contrast between average brain responses) to the ingredient B (at two concentrations: B1 and B10, which was ten-fold increased compared to B1) compared to control stimulations, and between the two concentrations of the ingredient B (B10 vs. B1). P-value threshold=0.05, $k > 4$, DV: dorsal-ventral position in mm related to the posterior commissure. The part of the brain that was not covered with the average BOLD-based statistical maps is superimposed in dark grey on the anatomical maps (frontal cortices). (B) SVC-based statistics: related ROIs with uncorrected p-value that reach the criteria of $p < 0.05$ after Bonferroni correction for multiple ROIs comparison for sixteen regions of interest (ROI) from Saikali's atlas (Saikali et al., 2010)

chosen upon *a priori* hypothesis. Statistical differences are depicted in bold, and tendency toward statistical differences are depicted in italic. L: ROI in the left hemisphere, and R: ROI in the right hemisphere. ROI abbreviations are detailed at the bottom of the panel A.

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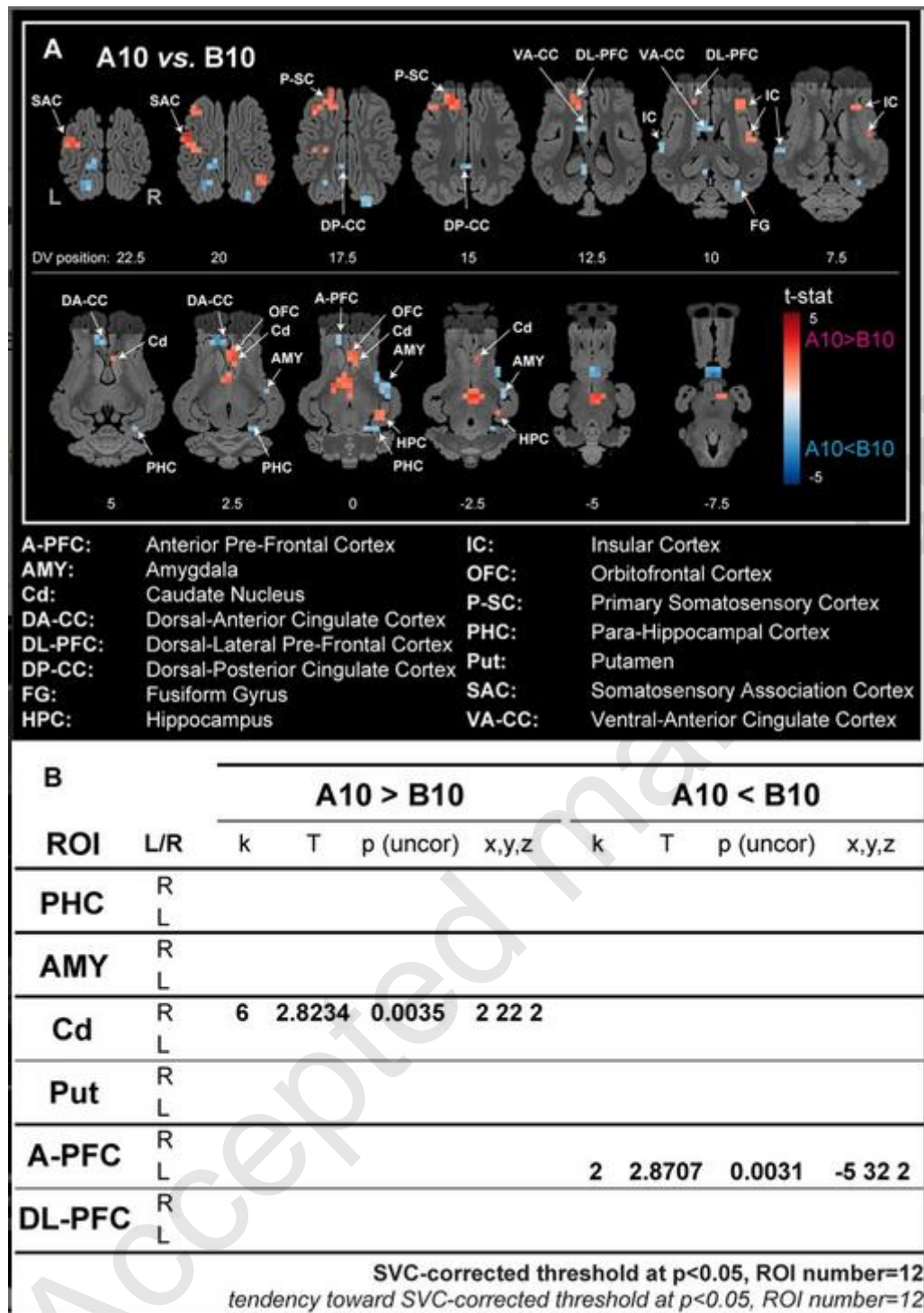


Figure 3-(A) Horizontal maps of global brain responses (contrast between average brain responses) between the ingredients A and B stimulations at the higher concentrations (A10 vs. B10). P-value threshold=0.05, DV: dorsal-ventral position in mm related to the posterior commissure. P-value threshold=0.05, $k > 4$, DV: dorsal-ventral position in mm related to the posterior commissure. The part of the brain that was not covered with the average BOLD-

based statistical maps is superimposed in dark grey on the anatomical maps (frontal cortices).

(B) SVC-based statistics: related ROIs with uncorrected p-value that reach the criteria of $p < 0.05$ after Bonferroni correction for multiple ROIs comparison for twelve regions of interest (ROI) from Saikali's atlas (Saikali et al., 2010) chosen upon *a priori* hypothesis. Statistical differences are depicted in bold, and tendency toward statistical differences are depicted in italic. L: ROI in the left hemisphere, and R: ROI in the right hemisphere. ROI abbreviations are detailed at the bottom of the panel A.