

## PAPER

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## Roasting carob flour decreases the capacity to bind glycoconjugates of bile acids

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Accepted 2nd July 2020

DOI: 10.1039/d0fo01158d

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Carob is the fruit obtained from *Ceratonia siliqua* L. and it is a source of bioactive compounds that have been linked to several health promoting effects, including lowering blood cholesterol concentration. The objective of this study was to connect the physicochemical changes of carob flour occurring during roasting with its capacity to bind glycoconjugates of bile acids. Carob flour samples were roasted for different times at 150 °C and chemically characterized by measuring the concentrations of tannins and polyphenols. Data showed that carob flour binds high amounts of bile acids: 732.6 µmol of bound bile acid per g of carob flour which is comparable to the 836.2 µmol per g bound by cholestyramine, a known cholesterol lowering drug. The carob flour ability to bind cholesterol decreases up to 40% during roasting. Data suggested that tannins and insoluble components play a major role in binding bile salts, as a result of hydrophobic interactions.

## Introduction

The carob pod, also known as algarroba, locust pod and St. John's bread, is the fruit obtained from a leguminous evergreen tree (*Ceratonia siliqua* L.). Carob pods are widely cultivated in the Mediterranean area, including Spain, Italy, Portugal, Greece and Morocco as principal producing countries.<sup>1</sup> Its production is estimated at approximately 310 000 tons per year.<sup>2</sup> The carob pod consists of two major parts: approximately 90% pulp and 10% seeds (see Fig. 1). The seeds contribute to more than 60% of the pod market price and are used for the production of locust bean gum, which is obtained from the endosperm.<sup>3</sup> Locust bean gum forms a viscous solution at relatively low concentration and is added as a thickener or stabilizer in many food products. The carob pulp has lower value and, up to now, is mainly used for animal feed and the production of sugar syrup.<sup>4,5</sup> Recently some papers try to valorize carob pulp as source of many valuable components such as dietary fiber, polyphenols, minerals, cyclitols and contains low amounts of fat.<sup>6–10</sup> Consumption of carob pulp was also related to several health promoting effects: it exhibits high antioxidant activity, improves glucose tolerance and attenuates cholesterol levels.<sup>11–16</sup> The chemical composition of the pulp strongly depends on the cultivar, origin and harvesting time.<sup>3,17</sup> Due to the high free sugar content (>48% of which more than 65% sucrose), the pulp is gaining popularity as a natural sweetener in foods, with a flavor profile and

appearance similar to cocoa powder.<sup>18</sup> In this vein, carob pulp is used as an alternative ingredient in some chocolate products leveraging on the peculiarity that it does not contain psychostimulants such as caffeine and theobromine.

After dehulling and deseeding, the pulp is dried in ventilated ovens and mechanically milled to obtain commercial carob flour (see Fig. 1). Optional processing steps include further thermal treatment of the carob flour (*e.g.* roasting) to improve flavour and wettability. Thermal treatment may have major effects on the technological properties of the food products as a result of the Maillard reaction (MR), caramelization reactions and thermal degradation of polyphenols.<sup>19–21</sup> The MR can be used to design ingredients having specific technological functionalities or to target some health function.<sup>22</sup> An interesting health function is the capacity of some food components to reduce blood cholesterol absorption by interfering with the bile acids' enterohepatic cycle. Studying the capacity of carob flour to bind bile acids is physiologically relevant, as high amounts of adsorbed bile acids *in vitro* are related to a decrease in LDL cholesterol *in vivo*.<sup>15,23</sup>

Dongowski, (2007)<sup>24</sup> reported bile acid absorption by dietary fiber-rich preparations, including carob fiber. This paper does not specify which carob flour component is responsible for bile acid absorption; however, considering its composition the role of insoluble tannins can be hypothesized. Up to now it is also not yet investigated which phenolic compounds may influence bile acid binding and which are the potential mechanisms. The physical absorption of steroidal moiety (like the one of bile acids) on the hydrophobic region of condensed polyphenol can be one of the possibilities.<sup>25</sup>

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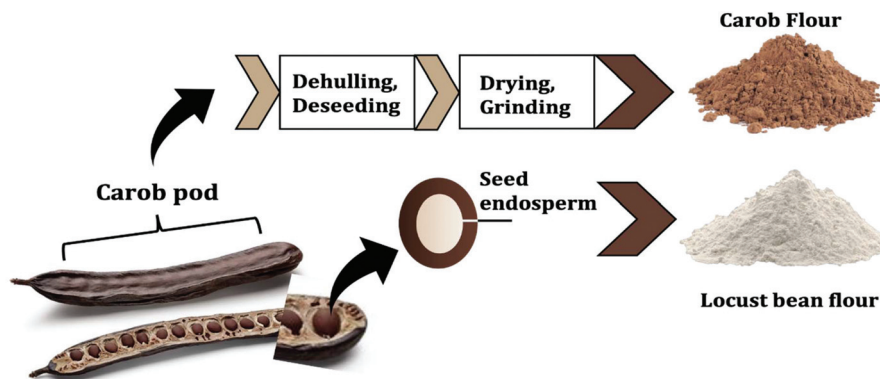


Fig. 1 The different parts of carob pods and their respective production into carob and locust bean flour.

The objective of this study was to connect changes in physicochemical properties of carob flour during roasting with its capacity to bind glycoconjugates of bile acids *in vitro*. Changes in physicochemical properties during roasting were investigated by measuring the difference in solubility, color, viscosity, formation of 5-hydroxymethylfurfural (HMF), total polyphenol and tannin content. In addition, changes in the fraction of bound and free polyphenols were examined.

## Materials and methods

### Material

Carob flour, obtained from BioFertil (C. da Renna, Itali). Standards of phenolic compounds were purchased from Sigma Aldrich (Steinheim, Germany): gallic acid (G-7384) and 5-(hydroxymethyl)furfural (H-40806). Cholestyramine resin and glycodeoxycholic acid (GDCA) were purchased from Sigma, St Louis, MO, USA. The Sigma bile acid assay kit was used to analyze the concentration of free bile acids.

### Roasting process

Carob flour was roasted in a pre-heated oven at five different time intervals: five grams of powder were placed in an aluminium tray and roasted for 0, 5, 10, 30 and 60 minutes at 150 °C.

### Color evaluation of roasted carob flour

Reflectance measurements were performed with a Colorflex EZ (Elscolab, Netherlands). The instrument was calibrated with a supplied white standard traceable to the National Bureau of Standard's perfect white diffuser. CIELAB parameters were calculated for the CIE illuminant D65 and 10° standard observer conditions. Parameters used were  $L^*$  (lightness),  $a^*$  (from green to red),  $b^*$  (from blue to yellow). Universal software 4.1 was used for analysis. All samples were measured in triplicate. To determine whether the flours were roasted homogenously, standard deviations were calculated and the significance of the differences of the means were determined for  $P < 0.05$  using analysis of variance one-way (ANOVA). The browning index of

each sample was also calculated to estimate the color change during roasting, as performed by Medeni, 2001:<sup>26</sup>

$$BI = \frac{[100(x - 0.31)]}{0.17}$$

where

$$x = \frac{(a + 1.75L)}{(5.645L + a - 3.012b)}$$

### Effect of roasting on the solubility of carob flour

For best comparison with the quantification of bound bile acids, the same extraction solvent was used to quantify the insoluble content; carob flour was mixed with a buffer solution (0.1 M phosphate buffer at pH 7.0) to obtain a 20 mg mL<sup>-1</sup> suspension. The suspension was extracted at 37 °C for 15 minutes. Subsequently, the soluble fraction was separated from the insoluble moiety by centrifuging for 6.30 minutes 4700g. Procedure was repeated until a clear solution was obtained. Subsequently, the insoluble fraction was dried at 85 °C for 5 h. The difference in weight before and after extraction was recorded and the procedure was performed in duplicate.

### Determination of changes in viscosity

The rheological behavior was measured by applying an in-house developed method using a rotational rheometer Physica MCR 501 (Anton Paar, Austria) with a titanium double gap concentric cylinder DG26.7 geometry. The geometry has an outer cylinder with internal and external radius of 11.910 mm and 13.801 mm, inner cylinder with an internal and external radius of 12.329 mm and 13.335 mm, and an effective bob length of 40 mm. This geometry measures differences already at a very low viscosity and was therefore most suitable. Samples were allowed to equilibrate at 37 °C for 3 minutes before measurements. A temperature of 37 °C was used to simulate viscosity at body T. A constant shear rate of 20 s<sup>-1</sup> was applied and viscosities were measured for 0.1, 0.5 and 1.0% w/w of low (0 min), medium (30 min) and high (60 min) roasted carob flour. A comparison was made with Locust bean meal, obtained from NatuurlijkNatuurlijk (Ede, Netherlands).



Locust bean meal (from the seeds) was used as comparing material to the carob flour. We wanted to check if viscosity could also influence bile acid binding. With this test, it was found that viscosity of carob flour does not play a role in bile acid binding.

#### Quantification of the amount of free and protein-bound HMF

For quantification of free and protein-bound 5-hydroxymethyl-furfural (HMF), the same method as performed by Fiore *et al.*, 2012<sup>27</sup> was used according to the quantification of free and bound phenolic compounds (as described below). In short, carob flour was extracted using acidified methanol in a 1 : 10 solid to liquid ratio (g mL<sup>-1</sup>) and analyzed with HPLC-UV. Protein-bound HMF was analyzed after 4 h alkaline hydrolysis (2 M NaOH : methanol 50 : 50 v/v).

#### Spectrophotometric quantification of total polyphenols

Carob flour was extracted with acidified methanol (trifluoroacetic acid, pH 2.5) to prevent fast oxidation of oxidative sensitive phenolic compounds. A liquid to solid ratio (mL g<sup>-1</sup>) of 10 : 1 was used. The mixture was extracted using ultrasound-assisted extraction (UAE) at 280 W with an extraction temperature of 30 °C for 30 minutes. Subsequently, the mixture was centrifuged at 15 133g for 5 minutes. The total polyphenol content (TPC) of extracts were determined according to the Folin-Ciocalteu colorimetric method. A calibration curve was obtained using gallic acid in a concentration range of 31.25–1000 µg mL<sup>-1</sup>. Results were expressed as mg per g of gallic acid equivalents. Measurements of sample extracts were performed in duplicate.

#### Estimation of total hydrolysable tannin content

The total hydrolysable tannin content (TTC) (gallo- and ellagitannins) was determined after a reaction with potassium iodide, as reported by Hartzfeld, Forkner, Hunter, & Hagerman, (2002)<sup>28</sup> with some modifications. Carob flour was mixed with 50/50 2 M NaOH/methanol. A solid to liquid ratio (g mL<sup>-1</sup>) of 1 : 10 was used, under strict oxygen free conditions by removing air with nitrogen preliminary to hydrolysis. After 4 h all samples were brought to a pH of 5.5 ± 0.1 with 2 M HCl and 3 M acetate buffer. Subsequently, samples were allowed to react 50 min with potassium iodide at 30 °C. A calibration curve was obtained using tannic acid in a concentration range of 10–100 µg mL<sup>-1</sup>. The absorbance was recorded at 525 nm and results were expressed as tannic acid equivalents (TAE). Measurements of sample extracts were performed in duplicate.

#### Quantification of the amount of free and bound phenolic compounds by HPLC

Since the HPLC data was compared to the data obtained from the spectrophotometric assays, the same extraction solvent was used; carob flour was extracted with acidified methanol (trifluoroacetic acid, pH 2.5) using ultrasound-assisted extraction (UAE) at 280 W with a temperature of 30 °C for 30 minutes. A solid to liquid ratio of 1 : 10 (g mL<sup>-1</sup>) was used. The extract was collected in vials after filtration with 25 µm PTFE membranes

and polyphenols were measured by HPLC-UV (Thermo Scientific Dionex Ultimate 3000 model, Ultimate 3000 pump and Diode 3000 Array Detection). The column used was a Varian Polaris C18A, 150 × 4.6 mm (Phenomenex, UK). Column temperature was maintained at 30 °C. Sample injection was 20 µL and elution was performed at a flowrate of 1 mL min<sup>-1</sup>. For the mobile phase a mixture of water and trifluoroacetic acid (TFA) at a pH of 2.5 was used as a modifier to prevent ionization of phenolic groups (A) and acetonitrile (B). The samples were eluted by the following gradient: 100% A as initial conditions with a linear gradient to 42% B after 20 minutes. 48% A and 42% B were maintained from 20–25 minutes. Detection was performed at 290, 330, 350 and 530 nm. Standards of gallic acid and HMF were prepared as stock solutions with a concentration between 5–200 and 0.1–20 µg mL<sup>-1</sup> respectively. Pyrogallol was identified as a main phenolic compound in carob flour, as already reported by previous authors.<sup>2,6,13</sup>

After the extraction of soluble phenolic compounds, the pellet was rinsed with methanol to eliminate residues and dried under nitrogen gas. After that the pellet was hydrolyzed with 2 M NaOH and methanol (50 : 50 v/v) under nitrogen for 4 h. After alkaline hydrolysis, solutions were neutralized with 2 M HCl and 1 M phosphate buffer at pH 7. The mixture was centrifuged at 20 600g for 10 minutes. The extract was collected in vials after filtration with 25 µm PTFE membranes and analyzed by HPLC-UV as described above.

#### Carob flour bile acid binding capacity

The binding capacity of carob flour was determined according to the kit manufacturer procedure.<sup>29</sup> A buffer solution (0.1 M phosphate buffer at pH 7.0 or 0.1 M acetate buffer pH 5.5) containing 2 mM glycodeoxycholic acid (GDCA) was added to carob flour and cholestyramine (positive control) to obtain a 5 mg mL<sup>-1</sup> suspension. Suspensions were incubated at 37 °C for 2 h under shaking. After incubation, the samples were centrifuged at 10 500g for 5 minutes. Three times 40 µL of sample solution was transferred to new tubes and solutions were diluted to be in a range of 2–150 µM bile acid concentration. A calibration curve was made using a linear range of 10, 20, 40, 80 and 100 µM GDCA solutions. All of the above solutions were prepared in ultrapure water.

Concentrations of GDCA were measured using the Sigma bile acid assay kit. All samples were performed at least in duplicate. The fluorescence intensity was measured at  $\lambda_{\text{ex}} = 530 \text{ nm}$ / $\lambda_{\text{em}} = 585 \text{ nm}$ . Black flat bottom multi well plates with a clear bottom were used for analysis. The calibration curve was used to obtain the conversion factor of GDCA.

The concentration of unbound bile acids of a sample was measured as:

$$[\text{Bile acids}](\mu\text{M}) = \frac{F_{\text{sample}} - F_{\text{blank}}}{F_{\text{standard}} - F_{\text{sample}}} \times 20 \times n$$

where  $F_{\text{sample}}$ ,  $F_{\text{standard}}$  and  $F_{\text{blank}}$  are the fluorescence intensity values of the sample, internal standard and sample blank wells. 20 µM is the effective concentration of the internal stan-



dard (internal standard is 0.25 the volume of the sample).  $n$  is the dilution factor used.

Extraction of untreated carob flour (0 minutes of roasting) was performed using ultrapure water. Fractions were separated by centrifuging for 10 minutes at 4700g. The pellet was rinsed until the liquid fraction became clear. The pellet (IC) and liquid fraction (SC) were both freeze dried (Christ Alpha 2–4 LD plus) and analyzed for the capacity to bind GDCA separately.

### Statistics

Measurements were performed at least in duplicate. Results were expressed as mean values  $\pm$  SD. One-way analysis of variance (ANOVA) was applied to determine significant differences with at least  $P < 0.05$ . Tukey's test was conducted to determine significant differences between all possible pairs of means.

## Results and discussion

### Physicochemical characterization of roasted carob flour

**Color.** Changes in CIE  $L^*a^*b^*$  and browning index (BI) values of carob flour at different roasting times are shown in Table 1. The one-way ANOVA indicated that time significantly ( $p < 0.05$ ) affected the color of carob flour during roasting. The small standard deviations indicate that the flours were roasted homogeneously. The  $L$ -value of carob flour decreased with increased roasting time. The decrease in lightness is associated with the formation of brown pigments in carob flour due to the Maillard reaction and caramelization reactions.<sup>30</sup> The  $a$ - and  $b$  values represent the redness and yellowness of a product. The  $a$ - and  $b$  values initially increased with roasting and decreased after 10 and 5 minutes, respectively. Woo *et al.*, 2015<sup>20</sup> reported that changes in CIE  $\times Lab$  values were related with the formation of MRP's in food, particularly HMF. In general, the decrease in lightness coincided with an increase in BI. The BI is a measure of the purity of brown color that develops during enzymatic and non-enzymatic roasting. However, upon severe roasting conditions (30 min) BI start to decrease. Interestingly, the decrease in BI is not parallel with that of  $L^*$ . This discrepancy might have different causes and can be related to the intrinsic properties of the various MRPs formed during the MR cascade but also related to the breakdown polymerized tannins. This result was in line with the observation made on coffee by Shan, Zzaman, & Yang, 2016.<sup>31</sup>

These authors found a decrease in BI of coffee beans roasted at 250 °C for 20 minutes. Different results were reported by Sahin *et al.*, 2009<sup>32</sup> where increased roasting of carob coincided with an increase in BI only. However in this paper the roasting process was performed directly on the carob kibbles before milling; this would have likely resulted in a less efficient heat transfer respect to the carob flour.<sup>32</sup>

**Water solubility.** Fig. 2 shows the content of insoluble material (% w/w) as a function of roasting time. The decrease in insoluble compounds as a result of roasting clearly show the increase in carob flour solubility determined by roasting treatment significantly ( $P < 0.05$ ). As shown in Table 2 a significant correlation was found between the total tannins (TTC), total gallic acid (TGA), free gallic acid (FGA) and total insoluble content (TIC). It is possible that thermal degradation of (insoluble) cellular structures during roasting led to a decrease in insoluble compounds. Consequently, it led to the solubilization in water of different compounds previously entrapped in the insoluble matrix. In particular, the degradation of insoluble tannins into more soluble phenolic compounds is a well-known phenomenon that can explain this macroscopic observation.<sup>33</sup> We will show later that the amount of total hydrolysable tannins sharply decreased with roasting time. On the other hand, insoluble non-starch polysaccharide fibers (e.g. cellulose and hemicellulose), non-polysaccharide fibers (e.g. lignin) and resistant starch may also have been degraded into more soluble products during roasting, as previously reported on cellulose, starch blends and hydrolytic lignin.<sup>34–36</sup>

**Viscosity.** The viscosity of Locust bean flour increased exponentially with concentration. In contrast, increased concentration of carob flour does not increase viscosity. Also roasting did not alter viscosity. Compared to Locust bean flour, known for its thickening effect, carob flour does not contain galactomannan. Although roasting led to the formation of more soluble products, polysaccharides in carob flour are not able to form a viscous network. This is relevant as the observed changes in the bile salt binding capacity (see below) cannot be attributed to modification of solution viscosity.

**HMF formation.** The formation of total, free and protein-bound HMF as a function of roasting time, is reported Fig. 3, Panel A. The formation of total HMF follows a zero-order model:  $[HMF_{tot}] = 0.01 + 0.0062 \times t$  (see Fig. 3, Panel B.). The unroasted carob flour (0 min) contained a low amount of free HMF probably formed during the manufacturer drying process and increased up to 36 mg kg<sup>−1</sup> at 60 min. Protein-bound

**Table 1** CIE-Lab color determination with  $L^*$  representing the lightness, browning index (BI),  $a^*$  and  $b^*$  for the red-green and blue-yellow color components respectively. Mean values are obtained after 0, 5, 10, 30 and 60 minutes of roasting  $\pm$  Standard deviation. Significant differences between samples were obtained ( $P < 0.05$ )

Roasting time (min)	$L^*$	$a$	$b$	BI
0	55.23 $\pm$ 0.06	11.35 $\pm$ 0.02	26.08 $\pm$ 0.05	76.89 $\pm$ 0.25
5	47.79 $\pm$ 0.05	12.51 $\pm$ 0.03	27.13 $\pm$ 0.10	99.02 $\pm$ 0.54
10	40.98 $\pm$ 0.04	12.87 $\pm$ 0.04	25.12 $\pm$ 0.09	112.26 $\pm$ 0.63
30	34.08 $\pm$ 0.03	11.31 $\pm$ 0.04	20.03 $\pm$ 0.03	108.17 $\pm$ 0.33
60	26.32 $\pm$ 0.03	9.11 $\pm$ 0.07	13.94 $\pm$ 0.03	97.55 $\pm$ 0.26





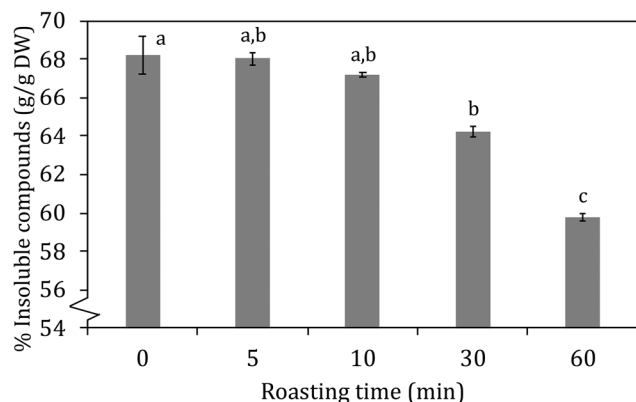


Fig. 2 Insoluble compounds (% w/w) in carob flour as a function of roasting time at 150 °C. Different letters indicate significant differences ( $p < 0.05$ ).

Table 2 Correlation matrix of the chemical properties of roasted carob flour; total tannin content (TTC), total insoluble content (TIC), total and free gallic acid (TGA and FGA)

	TTC	TIC	TGA	FGA
TTC	1			
TIC	0.9872	1		
TGA	0.9845	0.9998	1	
FGA	-0.9363	-0.9734	-0.9768	1

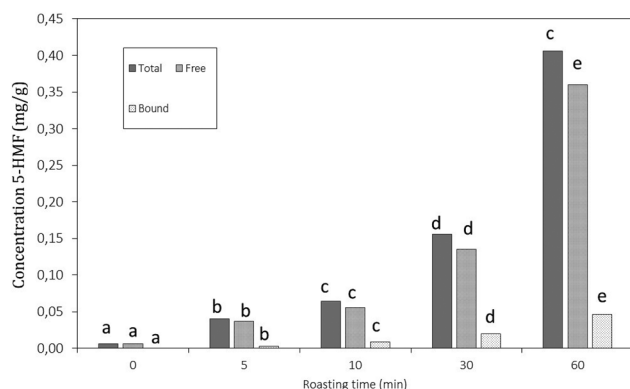


Fig. 3 Concentration of total, free and protein-bound 5-HMF as a function of roasting time. Different letters in the same series indicate significant differences ( $p < 0.05$ ).

HMF was clearly detectable already after 5 minutes of roasting and increased linearly over time being always about 10% of the total. Interestingly, this HMF value is in the range of the reference value reported for honey being 40 mg kg<sup>-1</sup> the maximum HMF content allowed for it.<sup>37</sup> Increased protein-bound HMF is the result of thermal processing and the continuation of the Maillard reaction, when the compound is incorporated in *e.g.* melanoidins. There was also a strong correlation ( $>-0.9$ ) found with  $L^*$ , but not with BI and  $a^-$  and  $b^-$  values. According to Agila & Barringer, 2012<sup>38</sup> there was a strong correlation found

between  $L^*$  value, chroma and the formation of HMF. Similar results in the ratio of protein-bound HMF with HMF-casein solutions and increased binding with heating was found also by Oral, Mortas, Dogan, Sarioglu, & Yazici, 2014.<sup>39</sup>

**Total content of phenolic compounds.** Changes in the total polyphenol content (TPC) and total hydrolysable tannin content (TTC) as a function of roasting time is shown in Fig. 4. Significant changes ( $p < 0.05$ ) in TPC and TTC as mg GAE/TAE per g of carob flour were observed with increased roasting. TPC increased up to 10 minutes of roasting and decreased subsequently. No initial increase was found with TTC during roasting. The decrease in TTC during roasting was in line with the data obtained by quantifying gallic acid. As reported in Fig. 5 gallic acid that is chemically bound to cellular material and tannins decreased upon roasting (from 85.5% at 0 minutes of roasting to 84.2, 83.6, 80.4 and 68.8% with 5, 10, 30 and 60 minutes respectively).<sup>40-42</sup> This decrease coincided with an increase in free gallic acid and the release of phenolic compounds was also observed in grape seeds.<sup>43</sup>

The decreases in total hydrolysable tannins (Fig. 4B) together with the shift of bound towards free gallic acid (Fig. 5) indicated that a thermally induced degradation of gallo- and ellagitannins happened in the carob flour. The

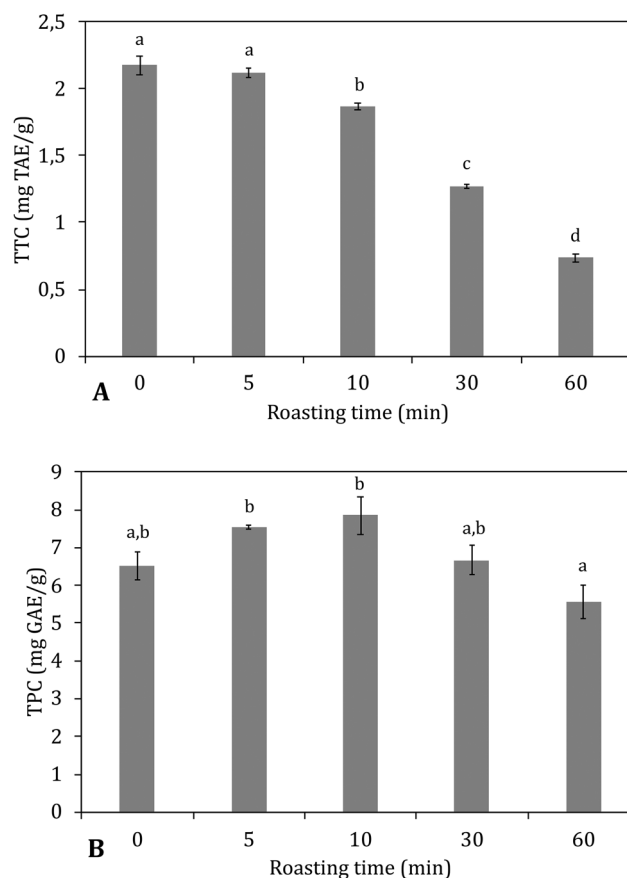
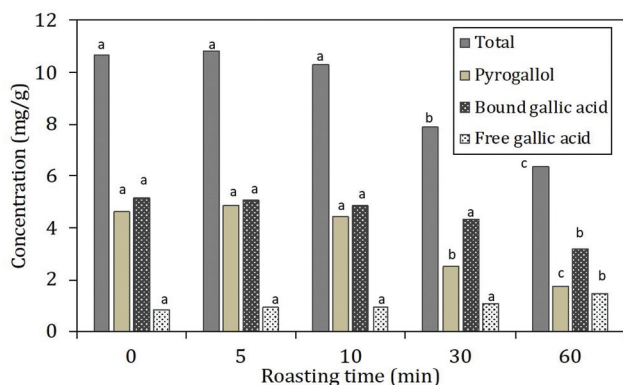


Fig. 4 Changes in the total hydrolysable tannin content (A) and total phenolic content (B) of carob flour obtained after roasting at 150 °C. Different letters indicate significant differences ( $p < 0.05$ ).





**Fig. 5** Phenolic content obtained after ultrasonic extraction with acidified methanol (pH = 2.5). The total, pyrogallol, bound and free gallic acid content were expressed as mg g<sup>-1</sup> of carob flour as a function of roasting. Different letters in the same series indicate significant differences ( $p < 0.05$ ).

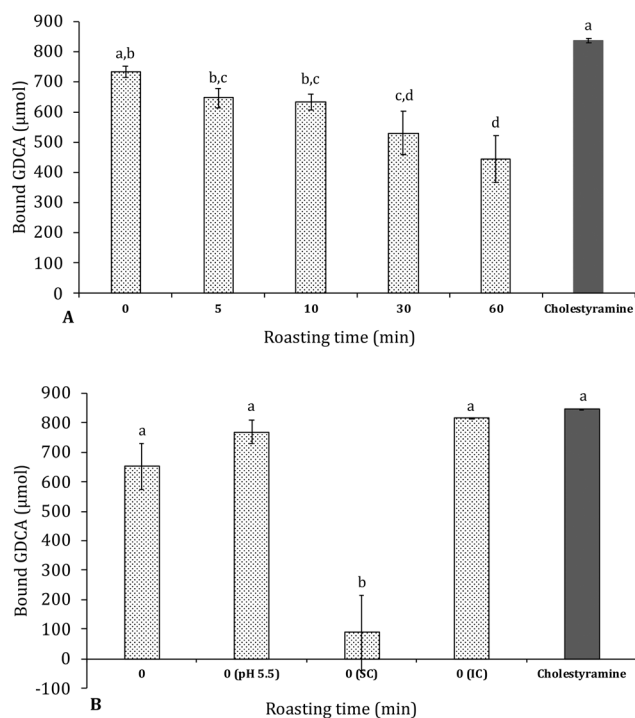
initial increase in TPC after extraction with acidified methanol could be explained by (i) degradation of insoluble and bound phenolics (e.g. tannins and lignin as seen in grape seed<sup>21,44</sup>) during roasting led to the formation of more soluble products, free phenolics and derivatives that were detected by this assay, in line with TTC and bound/free gallic acid. (ii) Formation of MRP's with a phenolic type of structure induced roasting. The TPC measured by the Folin–Ciocalteu method lacks specificity and detects all reducing compounds. The MR in sugar-rich environment favour the formation of reductones, a class of MRPs with a general reducing capacity,<sup>45</sup> which are likely to be detected by the reagent used.

Similar findings were already observed by Şahin *et al.*, 2009,<sup>32</sup> who showed roasting led to increased TPC and a positive correlation between BI and TPC. Papagiannopoulos *et al.*, 2004<sup>9</sup> not only reported an increase in TPC for roasted carob products compared to raw carob kibbles, but also a decrease in the amount of hydrolysable tannins that paralleled an increase in free gallic acid. Finally Tomaz *et al.*, 2019<sup>46</sup> reported a decrease in the tannin content and an increase in the gallic acid concentration with increased processing.

Next to free and bound gallic acid, also pyrogallol was identified by HPLC analysis after extraction of the carob flour samples. The amount of pyrogallol slightly increased at 5 min of roasting and decreased subsequently. The increase in pyrogallol at the onset of roasting is the consequence of thermal decarboxylation of free gallic acid.<sup>47</sup> The total phenolic content decreased after 5 minutes of roasting when decarboxylation reactions ultimately led to the degradation of gallic acid and pyrogallol. These results were in line with the trend of TPC reported in Fig. 4: after 10 minutes of roasting, where the decrease could be explained by the thermal degradation of polyphenols due to the decarboxylation reactions.<sup>48</sup> Moreover, the incorporation of phenolic compounds/phenolic type of MRP's in melanoidins<sup>49</sup> should be considered.

### Carob flour binding to bile acids glycoconjugates

In Fig. 6 the amount of glycodeoxycholic acid (GDCA) bound to the carob powder is shown in comparison to that of cholestyramine, a resin which is a known binder of bile acids and it is used as cholesterol-lowering drug. Carob flour showed a good ability to bind bile acids glycoconjugates. Unroasted carob flour bound 92% of GDCA compared to cholestyramine. Increased roasting resulted in significantly ( $p < 0.05$ ) decreased bile acid binding; from 92 to 81, 79, 67 and 55% after 5, 10, 30, and 60 minutes of roasting, respectively. Decreasing from physiological pH (pH = 7), to pH = 5.5 resulted in an increase in bile acid binding which fail to reach statistical significance (see Fig. 6, Panel B). Bile acids were mainly bound by the insoluble compounds (IC) compared to the soluble compounds (SC) being the latter almost negligible: as shown in Table 3 there is a high correlation between the total bound bile acids (TBBA), tannins (TTC) and insoluble compounds (TIC) as a function of roasting. Relevant factors contributing to the high GDCA binding capacity can be (i) tannins and insoluble compounds in general, which have a large impact on the capacity to bind bile salts, (ii) minerals and (iii) other components that may be involved in the stabilization of binding bile acids such as melanoidins or cellulose. Carob fiber treatment was able to decrease total and LDL



**Fig. 6** Effect of roasting on the amount of bound glycodeoxycholic acid (GDCA) at a concentration of 2 mM at pH = 7 per g of carob flour. In Panel A the effect of roasting on bile acid adsorption is reported while in Panel B the effect of pH of unroasted carob powder (time 0) on bile acid adsorption is shown. The ability of soluble- and insoluble content (IC/SC) were measured separately. Different letters are significantly different  $p < 0.05$ . Cholestyramine represents 100% bile acid adsorption as a positive control.



**Table 3** Correlation matrix of the chemical properties of roasted carob flour and the relation with bile acids; total bound bile acids (TBBA), total tannin content (TTC) and total insoluble content (TIC)

	TBBA	TTC	TIC
TBBA	1		
TTC	0.9697	1	
TIC	0.9480	0.9872	1

cholesterol in hypercholesterolemic patients.<sup>23</sup> Minerals in carob flour could play an important role in enhancing the binding of bile salts, especially as carob flour is rich in calcium and iron.<sup>6,17,50</sup> Other components that could have supported the adsorption of bile acids were cellulose and melanoidins. Ragot, Russell, & Schneeman, 1992<sup>51</sup> reported already that melanoidins from several heated protein–glucose solutions could interact with GDCA. It was observed that especially high molecular weight melanoidins interacted with bile acids. This is in line with data from tannins; especially HMW insoluble tannins are able to interact with bile acids. Finally cellulose could have contributed to support of binding bile acids, although actual values on the capacity to bind bile acids were low.<sup>52,53</sup>

Another paper by Dongowski, (2007)<sup>24</sup> before ours investigated the relation between tannins in carob and the capacity to bind bile acids. They found high amounts of bile acids were bound in a commercial carob fiber preparation and removal of tannins resulted in a sharp decrease in the amount of bound bile acids. This relation was not deeply investigated in the Dongowski's paper; however, our data reported in Table 3 and Fig. 6 clearly suggested that insoluble hydrolysable tannins can play a significant role. When these hydrolysable tannins are broken down by roasting, the degree of bound bile acids drops significantly. Despite the capacity of melanoidins to bind GDCA, roasting carob flour resulted in lower amounts of bound bile acids indicating the ability of insoluble tannins are far better than that of the neo-formed melanoidins.

It was reported that there are three mechanisms explaining the incomplete re-absorption of bile salts into the enterohepatic circulation at the end of the small intestine: the first mechanism explains that (in)soluble dietary fiber increases the barrier properties of the unstirred water layer between micelles and intestinal absorptive cells.<sup>54</sup> Secondly, it was reported that (in)soluble compounds might interact at a molecular level, including hydrophobic interactions.<sup>55</sup> Last, soluble dietary fiber might form a local viscous or gelatinous matrix that entraps the bile salt micelles.<sup>56</sup> All these mechanisms entail an increased loss of bile salts in the feces; therefore, they must be newly synthesized in the liver and this in turn leads to a reduction of the free cholesterol pool.

The probability that the formation of a local viscous or gelatinous network could contribute to lower LDL cholesterol *in vivo* was excluded, since increasing the concentration of carob flour, nor roasting had an effect on viscosity. On the other hand, although bile acids have amphipathic properties, hydrophobic interactions play a major role. Not only because

bile acids mainly bind to insoluble material and tannins in carob flour, but they also bind stronger at lower pH. Similar findings of higher amounts of bound bile acids at lower pH were reported by Drzikova *et al.*, 2005.<sup>55</sup> Camire, Zhao, & Violette, (1993)<sup>57</sup> and Dziedzic, Górecka, Kucharska, & Przybylska, (2012)<sup>58</sup> reported that trihydroxy bile acids (GDCA) had lower adsorption capacities compared to dihydroxy bile acids (cholic acid).

## Conclusions

The objective of this study was to investigate the mechanisms underpinning the cholesterol lowering effect of carob flour and the role of insoluble tannins. To this purpose the influence of roasting on the capacity of carob flour to bind glycoconjugates of bile acids *in vitro* was investigated. We found a high correlation between the capacity of carob flour to adsorb bile acids and their respective tannin and insoluble content. This study supports the evidence that carob has the capacity to bind high amounts of bile salts and that tannins and insoluble compounds are important for interactions between bile salts and carob flour. Roasting decreases the capacity of carob flour to adsorb bile acids (GDCA), mainly as roasting breaks down the HMW tannins in carob flour. Therefore, the bile acid lowering capacity of carob flour is mainly due to hydrophobic interactions exerted by insoluble tannins.

## Abbreviations

LDL	Low density lipoprotein
BI	Browning index
GDCA	glycodeoxycholic acid
TTC	Total hydrolysable tannin content
TPC	Total phenolic content
TFA	Trifluoroacetic acid
UAE	Ultrasound-assisted extraction
IC	Insoluble compounds
SC	Soluble compounds
HMF	5-Hydroxymethylfurfural
TGA	Total gallic acid
FGA	Free gallic acid
TBBA	Total bound bile acids

## Conflicts of interest

There are no conflicts to declare.

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