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Additional Information

1 **UNDERSTANDING EMULSIFIERS EFFECT ON BREAD AERATION**  
2 **DURING BREADMAKING**

3

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5

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13

14 **Running title: Emulsifiers role in bread aeration**

15 **Abstract**

16 BACKGROUND: Much research has been done to explain emulsifiers action during  
17 breadmaking, but there is still plenty unknown to elucidate their functionality despite  
18 their diverse chemical structure. The aim of the present study was to provide some light  
19 about the role of emulsifiers on air incorporation into the dough and gas bubbles  
20 progress during baking and their relationship with bread features. Emulsifiers like  
21 diacetyl tartaric acid ester of monoglycerides (DATEM), sodium stearyl lactylate  
22 (SSL), distilled monoglyceride (DMG-45 and DMG-75), lecithin and polyglycerol  
23 esters of fatty acids (PGEF) were tested in very hydrated doughs. RESULTS:  
24 Emulsifiers increased the maximum dough volume during proofing. Emulsifiers  
25 increased the number of bubbles incorporated during mixing, observing higher number  
26 of bubbles, particularly with PGEF. Major changes in dough occurred at 70 K when  
27 bubble size augmented, becoming more heterogeneous. DMG-75 produced the biggest  
28 bubbles. As a consequence, emulsifiers tend to increase the number of gas cells with  
29 lower size in the bread crumb, but led to greater crumb firmness, which suggested  
30 different interactions between emulsifiers and gluten, affecting protein polymerization  
31 during baking. CONCLUSION: Bubbles progress during baking allowed discriminate  
32 among emulsifiers, which could explain their performance in breadmaking.

33

34 **Keywords:** emulsifier, image analysis, bubble, dough aeration, bread, crumb

35

## 36 **Introduction**

37 Bakery products are extensively consumed worldwide due to their nutritional and  
38 physical characteristics. <sup>1</sup> Among the diversity of bakery products obtained from either  
39 different raw ingredients or making processes, the most appreciated products are the  
40 sponge baked wheat bread, with low density and soft crumb. In the course of flour and  
41 water mixing, gluten formation and aeration brought about during kneading will be  
42 responsible of the subsequent cellular structure of the baked bread. <sup>2</sup> Air incorporated  
43 into the dough during mixing must be kept through the breadmaking process to attain  
44 low density breads. Bread contains about 70% of gas that comes from the initial  
45 aeration and the fermentation, both are important stages to take into account during the  
46 making process. <sup>3</sup> Because of that air bubbles incorporation during mixing have been  
47 the focus of many studies that stated the influence of mixer type and mixing time, <sup>4,5</sup>  
48 besides the important role of ingredients. <sup>6,7</sup> Certainly, the progress of those initial  
49 nuclei bubbles throughout fermentation when carbon dioxide is generated <sup>8</sup> and final  
50 expansion of the gases occluded into the bubbles during baking determines the diversity  
51 of cellular structures encountered on bread crumbs. <sup>3</sup> Bubbles are very fragile and  
52 whatever changes in their number and size will have a direct impact on the internal  
53 crumb structure. <sup>9</sup>

54 Nowadays, large-scale production and consumers demand for higher quality,  
55 homogeneity and longer shelf life that have been achieved with the use of processing  
56 aids such as enzymes, hydrocolloids, emulsifiers, etc. to adjust doughs properties. These  
57 additives are essentials for improving dough properties and final quality of fresh  
58 product. <sup>10</sup> Specifically, emulsifiers are active surfactant composites used in  
59 breadmaking for their ability to stabilize dough, a thermodynamically unstable system,  
60 through their interactions with gluten proteins. <sup>11</sup> During mixing, the use of emulsifiers

61 increases the strength and the extensibility of the dough; in the fermentation stage they  
62 improve gas retention and avoid dough collapse,<sup>11, 12</sup> leading to softer bread crumbs,<sup>13</sup>  
63 although their effect is greatly dependent on the wheat flour protein content<sup>14</sup> and  
64 proofing duration.<sup>15</sup> Those studies confirmed the effect of different emulsifiers in  
65 breadmaking processes, specifically in improving the internal structure of bread.<sup>16</sup> In  
66 spite of the knowledge acquired on emulsifiers action during breadmaking, they are still  
67 attracting research due to there is still much unknown to explain their functionality  
68 despite their chemistry diversity. For instance, despite the impact of dough aeration into  
69 bread crumb features, there is no information about the role of emulsifiers on dough  
70 aeration and the bubbles number and size along the process. To understand the role of  
71 emulsifiers on determining the cellular structure of bread crumb, the main objective of  
72 this study was to assess the amount of gas occluded into the dough and bread along  
73 bread making and how several emulsifiers with diverse chemical structure affected the  
74 bubble size distribution.

75

## 76 **Materials and methods**

77 Breadmaking wheat flour was supplied by Harinera La Meta (Lleida, Spain) and  
78 compressed yeast by (DHW Europe, Germany). The selected emulsifiers included:  
79 diacetyl tartaric acid ester of monoglycerides (DATEM), sodium stearoyl lactylate  
80 (SSL) and distilled monoglyceride (with potassium citrate added) with two different  
81 particle sizes 45 microns (DMG-45) and 75 microns (DMG-75), which were provided  
82 by Danisco (Grindsted, Denmark), defatted hydrolyzed sunflower lecithin (Tricalcium  
83 phosphate) from Lasenor (Barcelona, Spain), and Polyglycerol esters of fatty acids  
84 containing polysorbate 80 (PGEF) from Palsgaard (Juelsminde, Denmark).

85

86 *Gas bubbles during fermentation and baking*

87 A very hydrated dough recipe containing wheat flour, water (900 ml kg<sup>-1</sup> based on  
88 wheat flour weight) and 10 g kg<sup>-1</sup> compressed yeast was used. Emulsifiers were added  
89 at 5 g kg<sup>-1</sup> (f.b.) whenever tested. Ingredients were mixed during 3 minutes at 328 rpm  
90 in a mixer (RZR-1 Heidolph, Schwabach, Germany).

91 The gas released and dough development characteristics during fermentation were  
92 recorded using the Rheofermentometer F3 (Chopin, France), slightly modifying the  
93 instructions given by supplier. Briefly, hydrated dough (315 g) were confined in a glass  
94 recipient. The tests were performed on dough at 30 K for 3 hours, with a slight  
95 cylindrical weight. Registered parameters included: Hm (mm), maximum dough  
96 fermentation height; T1, the time (min) at which Hm is attained; H'm (mm) maximum  
97 height of gaseous release; T'1, the time (min) at which H'm is reached; Tx, the time  
98 (min) at which gas starts to escape from the dough, thus when porosity of dough  
99 develops. All determinations were made at least in duplicate, and the average values  
100 were adopted.

101 A microscope was used to follow bubble changes of dough during baking as previously  
102 described Rodríguez-García, Salvador and Hernando <sup>17</sup> For that purpose, doughs were  
103 prepared as described before but without the addition of yeast to follow behavior of  
104 bubbles from air incorporation. Microbaking was performed using a system controller  
105 unit for heating and freezing stages (Analysa-LTS350, Linkam, Surrey, UK) mounted  
106 under the lens of a light microscope (Nikon ECLIPSE 80i, Nikon Co., Ltd., Tokyo,  
107 Japan). The temperature profile settings were from 30 K to 105 K increasing at 1.5 K  
108 min<sup>-1</sup>. Samples were captured at ×4 magnification (objective lens ×4/0.13∞/- WD 17.1,

109 Nikon). During microbaking, a video film was recorded with an attached camera  
110 (ExWaveHAD, model no. DXC-190) and images were acquired every 10 K. The  
111 analysis software (Linksys 32, Linkam) was directly interfaced with the microscope,  
112 enabling temperature control and image recording control. Duplicates were recorded.  
113 The number, size and distribution of the bubbles in the dough were analyzed using the  
114 ImageJ software (National Institutes of Health, Bethesda, MD, USA).

115

### 116 *Bread making and characterization*

117 A scale-down breadmaking method was carried out<sup>18</sup> to identify the emulsifiers effect.  
118 Recipes were prepared as described before, and then four grams of dough were placed  
119 in previously oiled cylindrical glass molds (17 mm x 300 mm, diameter x height). They  
120 were fermented for 100 min at 30 K and finally baked at 130 K for 11 min. Two batches  
121 were run for each sample.

122 Texture profile analysis of crumbs was carried out in a TA-XTPlus (Stable Micro  
123 Systems, Surrey, UK). A 10 mm thick slices were compressed twice with a 0.6 mm  
124 diameter probe up to 50% at 1 mm s<sup>-1</sup> speed. The registered parameters were crumb  
125 hardness (g), springiness, cohesiveness, chewiness (g) and resilience. In order to study  
126 cell crumb distribution and morphogeometric characteristics of the loaves, both cross  
127 and longitudinal sections of breads were captured using a scanner (HP Scanjet G3110,  
128 Hewlett-Packard, USA) with 600 dpi resolution. The 2D area and perimeter of  
129 longitudinal section was assessed using ImageJ software. The same software was used  
130 to analyze the cell crumb distribution in 10x10 mm crumb cross-sections. Image section  
131 was improved by splitting RGB channels and selecting the channel with greater contrast  
132 between background and object. Finally, Otsu algorithm (predefined by the software)

133 was applied to convert image into a binary image and particle analysis of the image was  
134 carried out. The parameters assessed were cell/cm<sup>2</sup>, mean area (mm<sup>2</sup>) and circularity  
135 (from 0, rectangle, up to 1, perfect circle). Six slices were used for each determination.

136

### 137 *Statistical analysis*

138 Experimental data were statistically analyzed by analysis of variance (ANOVA) using  
139 Statgraphics Centurion XVI.I 16.1 software (Statistical Graphics Corporation, UK), to  
140 identify significant differences among them. Cluster analysis and principal components  
141 (PCA) were also performed to discriminate among emulsifiers with the tested variables.

142

## 143 **Results and Discussion**

### 144 **Dough development and gaseous release characteristics**

145 To evaluate the action of diverse emulsifiers on dough performance during  
146 breadmaking, very hydrated doughs were used. The effect of emulsifiers on gas  
147 retention during dough fermentation was recorded in the rheofermentometer plots  
148 (Figure 1). After an initial elapsed time, a steady increase of dough volume was  
149 displayed, but certain variation was observed in the presence of emulsifiers (Figure 1a).  
150 Lecithin and PGEF delayed the onset of volume increase compared to the control and  
151 the other emulsifiers. All emulsifiers increased the proofing rate, calculated as the initial  
152 slope of dough volume increase (Table 1). The maximum dough development (Hm)  
153 reached in the presence of the emulsifiers was higher than the one observed in the  
154 control dough, being greater in the case of PGEF (34.6 mm), followed by SSL and  
155 DMG-75 (33.0 mm and 32.4 mm, respectively). The presence of polysorbate blended

156 with the PGEF might contribute to the high volume obtained due to its action as  
157 dispersing agent. This result agrees with those obtained by Gómez et al., Gómez, del  
158 Real, Rosell, Ronda, Blanco and Caballero.<sup>19</sup> where polysorbate addition to low  
159 hydrated doughs led to higher dough volumes than other emulsifiers as DATEM and  
160 SSL. Nevertheless, the time (T1) required to reach the maximum dough development  
161 was higher in the presence of emulsifiers than in the control, confirming that emulsifiers  
162 are much more effective when longer dough fermentations are applied.<sup>19</sup> This  
163 improvement has been ascribed to the emulsifiers ability for strengthening the gluten  
164 network, increasing dough extensibility<sup>19</sup> and dough volume,<sup>16</sup> which in turn was  
165 attributed to the formation of aggregates with gluten proteins.<sup>20</sup> However, that effect  
166 cannot be explained only by the emulsifier chemical structure, given that distilled  
167 monoglycerides with different particle size (DMG-45 and DMG-75) produced different  
168 responses. Dough stability during fermentation was greatly dependent on the emulsifier  
169 tested, and only lecithin and DMG-45 extended the stability of the dough longer than  
170 the control.

171 Regarding the gas production during fermentation (Figure 1b), the most evident effect  
172 was the decrease in the initial CO<sub>2</sub> production when emulsifiers were present. It seems  
173 that emulsifiers, independently of their chemical structure, affected the initial release of  
174 carbon dioxide. Taking into account that no sugar was added in the recipe, possible  
175 explanations could be related to either some interactions between emulsifiers and the  
176 free sugars, available in the flour for proofing, that decrease their readiness for the yeast  
177 or due to physical constraints derived of the more ordered and stronger protein structure  
178 in the presence of emulsifiers.<sup>21</sup> As the proofing progresses, main difference was  
179 observed during the last hour of fermentation when a decrease on the CO<sub>2</sub> production  
180 was observed, due to dough permeability to gas in some of the doughs. Doughs with

181 DMG-45, DMG-75 and lecithin showed greater permeability than the control, which  
182 resulted in a decrease of the ability to retain CO<sub>2</sub> at the end of the fermentation. The  
183 highest CO<sub>2</sub> production was with DATEM addition.

184

### 185 **Microscopy and analysis image of simulated microbaking**

186 The ability of the emulsifiers to stabilize the gas bubbles, incorporated into the doughs  
187 during mixing, was continuously monitored under a microscope. Very hydrated doughs  
188 were subjected to a steady temperature increase to simulate the baking process and  
189 consequently the capacity of the dough to hold the gas. Turbin-Orger, Boller, Chaunier,  
190 Chiron, Della Valle and Réguerre<sup>22</sup> suggested that the liquid fraction present in the  
191 dough influence the cellular structure by affecting the connectivity of bubbles and their  
192 possible coalescence. In this study, very hydrated doughs were used to discard the  
193 possible interference of liquid effect. The captured images of doughs along temperature  
194 increase are shown in Figure 2. Initially, differences in the structure of the doughs were  
195 barely visible. Junge, Hosney and Varriano-Marston<sup>23</sup> reported that emulsifiers  
196 increase the incorporation of gas bubbles during mixing, but they did not find  
197 modifications during the baking stage. However, dough images (Figure 2) showed  
198 progressive changes with the temperature increase and major changes were observed  
199 when reaching 70 K. Babin, Della Valle, Chiron, Cloetens, Hoszowska, Pernot,  
200 Réguerre, Salvo and Dendievel<sup>24</sup> reported that the cell structure stabilization occurs  
201 with the temperature range 50–70 K when main changes associated to starch granule  
202 swelling and gluten cross-linking are produced. In all cases, the bubble size augmented  
203 as the temperature increased and their number and size were dependent on the type of  
204 emulsifier. The most important differences were observed when using DMG-75: bigger  
205 bubbles were observed at low temperature (40 K) if compared to the doughs prepared

206 with the other emulsifiers, and these bubbles were really big at 70 K, giving place to the  
207 biggest bubbles at 100 K.

208 Quantitative analysis of the bubbles distribution and size is shown in Figure 3, where  
209 distributions were ordered from smaller to larger bubble width when temperature  
210 increased. In all the samples, the addition of emulsifiers increased the number of  
211 bubbles incorporated during mixing if compared to control, which may be due to the  
212 lower surface tension induced by the addition of emulsifiers. Kokelaar, Garritsen and  
213 Prins <sup>25</sup> showed that addition of some emulsifiers as SSL and DATEM originated more  
214 and smaller bubbles during mixing, because of the lower surface tension of dough  
215 inducing the subdivision of the entrapped air bubbles. When comparing the doughs  
216 prepared with the different emulsifiers (Figure 3), the dough formulated with PGEF  
217 presented greater incorporation of air bubbles during mixing, as the diagram  
218 corresponding to this emulsifier shows greater frequency of bubbles at the beginning of  
219 the micro baking process. Through temperature rise, all the samples, including control,  
220 showed an increase in the amount of detected bubbles, and bubbles size distribution  
221 became more heterogeneous due to expansion and interaction of the bubbles. The  
222 doughs prepared with DATEM, Lecithin and DMG- 45 presented a frequency  
223 distribution similar to that obtained for the control dough; in fact, the size of the bubbles  
224 increased in a uniform, controlled way (Figure 2). All these doughs showed small  
225 bubbles at low temperatures and a tendency to regular distribution of bubbles during  
226 heating; moreover, bubbles exceeding 120.000  $\mu\text{m}^2$  were not generally detected  
227 regardless of the heating temperature. Nevertheless, the samples prepared with SSL,  
228 PGEF and DMG-75 exhibited bigger bubbles, over 120.000  $\mu\text{m}^2$ . Specifically, DMG-75  
229 dough diagram presented very big bubbles, which continued interacting and coalescing  
230 even at 100 K. When temperature reached 100 K the samples containing SSL, PGEF

231 and DMG-75 presented coarser distribution of bubbles, while DATEM, DMG-45 and  
232 lecithin had more bubbles but smaller ones. When baking temperature rises, the bubbles  
233 expand increasing the coalescence due to Ostwald maturation,<sup>26</sup> where big bubbles  
234 grow up at the expense of small ones, consequently there is an increase in its size. With  
235 the addition of the emulsifiers, this phenomenon often decreased, due to the  
236 stabilization of the interface.<sup>27</sup> However, in the present work, it can be observed that  
237 depending on the emulsifier used in the dough formulation, the expansion of bubbles is  
238 controlled in a different way, being DATEM, DMG-45 and lecithin more effective for  
239 controlling this mechanism. It must be stressed that besides the different chemical  
240 structure of the emulsifiers, their physical structure must be considered, since DMG-45  
241 and DMG-75 induced different bubble stabilization.

242

### 243 **Image digital analysis and texture profile of breads**

244 The effect of emulsifier addition on technological characteristics is summarized in  
245 Table 2. Compared to the control, significantly smaller longitudinal area was produced  
246 by PGEF addition, which meant a reduction in the size of the loaves. The rest of the  
247 emulsifiers did not significantly modify this parameter. Previous studies reported that  
248 adding SSL and DATEM resulted in higher area and volume of breads, due to the  
249 increase in dough aeration and volume.<sup>28,29</sup> Probably the use of high hydrated doughs  
250 is responsible for the differences with previous studies. In addition, a negative  
251 correlation ( $r=-0.8754$ ) was observed between the longitudinal area of the small scale  
252 breads and the maximum height of the proofed dough (Hm). This correlation indicated  
253 that dough volume increased during fermentation with emulsifiers addition but likely  
254 they did not confer enough resistance to improve final volume. Likewise, no significant

255 differences were found in the longitudinal perimeter, except with DATEM and PGEF  
256 that gave smaller values.

257

258 The analysis of the bread cross section revealed significant differences in the number of  
259 gas cells (cell cm<sup>-2</sup>) and mean cells area (mm<sup>2</sup>) on account of emulsifiers addition  
260 (Table 2, Figure 4). The more number of cells, the less mean cell area and vice versa.  
261 DMG-45, DMG-75, DATEM and Lecithin showed greater cell number with smaller  
262 area than the control. In the case of distilled monoglycerides emulsifier (DM) with  
263 different particle size (DMG-45 and DMG-75), no significant difference was observed  
264 in these parameters, showing that particle size did not affect the cell number and area.  
265 Emulsifiers did not induce a significant effect on the circularity compared to the control.  
266 However, significant differences between DMG-75 (0.60) and PGEF (0.74) were found.  
267 Perfect circularity is difficult to obtain in bread, due to the pressure differences in the  
268 gas bubbles and changes occurred during process.<sup>25</sup>

269 All samples showed significant differences in all texture parameters compared to  
270 control (Table 2). With the hydrated recipe used very soft crumbs were obtained, and  
271 emulsifiers increased the crumb hardness although variation ranged from 79 to 100 g.  
272 The highest hardness was obtained with the PGEF, followed by SSL and DATEM.  
273 DMG-45, DMG-75 and lecithin were the emulsifiers that less rise the crumb hardness.  
274 Hardness showed a strong positive correlation ( $r=0.9373$ ) with the maximum height of  
275 dough (Hm) during fermentation, but that contrasts with results obtained when optimum  
276 hydration of wheat flour (500-600 ml kg<sup>-1</sup>) was used.<sup>12</sup> Dough hydration affects the size  
277 of the bubbles diameter,<sup>30</sup> leading to higher bubbles, but since all recipes were prepared  
278 with the same hydration it should be expected no additional effect due to the liquid  
279 phase. Considering the high number of smaller cells mostly found in the breads

280 containing emulsifiers, the hardness increase must respond to the thickness of the cell  
281 walls. Therefore, in this study the interaction of emulsifiers with proteins and starch  
282 leading to the cell walls had greater impact on texture than the bubbles feature. It has  
283 been previously reported that a higher degree of gluten polymerization during baking  
284 results in higher firmness of the baked products.<sup>11</sup> At the same time, emulsifiers, like  
285 SSL or DATEM interact with gluten, changing the solubilization of polymeric  
286 aggregates and that interaction was dependent on the type of emulsifier,<sup>20</sup> particularly  
287 SSL reduces the incorporation of gliadins into the gluten network having a direct effect  
288 on the subsequent polymerization during baking,<sup>11</sup> and in turn affecting crumb  
289 firmness. Therefore, at the level of hydration used in the present study, emulsifiers  
290 contributed to increase dough aeration and in consequence the number of gas cells in the  
291 crumb, but simultaneously their interaction with gluten changed the proteins  
292 polymerization during baking affecting cell walls thickness and in turn crumb firmness.

293 Considering the other texture parameters, chewiness was significantly higher in the  
294 samples with emulsifiers, except with DMG-75, than in the control, and resilience  
295 decreased especially in samples with distilled monoglycerides (DMG-45 and DMG-75),  
296 which again differed than the previously reported with optimum hydrated doughs,<sup>12</sup>  
297 confirming the important role of water on the dough aeration and crumb texture.

298

### 299 **Statistical analysis**

300 In order to understand the effect produced by the emulsifiers and the differences or  
301 correlations between them, a cluster analysis (Figure 5) and an analysis of principal  
302 components (Figure 6) were carry out. Cluster analysis showed the discrimination  
303 between breads containing emulsifiers and the control by combining the two

304 observations that were closest to form the groups. Three well differentiated groups were  
305 drawn, with the control and DMG-75 being more separated from the rest of the samples.  
306 The other emulsifiers were closer in relation to the analyzed variables, evidencing more  
307 similar effects in the doughs and final product. According to their performance with  
308 dough and bread, the closest emulsifiers were DATEM and lecithin, followed by DMG-  
309 45, SSL and finally PGEF. In this study, the closeness observed between lecithin and  
310 DATEM was attributed to the production treatment of lecithin that included a  
311 hydrolysis stage, thus it behaves as a monoglyceride despite of being from a  
312 diglyceride.

313 Principal component analysis (PCA) of the experimental data obtained containing  
314 emulsifiers resulted in two principal components explaining 48.3 and 22.0 % of the data  
315 variation (Figure 6). Thus, the model explained 70.3% of the total variation in data. The  
316 first principal component weight (PC1) was defined by the longitudinal 2D area, the  
317 longitudinal 2D perimeter and cell  $\text{cm}^{-2}$  in the positive axis, and on the negative axis  
318 were located resilience, cohesiveness, springiness, Tx and mean cell area. Component 2  
319 (PC2) was defined by T1, bubbles  $\text{cm}^{-2}$ , longitudinal 2D area, H'm and the mean bubble  
320 area. DATEM, Lecithin, SSL, and PGEF were found in the negative PC1 component  
321 where the majority of dough and bread responses were located. As shown in cluster  
322 analysis (Figure 5), DMG-75 was the furthest emulsifier attending to its experimental  
323 responses, particularly longitudinal 2D area and perimeter, and H'm. Results obtained  
324 from DATEM and Lecithin were explained due to responses to chewiness,  
325 cohesiveness, resilience, cell circularity and Tx. However, PGEF and SSL, adjacent in  
326 cluster analysis, were related with the mean cell area, hardness, T'1 and Hm.  
327 Eventually, DMG-45 position was related to T1 and the number of gas cells  $\text{cm}^{-2}$ .  
328 Overall, emulsifiers could be grouped into four categories attending to the responses

329 obtained with dough and bread performance. In the first group, DATEM and Lecithin  
330 due to their effect on crumb texture and dough permeability; second group would  
331 include SSL and PGEF that showed bigger bubbles, with less and bigger gas cells and  
332 higher crumb hardness; third group with an intermediate behavior respect to the  
333 previous ones DMG-45; and finally DMG-75, with a more distant behavior than the  
334 control, which led to big bubbles.

335

## 336 **CONCLUSIONS**

337 Emulsifiers role on the progress of bubbles during proofing and baking was evaluated.  
338 Emulsifiers showed different functionality that was attributed to their diverse chemical  
339 structure and also physical characteristics (particle size). Furthermore, results shown  
340 that emulsifiers functionality was dependent on the dough hydration. All emulsifiers  
341 studied, increased the maximum dough volume during proofing, but showing different  
342 effect on dough permeability or ability to retain CO<sub>2</sub>. Digital image analysis of the  
343 recorded baking under microscope, allowed quantifying both bubbles number and size  
344 and understand emulsifiers role on aeration. Emulsifiers allowed greater air  
345 incorporation into the dough observing higher number of bubbles, particularly with  
346 PGEF. Major changes in dough occurred at 70 °C when bubble size augmented and  
347 became more heterogeneous, and emulsifiers affected the size and number of bubbles,  
348 with DMG-75 producing the biggest bubbles. In bread, emulsifiers tend to increase the  
349 number of gas cells with lower size, but that gave greater crumb firmness, which  
350 suggested different interactions between emulsifiers and gluten, affecting protein  
351 polymerization during baking. Despite the diverse chemical structure of the emulsifiers,  
352 experimental data following dough proofing and bread features allowed to discriminate  
353 among them.

354

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359

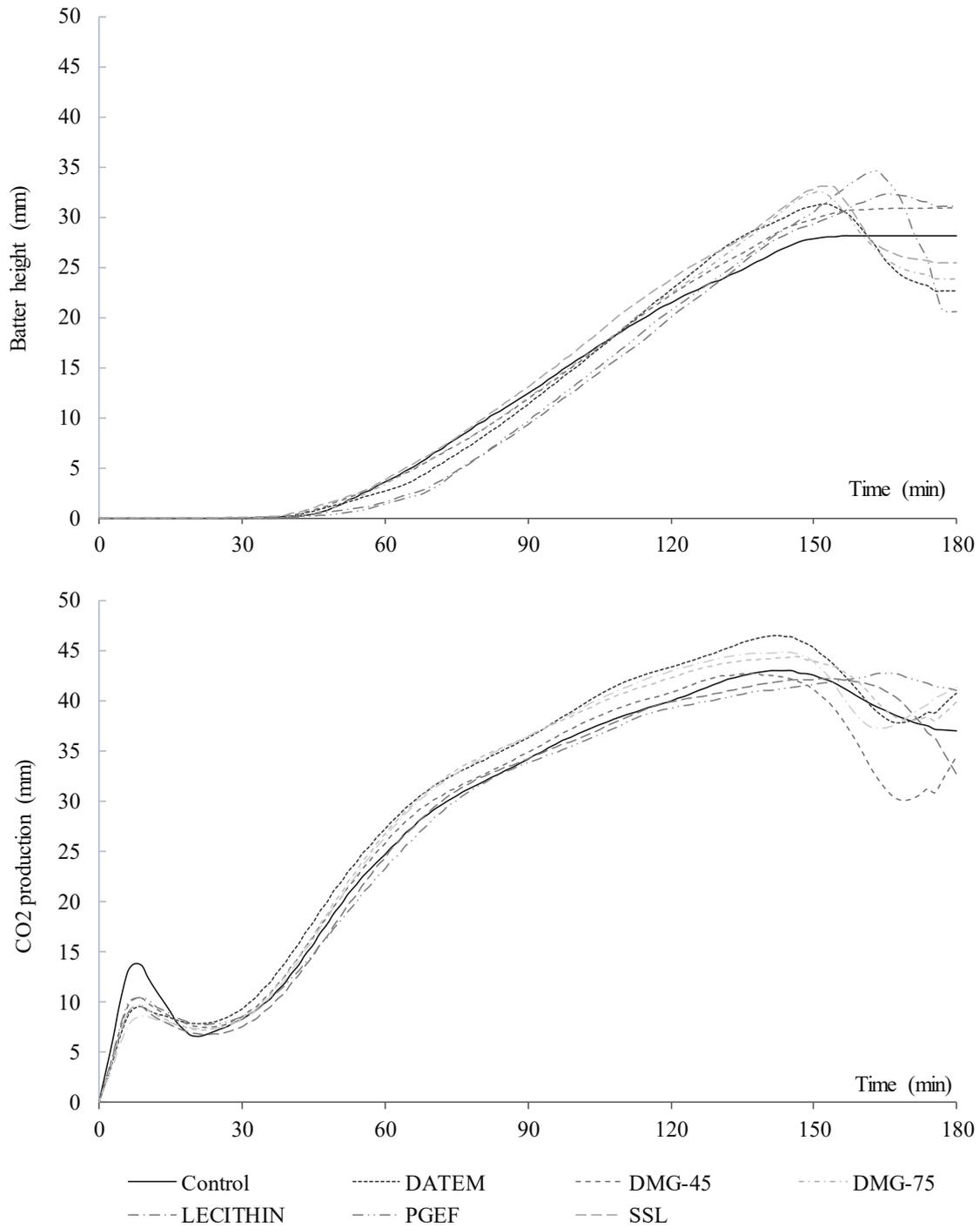
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445

446 **Figure 1.** Rheofermentometer curves consisted of dough development time curves (a)  
 447 and gas release curves (b).

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450 **Table 1.** Analysis of fermentation stage of batters containing emulsifiers by  
 451 reofermentometer.

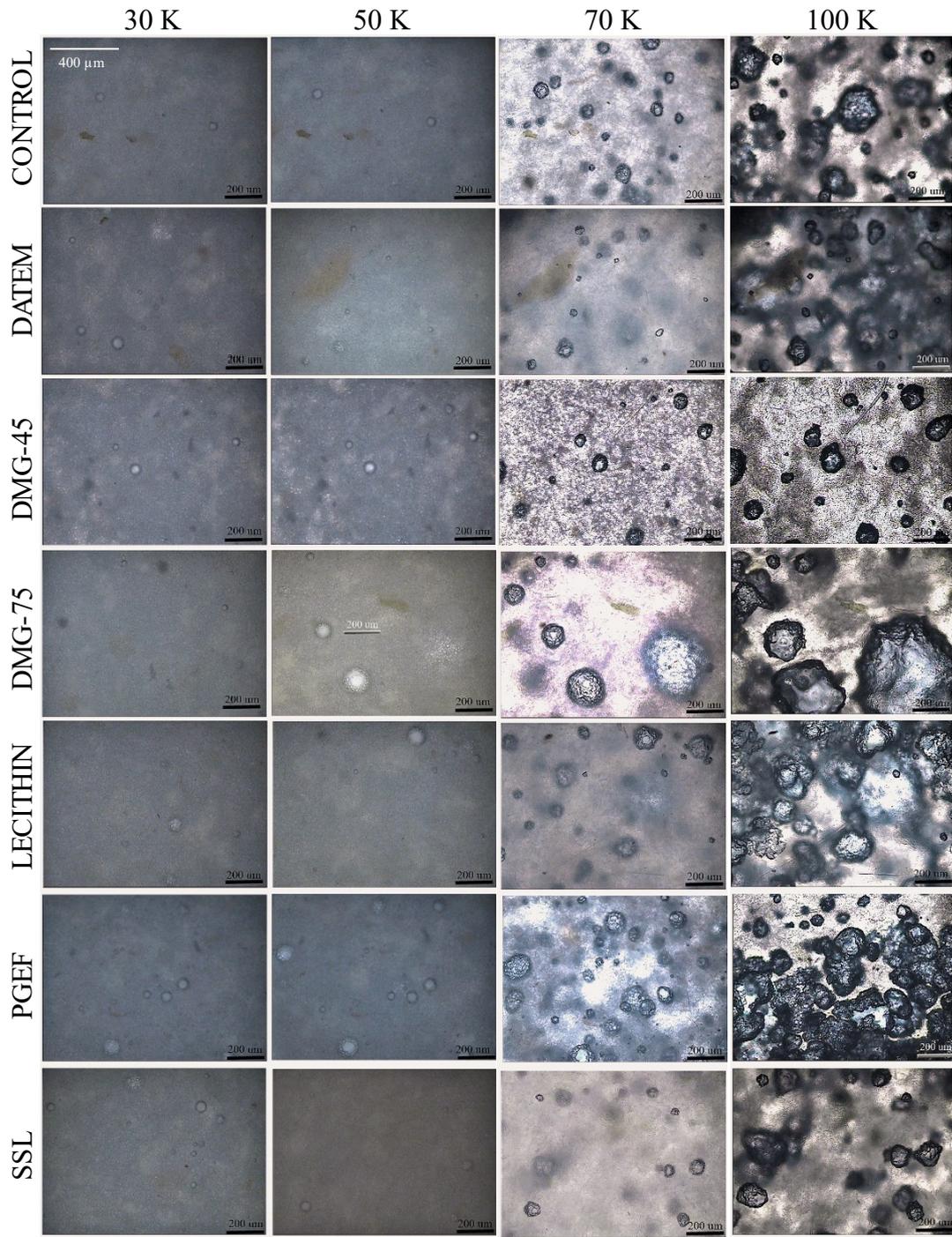
	Dough development			Gas behaviour		
	Hm (mm)	T1 (min)	Proofing rate (%)	H'm (mm)	T'1 (min)	Tx (min)
<b>Control</b>	28.5±0.5 <sup>a</sup>	145±0 <sup>a</sup>	30.79	43.7±1.0 <sup>a</sup>	146±3 <sup>a</sup>	140±1 <sup>b</sup>
<b>DATEM</b>	31.4±1.0 <sup>b</sup>	167±9 <sup>c</sup>	33.44	47.0±0.8 <sup>c</sup>	161±7 <sup>c</sup>	145±2 <sup>c</sup>
<b>DMG-45</b>	31.0±1.0 <sup>b</sup>	169±4 <sup>c</sup>	31.13	43.6±0.1 <sup>a</sup>	140±6 <sup>a</sup>	136±5 <sup>a</sup>
<b>DMG-75</b>	32.4±0.8 <sup>b</sup>	150±2 <sup>b</sup>	31.62	45.8±1.3 <sup>b</sup>	161±8 <sup>c</sup>	134±0 <sup>a</sup>
<b>Lecithin</b>	32.3±0.6 <sup>b</sup>	165±3 <sup>c</sup>	33.77	43.4±0.8 <sup>a</sup>	165±4 <sup>c</sup>	136±3 <sup>a</sup>
<b>PGEF</b>	34.6±0.9 <sup>d</sup>	164±6 <sup>c</sup>	34.77	44.0±0.5 <sup>ab</sup>	176±7 <sup>d</sup>	175±5 <sup>d</sup>
<b>SSL</b>	33.0±0.8 <sup>c</sup>	153±4 <sup>b</sup>	31.46	44.5±0.9 <sup>b</sup>	150±3 <sup>b</sup>	148±4 <sup>c</sup>

452 Mean ± standard deviation. Different letters within the same parameter differ  
 453 significantly ( $P<0.05$ )

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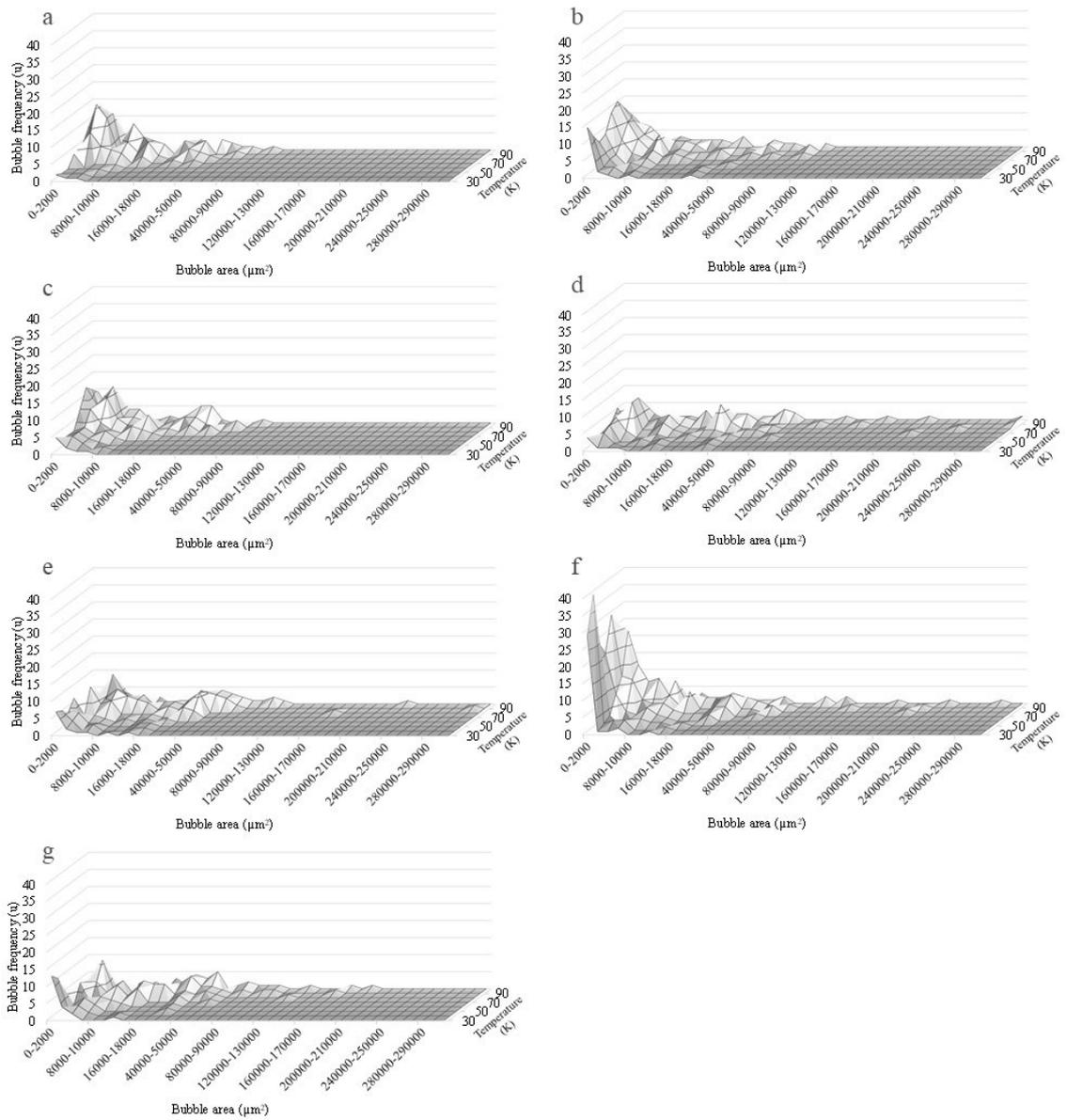
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458 **Figure 2.** Captured images of gas bubbles during simulated microbaking at microscope.



459

460 **Figure 3.** Bubble size distribution during baking for each emulsifier: Control (a),  
 461 DATEM (b), DMG-45 (c), DMG-75 (d), Lecithin (e), PGEF (f) and SSL (g).

462 **Table 2.** Emulsifiers effect on loaves morphogeometrics characteristics, cell crumb distribution and texture profile of small scale breads.

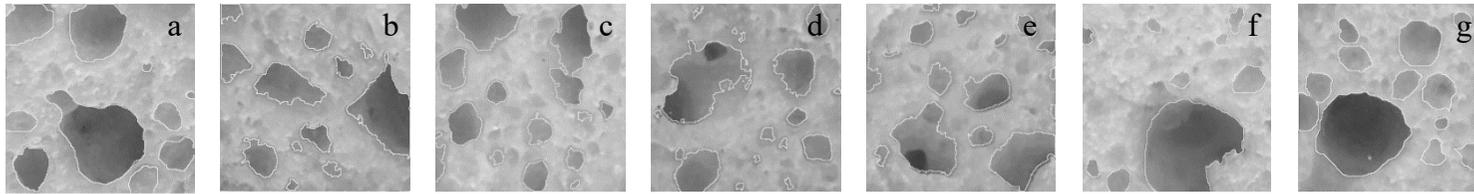
Sample	Control	DATEM	DMG-45	DMG-75	Lecithin	PGEF	SSL
<b>Longitudinal section</b>							
Area (cm <sup>2</sup> )	5.37 ± 0.48 <sup>bc</sup>	5.31 ± 0.24 <sup>bc</sup>	5.09 ± 0.19 <sup>ab</sup>	5.12 ± 0.32 <sup>ab</sup>	5.00 ± 0.24 <sup>ab</sup>	4.95 ± 0.20 <sup>a</sup>	5.50 ± 0.30 <sup>c</sup>
Perimeter (cm)	1.25 ± 0.04 <sup>c</sup>	1.17 ± 0.07 <sup>ab</sup>	1.23 ± 0.05 <sup>bc</sup>	1.25 ± 0.07 <sup>bc</sup>	1.23 ± 0.1 <sup>bc</sup>	1.11 ± 0.04 <sup>a</sup>	1.20 ± 0.08 <sup>bc</sup>
<b>Cross section</b>							
Number of cells cm <sup>-2</sup>	10 ± 2 <sup>a</sup>	13 ± 2 <sup>bc</sup>	15 ± 1 <sup>cd</sup>	16 ± 2 <sup>d</sup>	15 ± 2 <sup>cd</sup>	9 ± 2 <sup>a</sup>	12 ± 3 <sup>ab</sup>
Mean cell area (mm <sup>2</sup> )	3.31 ± 0.81 <sup>b</sup>	1.92 ± 0.89 <sup>a</sup>	1.57 ± 0.47 <sup>a</sup>	1.61 ± 0.40 <sup>a</sup>	1.88 ± 0.34 <sup>a</sup>	3.23 ± 1.70 <sup>b</sup>	2.48 ± 0.48 <sup>ab</sup>
Minimum cell area (mm <sup>2</sup> )	0.15 ± 0.02 <sup>c</sup>	0.14 ± 0.04 <sup>c</sup>	0.10 ± 0.03 <sup>ab</sup>	0.06 ± 0.02 <sup>a</sup>	0.12 ± 0.03 <sup>bc</sup>	0.13 ± 0.02 <sup>bc</sup>	0.12 ± 0.03 <sup>bc</sup>
Maximum cell area (mm <sup>2</sup> )	11.80 ± 3.61 <sup>bc</sup>	7.88 ± 1.79 <sup>ab</sup>	7.22 ± 1.94 <sup>a</sup>	9.12 ± 2.60 <sup>abc</sup>	10.13 ± 2.37 <sup>cd</sup>	17.73 ± 4.77 <sup>d</sup>	13.67 ± 3.65 <sup>abc</sup>
Circularity	0.68 ± 0.16 <sup>ab</sup>	0.68 ± 0.12 <sup>ab</sup>	0.62 ± 0.11 <sup>ab</sup>	0.60 ± 0.11 <sup>a</sup>	0.70 ± 0.06 <sup>ab</sup>	0.74 ± 0.07 <sup>b</sup>	0.72 ± 0.05 <sup>ab</sup>
<b>Crumb texture</b>							
Hardness (g)	55 ± 4 <sup>a</sup>	85 ± 6 <sup>c</sup>	79 ± 3 <sup>b</sup>	79 ± 3 <sup>b</sup>	80 ± 4 <sup>bc</sup>	100 ± 6 <sup>e</sup>	93 ± 5 <sup>c</sup>
Springiness	0.95 ± 0.01 <sup>c</sup>	0.93 ± 0.01 <sup>c</sup>	0.94 ± 0.02 <sup>a</sup>	0.86 ± 0.06 <sup>ab</sup>	0.94 ± 0.04 <sup>c</sup>	0.91 ± 0.02 <sup>bc</sup>	0.85 ± 0.08 <sup>c</sup>
Cohesiveness	0.83 ± 0.03 <sup>c</sup>	0.73 ± 0.02 <sup>b</sup>	0.72 ± 0.03 <sup>a</sup>	0.66 ± 0.05 <sup>a</sup>	0.73 ± 0.04 <sup>b</sup>	0.74 ± 0.03 <sup>b</sup>	0.64 ± 0.04 <sup>b</sup>
Chewiness (g)	35 ± 5 <sup>a</sup>	56 ± 6 <sup>bcd</sup>	53 ± 5 <sup>bc</sup>	47 ± 5 <sup>ab</sup>	56 ± 3 <sup>bc</sup>	58 ± 6 <sup>d</sup>	51 ± 7 <sup>b</sup>
Resilience	0.49 ± 0.03 <sup>c</sup>	0.38 ± 0.01 <sup>b</sup>	0.37 ± 0.04 <sup>a</sup>	0.29 ± 0.03 <sup>a</sup>	0.39 ± 0.04 <sup>b</sup>	0.40 ± 0.03 <sup>b</sup>	0.26 ± 0.01 <sup>b</sup>

463 Mean ± standard deviation. Different letters within the same parameter differ significantly ( $P < 0.05$ )

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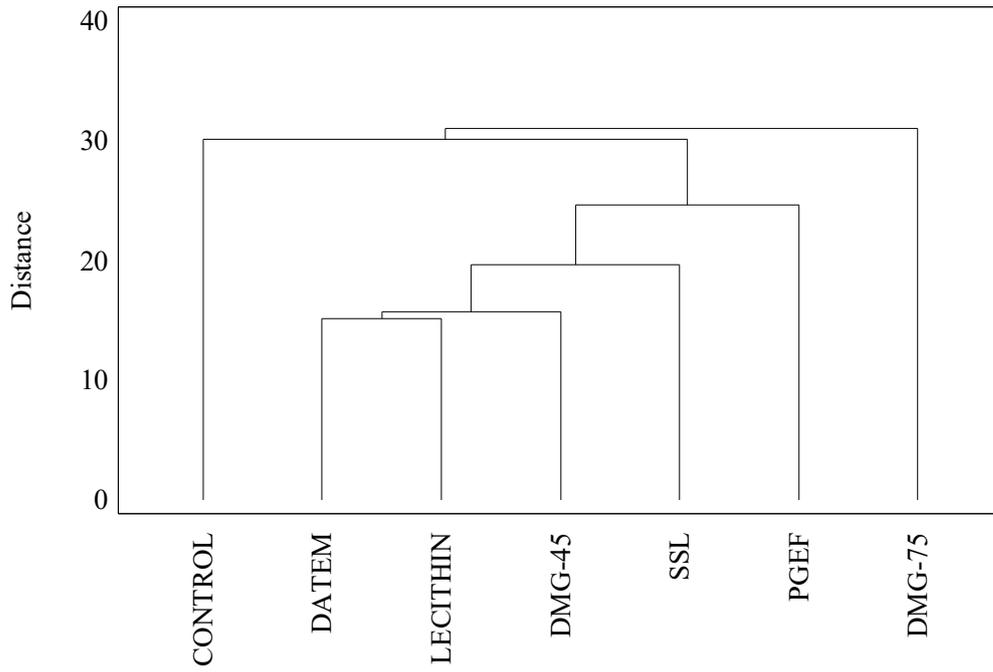
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468 **Figure 4.** Captured images and bubbles count of small scale breads. a: Control, b: DATEM, c:DMG-45, d:DMG-75, e: Lecithin, f: PGEF, g: SSL.



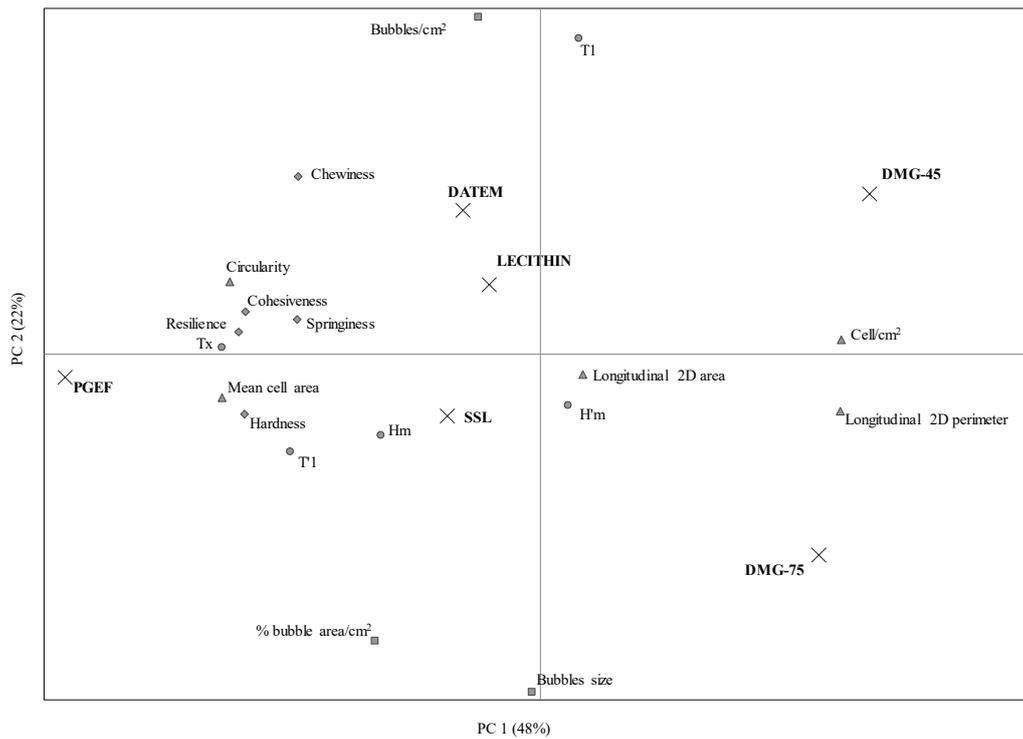
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471 **Figure 5.** Cluster statistical analysis by using closest neighbor method.

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476 **Figure 6.** Score plot from a principal component analysis of the combination of  
 477 components weight (■ simulated microbaking, ◆ texture properties, ●  
 478 rheofermentometer variables and ▲ digital image analysis of breads) and principal  
 479 components (× emulsifiers).

480