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1 **Potential of the microalgae *Nannochloropsis* and *Tetraselmis* for being**  
2 **used as innovative ingredients in baked goods**

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16 **Abstract**

17 The potential use of the microalgae species *Tetraselmis* and *Nannochloropsis* was  
18 investigated for the production of functional breads and crackers. Optimum flour  
19 substitution levels were 2.5% for baked crackers and 1.0 or 2.0% for breads containing  
20 *Nannochloropsis* or *Tetraselmis*, respectively. No major differences were observed in the  
21 physicochemical properties of the end products besides an expected darker and greener  
22 colour. Microalgae incorporation led to increased phenolic content and *in vitro*  
23 antioxidant capacity in both matrices. For example, the total phenolic content of crackers  
24 increased from  $24.6 \pm 1.5$  mg/100 g in the control to  $32.4 \pm 0.4$  or  $34.2 \pm 1.0$  mg/100 g in  
25 crackers containing *Tetraselmis* or *Nannochloropsis*, respectively. The amount of  
26 bioaccessible polyphenols after a simulated gastrointestinal digestion was also higher in  
27 microalgae-containing goods than in the controls. Sensory evaluation showed that  
28 microalgae-containing products were competitive with the controls with the added  
29 advantage of having an improved nutritional value and a “trendy” ingredient. Moreover,  
30 microalgae-containing products showed an increased emission of some volatile  
31 compounds such as p-cymene and (Z)-2-heptenal, which are responsible for fresh, citrus,  
32 terpenic, woody, and spicy or fatty, oily, and fruity odours, respectively.

33

34 **Keywords:** Functional foods, bread, crackers, *Nannochloropsis*, *Tetraselmis*.

## 35 **1. Introduction**

36 Humans are no strangers to the consumption of microalgae: already in the ninth century  
37 the Kanem Empire used *Arthrospira* as food in Africa (Oncel, Kose, Vardar, & Torzillo,  
38 2015). Nowadays, microalgae are generally marketed as nutritional supplements and  
39 promoted as “superfoods” that can be utilised as ingredients in the manufacture of  
40 “trendy” foods. For example, baked goods formulated using microalgae such as Wrawp  
41 (Wrawp Foods, CA, USA) and Helga Algae Crackers (Evasis Edibles GmbH, Bendorf,  
42 Austria) are currently commercially available.

43 A large number of scientific publications evaluated the potential of *Spirulina* and  
44 *Chlorella* for being used as ingredients in the manufacture of milkshakes, vegetable  
45 soups, snacks, pasta, yogurts, and baked goods including bread and biscuits (Lafarga,  
46 2019). This makes sense as both *Spirulina* and *Chlorella* are not only the most popular  
47 but also the most studied and cultivated microalgal strains (Garrido-Cardenas, Manzano-  
48 Agugliaro, Acien-Fernandez, & Molina-Grima, 2018). However, only a limited number  
49 of publications studied the effect of incorporating other species into this food group. For  
50 example, García-Segovia, Pagán-Moreno, Lara, and Martínez-Monzó (2017) reported  
51 that, although colour differences were observed when compared to the control, textural  
52 properties of the breads were not affected after incorporation of *Isochrysis galbana*,  
53 *Tetraselmis suecica*, *Scenedesmus almeriensis*, or *Nannochloropsis gaditana* at a  
54 concentration of 1.5% (w/w). Limited information is also available on the sensorial  
55 attributes of breads formulated using microalgae species different to *Spirulina* and  
56 *Chlorella*. Sensorial attributes of foods, especially flavour and aroma, are of key  
57 importance, as Western cultures do not seem to be willing to compromise taste for health.  
58 In addition, little is known on the effect of microalgae incorporation into other baked  
59 products different from bread. Some studies have been conducted on functional biscuits

60 enriched in: (i) eicosapentaenoic acid from *I. galbana* (Gouveia et al., 2008); (ii) fibre  
61 and protein from *S. platensis* (Singh, Singh, Jha, Rasane, & Gautam, 2015); and (iii)  
62 polyphenols and proteins from *S. platensis*, *C. vulgaris*, *T. suecica*, or *Phaeodactylum*  
63 *triconutum* (Batista et al., 2017). Biscuits or cookies are good delivery vehicles for health-  
64 promoting compounds because of their popularity and convenience. However, they  
65 contain a high sugar and/or fat (generally butter) content and it would be interesting to  
66 assess the effect of microalgae incorporation into other healthier products such as  
67 crackers.

68 Based on the current gap in knowledge, the aim of the current paper was to assess the  
69 potential of the species *Tetraselmis* and *Nannochloropsis*, which are currently  
70 underutilised in the food industry, for being used as novel ingredients for the production  
71 of functional breads and crackers. Studied quality parameters included volume, colour,  
72 texture, polyphenolic content, antioxidant activity, aroma volatile compounds, and  
73 sensorial attributes. In addition, the bioaccessibility of polyphenols after a simulated  
74 gastrointestinal digestion and the volatile profile of the products were also determined.

## 75 **2. Materials and methods**

### 76 **2.1 Preparation of the microalgae-containing breads and crackers**

77 Flour substitution levels evaluated in preliminary trials varied from 1 to 3% (w/w) for  
78 bread and from 1.25 to 3.75% (w/w) for crackers. Breads were produced following a  
79 straight dough baking procedure as described by Lafarga, Gallagher, Aluko, Auty, and  
80 Hayes (2016). Control wheat-only breads were labelled as BR-C. Breads containing  
81 *Tetraselmis* or *Nannochloropsis* at flour substitution levels of 2.0 or 1.0% (w/w) were  
82 labelled as BR-T and BR-N, respectively.

83 Crackers were produced following the methodology previously described by Lafarga et  
84 al. (2019a) with some modifications: in the current study, the doughs were sheeted to 2.0  
85 mm instead of 2.5 mm and were cut in 40 mm circles instead of squares. Control crackers  
86 were labelled as CR-C and crackers containing *Tetraselmis* or *Nannochloropsis* biomass  
87 at a concentration of 2.5% (w/w) were labelled as CR-T or CR-N, respectively.

### 88 **2.2 Physical analysis**

89 Colour recordings ( $L^*$ ,  $a^*$ , and  $b^*$  values) were taken using a Minolta CR-200 colorimeter  
90 (Minolta INC., Tokyo, Japan) and the D65 illuminant. Chroma ( $Ch$ ) and difference from  
91 the control ( $\Delta E$ ) were calculated in triplicate as described by Lafarga et al. (2019b) and  
92 determined on day 1 post-baking.

93 The weight and dimensions of ten crackers were averaged for each formulation and  
94 replicate. Cracker dimensions were measured at day 1 post-baking using a digital Vernier  
95 calliper (JP Selecta, Barcelona, Spain) and the spread ratios, specific volume, and density  
96 were calculated for each cracker as described by Jan, Panesar, and Singh (2018). Bread  
97 loaf volume was calculated using AACC Method 10-05.01.

98 Moisture content was determined using AACC Method 44-15.02. The water activity ( $a_w$ )  
99 of all samples was measured using an AquaLab meter (Decagon Devices Inc., WA, USA).  
100 Three measurements were taken for each formulation and replicate. The pH of 1 g of  
101 ground sample, added to 10 g of distilled water, was determined in triplicate per  
102 formulation and replicate using a Basic 20 pH-meter (Crison Instruments S.A., Barcelona,  
103 Spain).

104 Texture characteristics were assessed using a TA.XT2 Texture Analyser (Stable Micro  
105 Systems Ltd., Surrey, England) connected to Exponent software v.5.0.6.0. Texture profile  
106 analysis of the breads was conducted as described by Lafarga et al. (2019c) and using a  
107 P/20 aluminium compression probe. Crackers hardness was determined using a knife  
108 edge with slotted insert probe (HDP/BS) as described by Lafarga et al. (2019a). Ten  
109 samples were taken for each formulation and replicate.

### 110 **2.3 Total phenolic content**

111 The total phenolic content (TPC) of the breads and crackers was determined by the Folin  
112 Ciocalteu method, following the protocol described by Lafarga, Villaró, Bobo, Simó, and  
113 Aguiló-Aguayo (2019d). Extraction time was 2 h at room temperature. TPC was  
114 determined in triplicate and results were expressed as mg of gallic acid equivalents per  
115 100 g of dry weight (DW).

### 116 **2.4 Antioxidant activity**

117 The antioxidant capacity of the breads and crackers was determined using the same  
118 extract utilised for determination of TPC and using both the ferric reducing antioxidant  
119 power (FRAP) and the DPPH scavenging activity assays. The procedure followed was  
120 described previously by Lafarga et al. (2019d). Antioxidant capacity was determined in  
121 triplicate and results were expressed as mg of ascorbic acid equivalents per 100 g of DW.

## 122 **2.5 *In vitro* gastrointestinal digestion**

123 A simulated gastrointestinal digestion was performed in duplicate following the  
124 standardised static *in vitro* method previously described by Minekus et al. (2014). The  
125 method consists of three sequential states: (i) oral (37 °C, pH 7.0,  $\alpha$ -amylase, 2 min), (ii)  
126 gastric (37 °C, pH 3.0, pepsin, 2 h) and (iii) intestinal (37 °C, pH 7.0, pancreatin and fresh  
127 bile, 2 h). The pancreatin used contained enzymatic components including trypsin,  
128 amylase and lipase, ribonuclease, and protease. A blank was prepared using distilled  
129 water instead of sample. Determinations after the intestinal phase were performed in  
130 triplicate as described in previous sections.

## 131 **2.6 Sensorial analysis**

132 Sensory evaluation was undertaken by 30 semi-trained panellists (18 women, 12 men,  
133 age 18-50) recruited from IRTA Fruitcentre (Lleida, Spain) at day 1 post-baking. Sensory  
134 evaluation was conducted following the methodology described by Lafarga et al. (2019a).  
135 Each panellist assessed all the samples and was asked to indicate her or his opinion on  
136 the firmness, flavour, overall visual appearance, and overall acceptability of the products  
137 using a 9-point hedonic scale (from 1: extremely dislike to 9: extremely like). The  
138 acceptability index (AI) was calculated as described by Lucas, Morais, Santos, and Costa  
139 (2018). Finally, purchase intention (PI) was assessed using a 5-point hedonic scale which  
140 ranged from 1: “certainly would not buy” to 5: “certainly would buy”.

## 141 **2.7 Volatile compounds**

142 Extraction and determination of the volatile compounds emitted by the breads and  
143 crackers was performed using HS-SPME-GC/MS following the conditions previously  
144 described by Pico, Antolín, Román, Gómez, and Bernal (2018) with some modifications.  
145 Briefly, an amount of 1 g ( $\pm 0.005$  g) of ground sample was weighed into 20 mL vials and



146 mixed with 10 mL of 20% (w/v) sodium chloride at pH 3.0. The vials were immersed in  
147 a water bath at 60 °C and the SPME fibre (65 mm PDMS/DVB; Supelco Co., PA, USA)  
148 was exposed to the headspace for 60 min.

149 After extraction, the fibre was injected for thermal desorption into the injector port for 10  
150 min. The GC-MS analyses were performed using a 6890N gas chromatograph-mass  
151 spectrometer equipped with a HP-FFAP (50 m × 0.2 mm; 0.33 µm) column, both  
152 purchased from Agilent Technologies Inc. (CA, USA). Temperature conditions are  
153 described in the above cited publication. In this study, injector and detector temperatures  
154 were 240 °C. Mass spectra were obtained by electron impact ionisation at 70 eV and the  
155 scan mode was used to detect all the compounds in the range m/z 20-350. The preliminary  
156 identification of volatile compounds was verified by comparison of the mass spectral data  
157 obtained with those in NIST62 mass spectral database.

## 158 **2.8 Statistical analysis**

159 Results are expressed as mean ± standard deviation (S.D.). Differences between samples  
160 were analysed using analysis of variance (ANOVA) with JMP 13 (SAS Institute Inc.,  
161 Cary, USA). Where significant differences were present, a Tukey pairwise comparison  
162 of the means was conducted to identify where the sample differences occurred ( $p < 0.05$ ).

### 163 3. Results and discussion

#### 164 3.1 Preliminary baking trials

165 Incorporation of *Tetraselmis* and *Nannochloropsis* biomass into bread and crackers  
166 significantly affected colour parameters ( $p<0.05$ ):  $\Delta E$  was higher than 3 for all the  
167 formulated breads and crackers, suggesting that colour differences with the control were  
168 visible to the human eye. Higher microalgae content led in bread formulations to lower  
169  $L^*$  values for both crust and crumb ( $p<0.05$ ; Figure 1): a negative correlation was  
170 observed between microalgal biomass concentration and  $L^*$  values in crust (0.905; 0.05)  
171 and crumb (0.817; 0.05). Crackers with higher microalgal biomass concentration showed  
172 lower  $L^*$  values, suggesting a darker colour ( $p<0.05$ ; Figure 2). Similar results were  
173 reported previously (Figueira, Crizel, Silva, & Salas-Mellado, 2011; Menezes, Coelho,  
174 Meza, Salas-Mellado, & Souza, 2015). Although  $a^*$  values of the microalgae-containing  
175 breads were lower and  $b^*$  values were higher, when compared to the control,  
176 incorporation of higher concentrations of microalgae did not cause further differences in  
177  $a^*$  and  $b^*$  values. These results may seem unexpected but this same effect was reported  
178 in baked products containing *S. platensis* (Batista et al., 2017), *C. vulgaris* (Gouveia,  
179 Batista, Miranda, Empis, & Raymundo, 2007), and *I. galbana* (Gouveia et al., 2008) and  
180 has been attributed to pigment degradation during the baking process and/or to a pigment  
181 saturation effect above a certain microalgae concentration.

182 Before discussing the sensorial acceptance of the breads and crackers it is important to  
183 highlight that panellists were first asked if they would be willing to buy baked products  
184 enriched in microalgae and only those who answered “yes” conducted the sensorial  
185 analysis. Moreover, results on sensorial analysis must be taken with caution, especially  
186 those on overall acceptance and PI, as the ideal would have been to assess these  
187 parameters using ~100 consumers. For a product to be accepted in terms of sensorial

188 characteristics, it is necessary to obtain an AI greater than 70% (Lucas et al., 2018).  
189 Although microalgae incorporation into the bread led to lower overall acceptability scores  
190 ( $p<0.05$ ), formulated breads showed AI values ranging between 71.7 and 80.8%. These  
191 values are in line with those reported for other foods containing microalgae (Lafarga et  
192 al., 2019b; Lucas et al., 2018). Maximum AI was obtained for breads containing  
193 *Tetraselmis* at a concentration of 2.0% and *Nannochloropsis* at a concentration of 1.0%.  
194 These breads showed relatively high PI values: approximately 55% of the panellists said  
195 they “probably would buy” them. Approximately the same amount of panellists suggested  
196 that they “probably would buy” the control breads, although these showed a higher  
197 percentage of panellists who “certainly would buy” them. Overall acceptability of  
198 crackers was not affected after incorporation of microalgae into the recipe. All of the  
199 microalgae-containing crackers showed AI values over 70%. Crackers containing  
200 *Tetraselmis* and *Nannochloropsis* at a flour substitution level of 2.5% (w/w) showed AI  
201 values of 85.9 and 79.8%, respectively. Approximately 82 and 91% of the panellists  
202 scored these two crackers within the range 7-9 (between “like moderately” and “like  
203 extremely”) and their PI ranged between 4 and 5 (between “would probably buy” and  
204 “certainly would buy”). Microalgae concentration higher than 2.5% resulted in decreased  
205 AI and PI values ( $p<0.05$ ).

206 Based on these results, breads containing *Tetraselmis* at a concentration of 2.0% (w/w)  
207 or *Nannochloropsis* at a flour substitution level of 1.0% (w/w) were selected for further  
208 analysis. These were labelled as BR-T and BR-N, respectively. Moreover, crackers  
209 containing *Tetraselmis* or *Nannochloropsis* at a flour substitution level of 2.5% (w/w),  
210 which were labelled as CR-T and CR-N were also selected for further analysis.

## 211 **3.2 Physicochemical properties**

### 212 **3.2.1 Colour and volume**

213 No differences were observed between the colour attributes of the breads at day 3 post-  
214 baking (data not shown) when compared to those measured at day 1 (Figure 1) - except  
215 for a decrease in crust  $L^*$  values ( $p<0.05$ ), probably caused by a loss of moisture during  
216 storage. Moreover, no colour differences were detected during storage of crackers for 10  
217 days. Colour values during storage suggest a stable product in terms of visual appearance.  
218 In the current study, the specific volume of BR-T and BR-N was lower than that of BR-  
219 C ( $p<0.05$ ; Table 1). Results can be attributed to a dilution of starch and gluten after  
220 substituting flour with microalgae and a decrease in the amount of fully hydrated starch  
221 granules caused by the added powder competing for water with starch. The lower loaf  
222 volume obtained after incorporation of microalgae into the recipe led to higher density in  
223 BR-N when compared to BR-C ( $p<0.05$ ). In addition, microalgae-incorporation into the  
224 crackers formulation did not affect volume and density, suggesting that higher microalgae  
225 concentrations can be incorporated into crackers without negatively affecting the visual  
226 appearance of the products (when compared to bread). A high spread ratio, which is a  
227 quality measure, is desirable in baked products (Mudgil, Barak, & Khatkar, 2017). The  
228 spread ratio of the crackers was not affected after the incorporation of microalgae into the  
229 crackers' recipe. Previous studies suggested an increase of the spread ratio of crackers  
230 enriched in powdered broccoli co-products and reported a positive correlation between  
231 spread ratio and broccoli content (Lafarga et al., 2019a). Higher microalgae concentration  
232 could probably lead to higher spread ratios, although this would need to be assessed in  
233 further studies.

### 234 **3.2.2 Moisture and water activity**

235 Moisture content and  $a_w$  values of the breads was comparable to that measured in the  
236 crumb of commercially available bagels or breads (Schmidt & Fontana, 2008).  
237 Incorporation of microalgae into the bread formulations led to lower moisture content

238 ( $p<0.05$ ). The moisture content of BR-T was lower than that of BR-N ( $p<0.05$ ). A  
239 decrease in moisture was observed at day 3 post-baking because of bread staling ( $p<0.05$ ).  
240 Water loss during storage was calculated as 18.5, 9.0, and 5.3% for BR-C, BR-T, and  
241 BR-N, respectively. Microalgae incorporation into the crackers also led to reduced  
242 humidity at day 1 post-baking ( $p<0.05$ ). However, no significant differences were  
243 observed between the moisture content at days 1, 5, and 10 post-baking, suggesting stable  
244 products.  
245 Substituting wheat flour with microalgal biomass did not affect the pH and the  $a_w$  of the  
246 breads at day 1 post-baking (Table 1). A decrease in  $a_w$  was observed in bread samples  
247 during storage ( $p<0.05$ ). The observed decrease was bigger in BR-C when compared to  
248 BR-T and BR-N, probably caused by a higher moisture loss during storage. Similar  $a_w$   
249 values were also observed in breads enriched in bioactive ingredients (Lafarga et al.,  
250 2016). The  $a_w$  of CR-T and CT-N was lower than that CR-C, caused by the above  
251 mentioned lower moisture content. Storage for 10 days did not affect pH and  $a_w$  values  
252 for any of the cracker samples, suggesting once again stable products.

### 253 **3.2.3 Textural properties**

254 Figure 3 shows the textural properties of BR-C, BR-T, and BR-N at days 1 and 3 post-  
255 baking. A higher bread density has often been correlated with increased hardness.  
256 However, in the current study, no differences were observed in hardness, which is the  
257 peak force that occurred during the compression of the bread slices. Similar results were  
258 observed after incorporation of freeze-dried broccoli co-products into bread at a  
259 concentration of 2% (Lafarga et al., 2019c). The observed increase in hardness at day 3,  
260 when compared to the values obtained at day 1, can be attributed to bread staling and  
261 moisture loss. Moreover, no differences in springiness, cohesiveness, gumminess,  
262 chewiness, and resilience were observed between the microalgae-containing breads BR-

263 T or BR-N and BR-C, suggesting a comparable mouth-feel and a similar retention of the  
264 textural properties after compression at both days 1 and 3 post-baking. Results were in  
265 line with those reported by García-Segovia et al. (2017).  
266 Hardness of the control and microalgae-containing crackers, which is the force required  
267 to break or snap the cracker, is shown in Figure 3. Lower moisture content is correlated  
268 with increased hardness in crackers (Millar et al., 2017). However, no significant  
269 differences between the hardness of CR-T, CR-N, and CR-C. As shown in Figure 3, no  
270 differences in hardness were observed during storage. Similar results were observed  
271 previously in other food matrices where microalgae incorporation did not affect  
272 functional properties of the end products (De Marco, Steffolani, Martínez, & León, 2014;  
273 García-Segovia et al., 2017).

### 274 **3.3 Total phenolic content and antioxidant capacity**

275 Currently, algae-derived polyphenols are one of the top trends in functional foods for the  
276 prevention of cardiovascular diseases and diabetes (Murray, Dordevic, Ryan, & Bonham,  
277 2018). Microalgae incorporation led to increased TPC in both studied food matrices  
278 ( $p < 0.05$ ) and, as expected, to an increased antioxidant capacity (Figure 4). Results are not  
279 surprising as several studies reported the high antioxidant activity of microalgal biomass,  
280 which has been attributed to their high phenolic and carotenoid content (Goiris et al.,  
281 2012). A positive correlation was observed between TPC and antioxidant capacity of  
282 crackers at when assessed using the FRAP (0.884 at day 1 and 0.986 at day 5; 0.05) and  
283 DPPH (0.852 at day 1 and 0.991 at day 5; 0.05) methods. Previous studies also reported  
284 an increased content of polyphenols and a higher antioxidant capacity after incorporation  
285 of microalgae in, for example, pasta (De Marco et al., 2014) or broccoli soup (Lafarga et  
286 al., 2019b).

287 Results shown in Figure 4 demonstrate that the amount of polyphenols in the enzymatic  
288 digestive extracts obtained after a simulated gastrointestinal digestion is higher than that  
289 expected based on extractions made using methanol ( $p<0.05$ ). This was probably caused  
290 by a higher liberation of polyphenols because of the action of digestive enzymes. The  
291 longer extraction time can also partially explain these findings. Not only the phenolic  
292 content but also the antioxidant capacity of the enzymatic digestive extracts was higher  
293 than that of the methanolic extracts ( $p<0.05$ ). The observed increase in antioxidant  
294 capacity can be attributed to the higher phenolic content but also to the generation of  
295 bioactive peptides with antioxidant capacity, as previous studies demonstrated that  
296 microalgae are good sources of bioactive peptides (Ko et al., 2018; Wu, Xu, Sun, Yu, &  
297 Zhou, 2015). Food processing is a crucial step to improve bioaccessibility and produce  
298 products with beneficial nutritional properties (Barba et al., 2017). Cavonius, Albers, and  
299 Undeland (2016) suggested that cell disruption, and to a lesser extent, strong pH  
300 variations were needed to increase bioaccessibility of lipids while a pre-freezing step was  
301 required to improve accessibility of proteins derived from *Nannochloropsis oculata*. The  
302 bioaccessibility and bioavailability of other antioxidant compounds found in microalgae  
303 such as carotenoids have been shown to strongly depend on, for example, the food matrix  
304 and processing conditions (Kopec and Failla, 2018).

### 305 **3.4 Sensorial attributes and volatile profile**

306 Incorporation of microalgae into the bread formulation, at the concentrations studied  
307 herein, did not affect the visual appearance scores of the breads. Lafarga et al. (2019c)  
308 recently reported high visual acceptability scores of a green bread formulated using  
309 broccoli leaves. No differences were observed in the texture and flavour scores of BR-C  
310 and BR-T or BR-N, although some panellists (3 out of 30) described an unpleasant “fishy”  
311 taste. Incorporation of microalgae into the bread formulation led to lower aroma scores

312 ( $p<0.05$ ). The aroma score of BR-T was lower than that of BR-N ( $p<0.05$ ), probably  
313 caused by a higher microalgae content. Incorporation of microalgal biomass into the  
314 cracker formulation led to a decreased visual appearance when compared to CR-C  
315 ( $p<0.05$ ; Figure 5). Although green crackers containing *Chlorella* biomass, at a  
316 concentration of 5%, are currently being commercialised under the brand Helga Algen  
317 Cracker (Evasis Edibles, Austria), the number of green-coloured baked goods currently  
318 being commercialised in Europe is still limited. European consumers are not yet used to  
319 coloured baked products and this could be the cause of the observed lower visual  
320 appearance scores. Moreover, no differences were detected between the aroma scores of  
321 CR-T, CR-N, and the control CR-C. In turn, flavour and overall acceptance scores were  
322 higher in microalgae-containing crackers when compared to the control ( $p<0.05$ ) – don't  
323 forget that only panellists that would be willing to buy microalgae-enriched products  
324 carried out the sensorial analysis. As mentioned previously, these results are preliminary  
325 and a sensorial analysis with a larger number of consumers would better describe the  
326 sensorial attributes and the acceptance of these products.

327 Volatile compounds identified in the breads and crackers are listed in Table 2. A total of  
328 42 compounds including alcohols (5), aldehydes (12), ketones (4), esters (11), sulfur  
329 compounds (2), acids (1), terpenes (4) and hydrocarbons (3) were detected in the samples.  
330 The most abundant compounds in crackers were (listed in decreasing order) undecanone,  
331 hexanoic acid, dipropyl disulfide, nonanal, phenylacetaldehyde, 3-methyl-1-butanal, and  
332 2-methyl-1-butanal. Regarding aldehydes, the most important odorant was nonanal,  
333 which is formed from  $\beta$ -cleavage of the 10-OOH hydroperoxide and imparts green and  
334 fatty notes to flavour (Parker, 2015). Among the Strecker aldehydes of the amino acid  
335 methionine, phenylacetaldehyde and 2- and 3-methylbutanal were the most prominent in  
336 the aroma profile of crackers, which were also detected in previous HS-SPME studies on



337 wheat bread (Raffo, Carcea, Castagna, & Magrì, 2015). Most relevant alcohols were 3-  
338 methyl-1-butanol, 1-octen-3-ol, and 2-ethyl-1-hexanol. 3-Methyl-1-butanol is produced  
339 during dough fermentation (Mildner-Szkudlarz, Zawirska-Wojtasiak, Szwengiel, &  
340 Pacyński, 2011). In the group of ketones, besides identifying aroma compounds such as  
341 1-octen-3-one, 6-methyl-5-hepten-2-one, and 2-nonanone, we would like to highlight the  
342 identification of 2-undecanone, which was the most abundant ketone in this study and  
343 was not reported in previous HS-SPME analyses on bread (Pacyński, Wojtasiak, &  
344 Mildner-Szkudlarz, 2015). The groups of esters, acids, sulphides, terpenes, and  
345 hydrocarbons completed the list of identified compounds. The aroma profile of bread and  
346 crackers was similar. However, in bread the most significant aldehyde was (Z)-2-  
347 heptenal, which is a product from the degradation of the linoleic acid (Parker, 2015).

348 **4. Conclusions**

349 Overall, *Tetraselmis* and *Nannochloropsis* biomass show potential for being used as novel  
350 functional ingredients in bread and crackers. Results demonstrated that not only the *in*  
351 *vitro* phenolic content or antioxidant capacity of the products was improved after  
352 microalgae-incorporation but also the amount of bioaccessible polyphenols and the  
353 antioxidant capacity of the enzymatic digestive extracts, suggesting also healthier  
354 products. Their utilisation would also allow food processors to differentiate by using a  
355 “trendy” ingredient. Sensory evaluation showed that microalgae-containing breads and  
356 crackers, enriched at the concentrations studied in the current study, were competitive  
357 with the control breads and crackers with the added advantage of having an improved  
358 nutritional value.

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364 **References**

- 365 Barba, F. J., Mariutti, L. R. B., Bragagnolo, N., Mercadante, A. Z., Barbosa-Cánovas, G. V.,  
366 & Orlien, V. (2017). Bioaccessibility of bioactive compounds from fruits and  
367 vegetables after thermal and nonthermal processing. *Trends in Food Science &*  
368 *Technology*, *67*, 195-206.
- 369 Batista, A. P., Niccolai, A., Fradinho, P., Fragoso, S., Bursic, I., Rodolfi, L., . . . Raymundo,  
370 A. (2017). Microalgae biomass as an alternative ingredient in cookies: Sensory,  
371 physical and chemical properties, antioxidant activity and *in vitro* digestibility. *Algal*  
372 *Research*, *26*, 161-171.
- 373 Cavonius, L. R., Albers, E., & Undeland, I. (2016). *In vitro* bioaccessibility of proteins and  
374 lipids of pH-shift processed *Nannochloropsis oculata* microalga. *Food & Function*, *7*,  
375 2016-2024.
- 376 De Marco, E. R., Steffolani, M. E., Martínez, C. S., & León, A. E. (2014). Effects of spirulina  
377 biomass on the technological and nutritional quality of bread wheat pasta. *LWT*, *58*,  
378 102-108.
- 379 Figueira, F. d. S., Crizel, T. d. M., Silva, C. R., & Salas-Mellado, M. d. I. M. (2011). Elaboration  
380 of gluten-free bread enriched with the microalgae *Spirulina platensis*. *Brazilian Journal*  
381 *of Food Technology*, *14*, 308-316.
- 382 García-Segovia, P., Pagán-Moreno, M. J., Lara, I. F., & Martínez-Monzó, J. (2017). Effect of  
383 microalgae incorporation on physicochemical and textural properties in wheat bread  
384 formulation. *Food Science and Technology International*, *23*, 437-447.
- 385 Garrido-Cardenas, J. A., Manzano-Agugliaro, F., Acien-Fernandez, F. G., & Molina-Grima,  
386 E. (2018). Microalgae research worldwide. *Algal Research*, *35*, 50-60.
- 387 Goiris, K., Muylaert, K., Fraeye, I., Foubert, I., De Brabanter, J., & De Cooman, L. (2012).  
388 Antioxidant potential of microalgae in relation to their phenolic and carotenoid content.  
389 *Journal of Applied Phycology*, *24*, 1477-1486.
- 390 Gouveia, L., Batista, A. P., Miranda, A., Empis, J., & Raymundo, A. (2007). *Chlorella vulgaris*  
391 biomass used as colouring source in traditional butter cookies. *Innovative Food Science*  
392 *& Emerging Technologies*, *8*, 433-436.
- 393 Gouveia, L., Coutinho, C., Mendonça, E., Batista, A. P., Sousa, I., Bandarra, N. M., &  
394 Raymundo, A. (2008). Functional biscuits with PUFA- $\omega$ 3 from *Isochrysis galbana*.  
395 *Journal of the Science of Food and Agriculture*, *88*, 891-896.
- 396 Jan, K. N., Panesar, P. S., & Singh, S. (2018). Optimization of antioxidant activity, textural and  
397 sensory characteristics of gluten-free cookies made from whole indian quinoa flour.  
398 *LWT*, *93*, 573-582.
- 399 Ko, S.-C., Heo, S.-Y., Choi, S.-W., Qian, Z.-J., Heo, S.-J., Kang, D.-H., . . . Jung, W.-K. (2018).  
400 A heptameric peptide isolated from the marine microalga *Pavlova lutheri* suppresses  
401 PMA-induced secretion of matrix metalloproteinase-9 through the inactivation of the

- 402 JNK, p38, and NF- $\kappa$ B pathways in human fibrosarcoma cells. *Journal of Applied*  
403 *Phycology*, 30, 2367-2378.
- 404 Koec, M. L., & Failla, M. L. (2018). Recent advances in the bioaccessibility and bioavailability  
405 of carotenoids and effects of other dietary lipophiles. *Journal of Food Composition and*  
406 *Analysis*, 68, 16-30.
- 407 Lafarga, T. (2019). Effect of microalgal biomass incorporation into foods: Nutritional and  
408 sensorial attributes of the end products. *Algal Research*, 41, 101566.
- 409 Lafarga, T., Acién-Fernández, F. G., Castellari, M., Villaró, S., Bobo, G., & Aguiló-Aguayo,  
410 I. (2019b). Effect of microalgae incorporation on the physicochemical, nutritional, and  
411 sensorial properties of an innovative broccoli soup. *LWT*, 111, 167-174.
- 412 Lafarga, T., Gallagher, E., Aluko, R. E., Auty, M. A. E., & Hayes, M. (2016). Addition of an  
413 enzymatic hydrolysate of bovine globulins to bread and determination of hypotensive  
414 effects in spontaneously hypertensive rats. *Journal of Agricultural and Food*  
415 *Chemistry*, 64, 1741-1750.
- 416 Lafarga, T., Gallagher, E., Bademunt, A., Bobo, G., Echeverria, G., Viñas, I., & Aguiló-  
417 Aguayo, I. (2019a). Physicochemical and nutritional characteristics, bioaccessibility and  
418 sensory acceptance of baked crackers containing broccoli co-products. *International*  
419 *Journal of Food Science & Technology*, 54, 634-640.
- 420 Lafarga, T., Gallagher, E., Bademunt, A., Viñas, I., Bobo, G., Villaró, S., & Aguiló-Aguayo,  
421 I. (2019c). Bioaccessibility, physicochemical, sensorial, and nutritional characteristics  
422 of bread containing broccoli co-products. *Journal of Food Processing and*  
423 *Preservation*, 43, e13861.
- 424 Lafarga, T., Villaró, S., Bobo, G., Simó, J., & Aguiló-Aguayo, I. (2019d). Bioaccessibility and  
425 antioxidant activity of phenolic compounds in cooked pulses. *International Journal of*  
426 *Food Science & Technology*, 54, 147-157.
- 427 Lucas, B. F., Morais, M. G. d., Santos, T. D., & Costa, J. A. V. (2018). *Spirulina* for snack  
428 enrichment: Nutritional, physical and sensory evaluations. *LWT*, 90, 270-276.
- 429 Menezes, B., Coelho, M., Meza, S., Salas-Mellado, M., & Souza, M. (2015). Macroalgal  
430 biomass as an additional ingredient of bread. *International Food Research Journal*, 22,  
431 819-824.
- 432 Mildner-Szkudlarz, S., Zawirska-Wojtasiak, R., Szwengiel, A., & Pacyński, M. (2011). Use of  
433 grape by-product as a source of dietary fibre and phenolic compounds in sourdough  
434 mixed rye bread. *International Journal of Food Science & Technology*, 46, 1485-1493.
- 435 Millar, K. A., Barry-Ryan, C., Burke, R., Hussey, K., McCarthy, S., & Gallagher, E. (2017).  
436 Effect of pulse flours on the physicochemical characteristics and sensory acceptance of  
437 baked crackers. *International Journal of Food Science & Technology*, 52, 1155-1163.
- 438 Minekus, M., Alming, M., Alvito, P., Ballance, S., Bohn, T., Bourlieu, C., . . . Dupont, D.  
439 (2014). A standardised static *in vitro* digestion method suitable for food—an  
440 international consensus. *Food & Function*, 5, 1113-1124.

- 441 Mudgil, D., Barak, S., & Khatkar, B. (2017). Cookie texture, spread ratio and sensory  
442 acceptability of cookies as a function of soluble dietary fiber, baking time and different  
443 water levels. *LWT*, *80*, 537-542.
- 444 Murray, M., Dordevic, A. L., Ryan, L., & Bonham, M. P. (2018). An emerging trend in  
445 functional foods for the prevention of cardiovascular disease and diabetes: Marine algal  
446 polyphenols. *Critical Reviews in Food Science and Nutrition*, *58*, 1342-1358.
- 447 Oncel, S. S., Kose, A., Vardar, F., & Torzillo, G. (2015). From the ancient tribes to modern  
448 societies, microalgae evolution from a simple food to an alternative fuel source. In S.  
449 K. Kim (Ed.), *Handbook of marine microalgae: Biotechnology advances* (pp. 127-138).  
450 MA, USA: Academic Press.
- 451 Pacyński, M., Wojtasiak, R. Z., & Mildner-Szkudlarz, S. (2015). Improving the aroma of  
452 gluten-free bread. *LWT*, *63*, 706-713.
- 453 Parker, J. K. (2015). Thermal generation of aroma. In J. K. Parker, S. Elmore & L. Methven  
454 (Eds.), *Flavour development, analysis and perception in food and beverages* (pp. 151-  
455 186). MA, USA: Woodshed Publishing.
- 456 Pico, J., Antolín, B., Román, L., Gómez, M., & Bernal, J. (2018). Analysis of volatile  
457 compounds in gluten-free bread crusts with an optimised and validated SPME-  
458 GC/QTOF methodology. *Food Research International*, *106*, 686-695.
- 459 Raffo, A., Carcea, M., Castagna, C., & Magrì, A. (2015). Improvement of a headspace solid  
460 phase microextraction-gas chromatography/mass spectrometry method for the analysis  
461 of wheat bread volatile compounds. *Journal of Chromatography A*, *1406*, 266-278.
- 462 Schmidt, S. J., & Fontana, A. J. J. (2008). Water activity values of selected food ingredients  
463 and products. In G. V. Barbosa-Cánovas, A. J. J. Fontana, S. J. Schmidt & T. P. Labuza  
464 (Eds.), *Water activity in foods: Fundamentals and applications* (pp. 407-420). Iowa,  
465 USA: Blackwell Publishing.
- 466 Singh, P., Singh, R., Jha, A., Rasane, P., & Gautam, A. K. (2015). Optimization of a process  
467 for high fibre and high protein biscuit. *Journal of Food Science and Technology*, *52*,  
468 1394-1403.
- 469 Wu, H., Xu, N., Sun, X., Yu, H., & Zhou, C. (2015). Hydrolysis and purification of ACE  
470 inhibitory peptides from the marine microalga *Isochrysis galbana*. *Journal of Applied*  
471 *Phycology*, *27*, 351-361.