Foaming properties of soy protein isolate hydrolysates

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Abstract: Tensoactive species obtained by papain hydrolysis of soy protein were characterized structurally and physicochemically, and their foam-forming and -stabilizing capacity studied. Protein structural changes upon reaction ending were correlated with functional and interfacial properties and with the behaviour thereof with varying hydrolysis degree. Two different means of halting hydrolysis -pH reduction (pH=2) and `quick` freezing (-18 ºC), respectively- were studied. Distinct structural changes and associated functional properties were found according to reaction ending conditions. No improvement of foaming properties was found for partially-hydrolyzed isolates subject to freezing at reaction ending - with respect to the starting unhydrolyzed soy protein isolate. In contrast, pH treatment as a means of halting hydrolysis led to a significant enhancement of the foaming properties of soybean protein hydrolysates consistently for all studied hydrolysis degrees (0%, 1.8%, 2.5% and 6%).

Keywords: Soy Protein, Enzymatic Hydrolysis, Foaming Properties

1. Introduction

Soy protein is a protein that is isolated from soya bean. Soya bean (*Glycine max*) is a species of legume family (*Fabaceae*) cultivated for its seeds. Soy protein is made from soybean meal that has been dehulled, defatted and processed into three kinds of high protein commercial products: soy flour, concentrates, and isolates. Soy protein finds a variety of applications in the food industry for its functional properties and his popularity has increased due to its use in health food products. In view of its high digestibility and balanced aminoacid composition, soy protein constitutes a major protein source capable of substituting meat and dairy proteins in the manufacture of food products. As little as 1% of the soy protein production is currently used in the human food industry for the improvement of textural properties; the rest being used in the manufacture of animal feeds [1]. Soy protein is used predominantly as a functional ingredient in view of its gelling, and foam- and emulsion-forming and stabilizing capacity [1]. However, poor foaming properties have been attributed to a highly compact structure of soybean protein -compared with protein from other sources- not liable to adsorption or unfolding at the interface in a way such that it prevents adequate formation of interfacial film [2].

A limited extent of proteolysis has been suggested to have a positive effect on the surface activity and the foaming and emulsifying properties of soy proteins [3]. Previous studies were focused on the modifications resulting from enzymatic hydrolysis of soy proteins using papain and on the relationship between the structural changes undergone during hydrolysis and the functional properties of modified proteins [4], [5]. Nonetheless, recent advances in analytical foam formation and stabilization methods may be used in furthering research on the dependence between surface functional properties and protein structure. In particular, further research on the improvement of functional properties of soy protein at pH above 4 has been suggested with a view to the inclusion of these proteins into suitable applications in food systems [1].

Here, a structural and physicochemical characterization was made of the products of partial papain hydrolysis of soy protein using two different means of halting the reaction. The effects of low-temperature and low-pH ending conditions were thus compared with a view to studying the suitability of hydrolyzed soy protein for food applications. The behaviour of interfacial and foaming properties was correlated with the

structural changes undergone by the soy protein isolates during hydrolysis and according to means of reaction halting.

2. Materials and Methods

2.1. Preparation of Soy Protein Isolates

Soy protein isolates (SPI) were obtained from defatted soybean flour by solubilization in alkaline aqueous medium (pH 8.0), followed by isoelectric precipitation (pH 4.5), precipitate dispersion in alkaline medium (pH 8.0) and drying by liofilization [6].

2.2. Preparation of Soy Protein Hydrolysates

SPIs (at a concentration of c.a. 30 mg/ml) were incubated with papain solution (0.2 mg/ml) in a 4:1 (vol:vol) ratio under agitation in a thermostatized bath at 40ºC. SPIs had a protein content of 90.8 ± 0.4 g/100 g, a water content of 3.50 ± 0.02 g/100 g and a mineral ash content of 5.7 ± 0.02 g/100 g. Papain purchased from Sigma contained 28 units/mg –with one enzyme unit hydrolyzing 1.0 μmol of α-N-benzoyl-L-arginine ethyl ester (BAEE) per minute at pH 6.2 at 25°C. Substrate solutions or dispersions were prepared in 0.01M Na_3PO_3 buffer solution of pH 8.0. Different hydrolysis degrees (0%, 1.8%, 2.5 and 6.0 %) were obtained according to different times of reaction. Hydrolysis was halted by means of: 1) pH reduction in the dispersion of soy protein isolate hydrolysates (SPIHs) to pH 2 using 6 N HCl (HpH); and 2) quick freezing of the SPIH dispersion in a bath with ice / NaCl cryogenic mixture at –18 °C (HT). SPIHs thus obtained were subject to liofilization. SPIHs of equal hydrolysis degree were obtained from the same pool of SPIs for either means of reaction halting.

2.3. Hydrolysis Degree Determinations

Hydrolysis degrees (HDegs) were determined by determination of free amino groups by the trinitrobenzenesulphonic acid (TNBS) method as described by [7], following the modifications introduced by [8]. Determinations were made by triplicate measurements using a standard absorbance curve for l-leucine at 420 nm.

2.4. Hydrolysate Composition

The protein content of SPIHs was determined by the method proposed by Lowry [9] using 100 μg/ ml dispersions in 0.01 M Na_3PO_3 solution at pH 8.0. The water content was determined according to AOAC 14004 [10] and the ash content according to AOAC 31013 [10]. Determinations were based on triplicate measurements.

2.5. Electrophoresis

SDS electrophoresis was conducted on 7–15 % denaturating acrylamide gradient gel with 0.375 M Tris-HCl, 1 % (w/v) SDS buffer solution of pH 8.8. Samples were conditioned with 0.125 M Tris-HCl, 20 % (v/v) glycerol, 4 % (w/v) SDS, 0.2 % (w/w) bromophenol blue buffer solution of pH 6.8. The electrophoretic run buffer consisted of 0.025 M Tris-HCl, 0.192 M glycine, 0.1 % (w/v) SDS. The run was conducted at a constant voltage of 90 V applied to two 1.5 mm-thick gels. A Hoefer Scientific Instruments SE 640 electrophoresis unit was used. Protein molecular weight was estimated using a GIBCO BRL® 10064–012 molecular weight standard consisting of 12 genetically engineered proteins in the range of 10 kDa to 120 kDa separated at 10 kDa intervals in addition to another, 200 kDa such protein. Determinations were based on duplicate measurements.

2.6. Surface Hydrophobicity Determinations

1-anilino-8-naphthalene sulphonate (ANS) was used as fluorescent probe [11]. Surface hydrophobicity determinations were based on SPIH dispersions in 0.1 M Na_3PO_3 solution at pH 7.0. Fluorescence measurements were made on a Perkin Elmer 2000 spectrofluorometer, adjusting the relative fluorescence intensity to 80% of the full scale value using 15 μl of 8 mM ANS in 3 ml absolute methanol and 364 nm and 484 nm excitation and emission wavelengths, respectively. Surface hydrophobicity was determined as the initial slope of the relative fluorescence intensity versus protein concentration plot, according to [12]. Determinations were based on duplicate measurements.

2.7. Thermal Stability and Denaturation Degree Determinations by Differential Scanning Calorimetry

Temperature, denaturation enthalpy and denaturation degree determinations were made on Polymer Laboratories PL–DSC equipment (Rheometric Scientific DSC 2, Ltd., Epsom, England). Aqueous dispersions of 20% w/v of the above described SPIHs in distilled water were used as samples. 20 mg of each such sample were placed in hermetic aluminium pans for analysis. Runs were conducted at a heating rate of 10 ºC/min over the temperature range of 25 ºC to 120 ºC. A reference run was conducted using one sample previously subject to heating treatment as used during the analysis. For all samples, the dry-matter content was determined upon drying to constant weight -using perforated pans- in an oven at 105 ºC.

The denaturation degree was calculated according to the equation: DDeg = $[(\Delta H - \Delta H_T)/ \Delta H] \times 100$, DDeg being the denaturation degree, ΔH the denaturation enthalpy of the unhydrolyzed starting protein, and ΔH_T the denaturation enthalpy of the protein modified by hydrolysis [13].

Runs were analyzed by means of Plus V5.41 Software. Determinations were based on duplicate measurements.

2.8. Interfacial Tension Determinations

Surface tension (σ) measurements at the air/water interface were carried out on a Lauda TVT 2 drop volume tensiometer equipped with 2.5 ml injection syringe. Determinations by the steady drop method were conducted for an initial number of six measuring cycles with three drops per cycle (6/3), using a drop formation rate in the range of 0.07-0.10 s/μl, followed by 9/3 cycles/drops per cycle using a drop

formation rate in the range of 0.10-0.80 s/μl. The pendant drop method, was used for 6/2 cycles/drops per cycle in the 12 to 16.21 s/m range of drop formation rate. Dissolutions at pH 8 with a protein concentration of 1mg/ml were obtained by dissolution of the protein hydrolysates in 0.1 M Na_3PO_3. The parameters used for analysis of here-described the assays, σ_e, k_a and k_r, are those used in the kinetic model developed by [14], where σ_e is the equilibrium surface tension, k_a and k_r first order rate constants of the protein adsorption and molecular rearrangement processes, respectively, at the air/water interface. Determinations were based on duplicate measurements of duplicate replicates.

2.9. Foaming Capacity Analysis

Foam formation and stabilization assays were conducted according to the conductimetric method developed by [15], following minor modification. Foam was generated by means of air bubbling at a flow rate of 100 ml/min through a G2-type sintered glass plate until collection of a preset foam volume of 60 ml. The analysis was conducted using 10 ml of dispersions of the hydrolysates in 0.1 M Na_3PO_3, at pH 8, using protein concentrations of 1 mg/ml, 2 mg/ml and 5 mg/ml. Following the procedure described by [16], determinations of the initial liquid-to-foam transfer rate (v_i) were based on initial slope analysis of the liquid volume incorporated in the foam (V_{LF}) vs. time plot. The maximum value of liquid volume in the foam (V_{max}) was also determined. The destabilization kinetics was analysed based on the parameters k_g, V_g, k_d and V_d resulting from application of the kinetic model developed by [17], where k_g, V_g, k_d and V_d are the rate constants and the maximum liquid volumes in the foam associated with gravitational drainage and gas diffusion or disproportionation processes, respectively, the latter also known as Ostwald's ripening. Determinations were based on triplicate measurements of duplicate replicates.

2.10. Statistical Analysis

The fixed-effects model of analysis of variance (ANOVA) was used for statistical treatment of data, with $\alpha = 0.05$; followed by the comparison of mean values by the least significant differences (LSD) test, with $\alpha = 0.05$, using Statgraphics plus 7.0 software.

3. Results and Discussion

3.1. SPIH Composition

HTs had a protein content in the range of 81.8 to 86.6 g/100 g, a water content between 12.10 and 13.20 g/100 g, and an ash content between 3.07 and 5.98 g/100 g while HpHs had a slightly lower protein content, in the range of 79.1 to 80.4 g/100 g, a water content between 11.87 and 13.56 g/100 g and an ash content between 6.03 and 9.08 g/100 g. Except for the ash content of HpHs, the above contents are in agreement with the expected values considering the starting soybean flour (protein: 51.5 ± 0.3 g/100 g; water: 6.89 ± 0.03 g/100 g; ash: 6.20 ± 0.03 g/100 g)

and the soy protein isolate (protein: 78.6 ± 0.3 g /100 g; water: 10.09 ± 0.05 g /100 g; ash: 4.53 ± 0.02 g /100 g). The ash content of HpHs was higher than that of the starting material, on account of HCl additions used for halting the reaction.

3.2. Structural Characterization of SPIHs

3.2.1. Polypeptide Composition

Polypeptides characteristic of β-conglycinin and glycinin were found by SDS-PAGE electrophoresis analysis of SPI control samples (HDeg 0%) and SPIH samples of low hydrolysis degree (HDeg 1.8%). For both HTs and HpHs, an increased HDeg (2.5% and 6.0%) led to the disappearance of the α and α' bands of β-conglycinin and to a slight reduction of polypeptide A of glycinin (data not shown). In [18] it was reported that the action of papain on soybean protein isolates was initiated on β-conglycinin, followed by glycinin; the subunits of the former being hydrolyzed to a larger extent than the AB subunits of glycinin. Such results are in accordance with the above reported disappearance of the α and α' bands in SPIHs of high HDeg (2.5% and 6.0%).

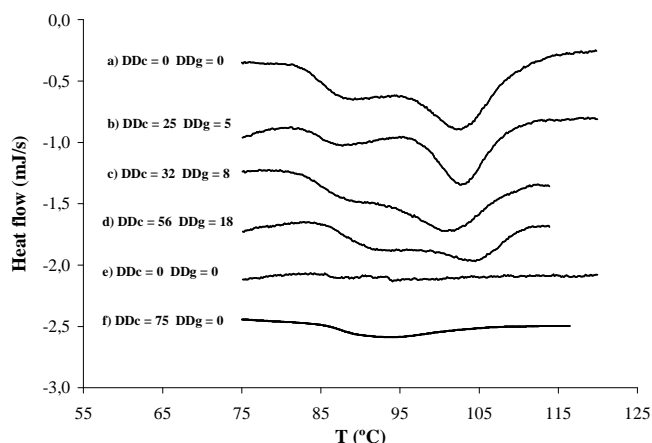#### 3.2.2. Analysis of Thermal Properties

Figure 1. *DSC thermograms of HT HDeg 0 (a), 1.8 (b), 2.5 (c) and 6.0% (d), HpH at pH 2 (e) and previously neutralized HpH (f). The first peak corresponds to β-conglycinin and the second one to glycinin. β-conglycinin (DDc) and glycinin (DDg) are expressed as percent fractions in all cases.*

Figure 1 shows the thermograms of the analyzed hydrolysates. HTs (pH 8.0) showed two endotherms associated with the denaturation of β-conglycinin and glycinin, respectively [19]. HTs of HDeg 6% showed an increase in maximum peak temperature (from 89 °C to 92 °C and from 102 °C to 104 °C) and a lesser extent of differentiation between both endotherms. Figure 1 shows, for HTs, an increase in DDeg with increasing HDeg, the effect on β-conglycinin being greater than on glycinin, consistent with the above reported results of electrophoresis assays. This suggests that a larger proportion of glycinin molecules may either retain or not show major disarrangement in their protein structure as a result of proteolysis.

HpH thermograms, shown in Figure 1, did not show the

endotherms corresponding to the denaturation of glycinin and β-conglycinin as a result of acid treatment used for halting the hydrolysis reaction. Similar results on the partial hydrolysis of β-conglycinin were previously reported [16], [20]. A partial reversal of the effect of acid treatment resulted from neutralization prior to thermal treatment of HpHs, leading to a DDeg of c.a. 75% consistent for all analyzed samples (see Figure 1).

3.2.3. Surface Hydrophobicity

HpHs were found to have hydrophobicity (Ho) nearly twice as high as that of HTs (HpH HDeg 0% Ho = 264 ± 2 and HT HDeg 0% Ho = 106 ± 2). In the range of low HDeg values, an increase in HDeg (to 1.8% for HpHs and to 1.8% and 2.5% for HTs) led to a significant increase in Ho (to 268 ± 2 and to 126 ± 2 and 112 ± 2, respectively) whereas an even higher HDeg (6.0%) led to a reduction in Ho to values nearly as low as those of the unhydrolyzed isolates (207 ± 2 and 85 ± 2, respectively). Protein dissociation and unfolding resulting from treatment at pH 2 led to increased exposure of sterically hindered hydrophobic zones of the unhydrolyzed proteins, with the resulting increase in surface hydrophobicity. Likewise, the unfolding of the protein structure resulting from hydrolysis further led to an increase in Ho. The increase in Ho as the reaction progressed may be attributed to the maintenance of major hydrophobic zones, such as polypeptide B of glycinin. In addition, the loss of hydrophilic peptides at the protein surface may also have led to an increase in hydrophobicity [21].

3.3. SPIHs Used for Surface Tension Modification

The parameters resulting from the model proposed by [14] were used for analysis of surface tension modification.

Table 1 shows the values of equilibrium surface tension (σ_e) and the protein adsorption and rearrangement first order rate constants at the air/water interface (k_a and k_r, respectively) for the assayed solutions.

Table 1. Surface tension variations in SPIHs [protein] = 1.0 mg/ml, pH = 8.0 and [NaCl] = 0 M

SPIH	$k_a \times 10$ (s^{-1})	$k_r \times 10^2$ (s^{-1})	σ_e (mN/m)
HT HDeg 0%	1.4 ± 0.1[a]	0.77 ± 0.09[b]	52 ± 1[a]
HT HDeg 1.8%	1.5 ± 0.1[a]	0.85 ± 0.08[b]	52 ± 1[a]
HT HDeg 2.5%	1.4 ± 0.1[a]	0.80 ± 0.08[b]	53 ± 1[a]
HT HDeg 6.0%	1.4 ± 0.1[a]	0.61 ± 0.09[c]	56 ± 1[b]
HpH HDeg 0%	1.6 ± 0.3[a]	1.3 ± 0.2[a]	50 ± 2[a]
HpH HDeg 1.8%	1.6 ± 0.3[a]	1.8 ± 0.2[a]	53 ± 2[a]
HpH HDeg 2.5%	2.0 ± 0.3[a]	1.7 ± 0.2[a]	51 ± 2[a]
HpH HDeg 6.0%	1.4 ± 0.3[a]	1.2 ± 0.2[a]	51 ± 2[a]

Mean values of triplicate measurements are shown
Mean values identified by the same letter do not differ significantly (α = 0.05)

No significant differences were found in k_a values for the treatments used to halt the reaction halting or for HDeg. k_r differed in significant amounts for either type of ending treatment; the values for HpH being higher than for HT. The

HDeg affected k_r only in the case of HTs, HDegs of 0%, 1.8% and 2.5% leading to higher values than for HDeg 6%. Likewise, no significant differences were found for σ_e according to ending treatment or HDeg, except for HT of HDeg 6% with a significantly higher σ_e value than for the other HTs (0%, 1.8% and 2.5%).

Above results showed that differences in the kinetics of surface tension modification found for HT and HpH are not to be ascribed to different equilibrium surface tension values for either type of hydrolysate, but, rather, to the rate of variation thereof (Figure 2). Such difference is reflected in the k_r value, meaning that differences may lie only on a different protein rearrangement capacity at the interface. Higher k_r values found for HpH may be attributed to a higher Ho and a lower molecular size of constituent polypeptides in view of the predominance of the 3S form of glycinin at acid pH [22], leading to a greater ease of rearrangement at the interface. The above is consistent with results reported by [23] on the ease and high rate of unfolding and rearrangement of 3S glycinin at the interface on account of a higher flexibility resulting from electrostatic repulsion forces within the molecule.
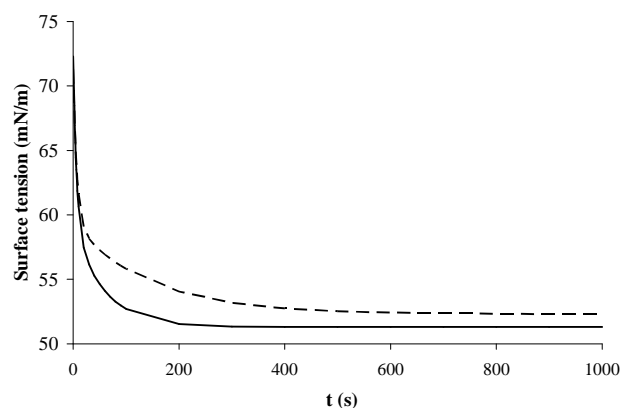

Figure 2. Surface tension vs. time plots showing the behaviour of HT (---) and HpH (-)

In [11] it was reported that β-conglycinin is the protein fraction with the greatest interfacial surface reduction capacity. As shown in Table 1 HT of HDeg 6% showed the lowest k_r value, a fact which may be attributed to the reduction in size and the loss of globular structure of β-conglycinin as a result of hydrolysis. Although such loss of structure was found consistently for all HpHs, a different behaviour from that of HTs was found. The fact that glycinin in HpHs –i.e. receiving pH 2 treatment- is dissociated, less affected by proteolysis and has greater flexibility may account for the invariance in the k_r value found for HpH HDeg 6% where the effect of the treatment used for hydrolysis halting was greater than that of the HDeg attained by reaction.

3.4. Foamability

The effects of the different variables used in the study on the initial liquid-to-foam transfer rate (v_i) and the maximum

liquid volume incorporated in the foam phase (V_{max}) are   shown in Table 2.

Table 2. v_i and V_{max} values in foams formed using dissolutions of different SPIHs at different protein concentrations, at pH 8.0

SPIH	[protein]: 1.0 mg/ml		[protein]: 2.0 mg/ml		[protein]: 5.0 mg/ml	
	v_i (ml/s)	V_{max} (ml)	v_i (ml/s)	V_{max} (ml)	v_i (ml/s)	V_{max} (ml)
HT HDeg 0%	0.18 ± 0.02^b	4.8 ± 0.3^b	0.23 ± 0.02^c	6.5 ± 0.3^d	0.23 ± 0.0^c	7.4 ± 0.3^a
HT HDeg 1.8%	0.16 ± 0.02^b	4.7 ± 0.3^b	0.22 ± 0.02^c	6.6 ± 0.3^d	0.25 ± 0.02^c	7.4 ± 0.3^a
HT HDeg 2.5%	0.17 ± 0.02^b	4.7 ± 0.3^b	0.24 ± 0.03^c	6.7 ± 0.3^d	$0.28 \pm 0.04^{a.c}$	7.5 ± 0.3^a
HT HDeg 6.0%	0.16 ± 0.02^b	3.6 ± 0.4^c	0.17 ± 0.04^b	4.5 ± 0.4^b	0.23 ± 0.02^c	$6.9 \pm 0.4^{a.d}$
HpH HDeg 0%	0.26 ± 0.02^a	7.6 ± 0.3^a	0.27 ± 0.01^a	7.4 ± 0.3^a	0.30 ± 0.03^a	7.7 ± 0.3^a
HpH HDeg 1.8%	0.30 ± 0.02^a	7.8 ± 0.3^a	0.28 ± 0.02^a	7.6 ± 0.3^a	0.29 ± 0.03^a	8.1 ± 0.3^a
HpH HDeg 2.5%	0.30 ± 0.02^a	7.8 ± 0.3^a	0.32 ± 0.02^a	8.0 ± 0.2^a	0.36 ± 0.03^a	8.2 ± 0.3^a
HpH HDeg 6.0%	0.26 ± 0.02^a	7.1 ± 0.3^a	0.28 ± 0.03^a	7.7 ± 0.3^a	0.30 ± 0.03^a	7.9 ± 0.3^a

Mean values of triplicate measurements are shown
Mean values identified by the same letter d

o not differ significantly ($\alpha = 0.05$)

v_i and V_{max} values found for HpH foams were significantly higher than for HT foams. The HDeg affected V_{max} significantly only for HT of HDeg 6%, leading to a significantly lower value than for the rest of the analyzed samples (0%, 1.8% and 2.5%) (Table 2). For HT foams, both v_i and V_{max} increased with increasing protein concentration in the aqueous dissolution used for foaming assays. The effect of protein concentration on V_{max}, led to a lesser extent of variation of the latter between HT and HpH foams.

The liquid volume drained by a foam is affected by the bubble size according to Laplace's Law and by a reduction in surface tension, in turn leading to a lower pressure difference between bubbles. The ΔP ratio for foam preparations using different protein solutions, assuming equal values of curvature radii, may be expressed as follows:

$$\Delta P_1 / \Delta P_2 = \sigma_1 / \sigma_2 \qquad (1)$$

The maximum difference between σ values found for HT and HpH preparations, during the kinetics of surface tension reduction, was that between 58 mN/m and 55 mN/m respectively -i.e. the maximum ΔP between HT and HpH bubbles being $58/55 = 1.05$. ΔP differences, and hence differences in v_i and V_{max} values between HTs and HpHs, may clearly not be accounted for in terms of σ values. A higher rate of incorporation of liquid into the foam phase may be associated with a greater resistance of HpH foams against incipient destabilization processes such as film rupture, Ostwald's ripening and gravitational liquid drainage. The above may also account for the higher V_{max} values found for HpH foams, as previously mentioned.

According to [23], whereas both 3S and 11S glycinin may form cross-linkage at the air/water interface, the latter is less rigid on account of a less compact structure. For proteins previously denatured by the effect of acid pH, glycinin -occurring predominantly in the 3S form- appears to have a greater ease and rate of rearrangement at the interface, in addition to the capacity of forming physical and covalent intermolecular interactions within the interfacial film. Hence,

the film formed by HpH proteins is expected to show a higher resistance to destabilizing processes.

In the case of HT of HDeg 6%, having a higher concentration of small-molecular-sized peptides adversely affecting foam stability [24], the above behaviour did not favour the formation of a cohesive film resistant to rupture and Ostwald's ripening. Acid treatment -leading to dissociation, denaturation and unfolding of HpH proteins- offset the effect of hydrolysis, resulting in a similar interfacial behaviour of HpH of HDeg 6% to that found for the rest of HpHs.

The increase in protein concentration may lead to a higher lamellar viscosity and the formation of a cohesive film composed of several layers of protein molecules at the interface, according to [25]. Such effects would result in a greater film resistance against rupture and Ostwald's ripening. Higher v_i and V_{max} values may thus have resulted from an increase in protein concentration. The effect of protein concentration was less significant for HpH foams (Table 2). The above may be explained in terms of an improved interfacial behaviour associated with the stabilization of the film layer; while the improvement derived from an increase in protein concentration appears to have reached an optimal level at a value of 1mg/ml.

3.5. Foam Stability

Assays of foam stability were based on the rate constants of liquid drainage in the foam. The effects of the different variables studied on the values of drainage and gravitational rate constants, k_d and k_g respectively, are shown in Table 3.

3.5.1. Gravitational Drainage

k_g values of HpH foams with a protein concentration of 1 mg/ml were significantly lower than for HT foams. With increasing protein concentration, differences between foams of both types of hydrolysate were reduced to insignificant values above at protein concentrations above 5 mg/ml. HT foams of HDeg 6% showed a significantly higher k_g value than for the rest of HTs (0%, 1.8% and 2.5%) (Table 3); whereas the HDeg did not have a significant effect on k_g in

HpH foams (Table 3). In HT foams, k_g decreased significantly with increasing protein concentration (Table 3).

Much research has been reported on an inverse relation between foamability and stabilizing properties of proteins [26], [27], [28], [29], [25], [30], [31]. The results reported here are in disagreement with a generalization of the above statement as such, since both a greater foamability as well as stabilizing properties were found for HpH foams, consistently with results also reported by [32] and [16].

As foam stabilizing mechanisms occur during the stage of foam formation [33], the proteins that account for the good foamability of a solution may also render it more stable. In [1] there is reported on the rapid formation of a thin rigid protein film –i.e. which may be considered as a gel layer- following the adsorption of 3S glycinin under appropriate occurrence conditions at the interface. HpH proteins therefore appear to form a gel-like protein layer leading to foam of a higher stability of a foam. Based on findings reported by [23], while both 3S and 11S glycinin have cross-linking capacity at the interface, the latter appears as less rigid in view of a greater degree of compaction, which may account for the fact that HT foams were found to retain a smaller amount of water in the lamella, consistent with a higher k_g value found for foams of this type of hydrolysate.

The gel forming capacity was found to decrease with decreasing molecular size. For HDeg of 6%, where the number of short-chain peptides was found to be greater than for the rest of the studied HDegs, the capacity of forming a gel-like film at the interface appears to have been reduced. The occurrence of low-molecular peptides may also have contributed to a lesser degree of resistance against the drainage process in foams of highly hydrolyzed protein.

Nonetheless, the effect of hydrolysis was offset by that of a high DDeg consistently for all HpHs -where glycinin was found under AB subunits- with the result that no significant variation of stabilizing properties against liquid drainage was found among foams of this type of hydrolysate.

As discussed above in reference to the foam formation capacity, an increase in protein concentration led to a higher viscosity of the lamella and may contribute to the formation a thicker and more cohesive film at the interface comprised of various layers of protein molecules [26], [34], [27], [29], [26]. Further, a higher protein concentration may result in a higher extent of interaction with water molecules, thus retarding the drainage process. Protein concentration did not have a significant effect on stability for HpH foams above an optimal protein concentration value found at 1mg/ml, as discussed above.

The amount of drained liquid is strongly influenced by the size of bubbles, as discussed above, the former decreasing with increasing bubble radius. Despite the smaller bubble radius of HpH foams, they were found to have a smaller rate of liquid drainage, leading to an enhancement of the stabilizing properties of these foams compared with those of HT foams.

3.5.2. Liquid Drainage Due to Ostwald's Ripening

HT foams were found to have a higher stability to Ostwald's ripening, consistent with the estimated k_d values amounting to several orders of magnitude below the values found for HpHs (Table 3). k_d values for HpH foams were found to decrease significantly with increasing protein concentration (Table 3).

Table 3. k_g and k_d values in foams prepared using SPIH solutions at pH 8.0 for different protein and NaCl concentrations.

SPIH	[protein]: 1.0 mg/ml		[protein]: 2.0 mg/ml		[protein]: 5.0 mg/ml	
	$k_g \times 10^3$ (ml^{-1}.s^{-1})	$k_d \times 10^4$ (ml^{-1}.s^{-1})	$k_g \times 10^3$ (ml^{-1}.s^{-1})	$k_d \times 10^4$ (ml^{-1}.s^{-1})	$k_g \times 10^3$ (ml^{-1}.s^{-1})	$k_d \times 10^4$ (ml^{-1}.s^{-1})
HT HDeg 0%	10 ± 1 [b]	0.005 ± 0.001 [b]	6.2 ± 0.9 [c]	0.005 ± 0.001 [b]	3.0 ± 0.2 [a]	0.005 ± 0.001 [b]
HT HDeg 1.8%	12 ± 3 [b]	0.005 ± 0.001 [b]	5.1 ± 0.5 [c]	0.005 ± 0.001 [b]	3.2 ± 0.3 [a]	0.005 ± 0.001 [b]
HT HDeg 2.5%	14 ± 2 [b]	0.005 ± 0.001 [b]	5.5 ± 0.9 [c]	0.005 ± 0.001 [b]	3.3 ± 0.9 [a]	0.005 ± 0.001 [b]
HT HDeg 6.0%	37 ± 7 [d]	0.005 ± 0.001 [b]	27 ± 5 [d]	0.005 ± 0.001 [b]	8.0 ± 0.9 [c]	0.005 ± 0.001 [b]
HpH HDeg 0%	3.4 ± 0.6 [a]	9 ± 2 [a]	3.8 ± 0.8 [a]	5 ± 1 [a]	2.7 ± 0.3 [a]	5 ± 1 [a]
HpH HDeg 1.8%	3.2 ± 0.3 [a]	9 ± 2 [a]	2.9 ± 0.3 [a]	4 ± 1 [a]	2.7 ± 0.4 [a]	5 ± 1 [a]
HpH HDeg 2.5%	2.5 ± 0.1 [a]	7 ± 2 [a]	3.0 ± 0.6 [a]	5 ± 1 [a]	2.8 ± 0.4 [a]	5 ± 1 [a]
HpH HDeg 6.0%	3.5 ± 0.4 [a]	7 ± 2 [a]	2.8 ± 0.2 [a]	5 ± 1 [a]	2.9 ± 0.6 [a]	5 ± 1 [a]

Mean values of triplicate measurements are shown
Mean values identified by the same letter do not differ significantly ($\alpha = 0.05$)

Disproportionation may be inhibited or retarded by a sufficiently thick layer of interfacial film as to resist the passage of gas [35], HpH foams thus being expected to show a greater resistance against disproportionation. The fact that HT foams showed a greater resistance to disproportionation, however, results from a strong dependence of such resistance on the bubble size of a particular foam, the latter being greater for HTs. However, as the increase in protein concentration led to the formation of increasingly resistant film, the above reported difference between hydrolysate types was lessened.

3.5.3. Volumetric Proportions of Gravitational Liquid Drainage and Ostwald's Ripening

In addition to the study of the above kinetic constants, the contributions of gravitational liquid drainage (V_g) and disproportionation (V_d) to the total drained liquid volume were analyzed according to variations in the study parameters (Table 4).

The proportion of liquid drained by gravitational drainage was significantly higher –at ratios no smaller than 0.80- than that due to disproportionation for all the assayed foams. The

V_g value was found not to depend significantly on protein concentration for foams prepared with HpHs (Table 4). In contrast, in foams prepared with HTs, V_g was found to increase with increasing protein concentration (Table 4), unlike k_g decreasing with increasing protein concentration (Table 3).

Foams prepared with HTs at a protein concentration of 5 mg/ml showed V_g values in the range of 0.98 and 1.00, the contribution of disproportionation to the total volume of drained liquid being negligible or nil (Table 4).

Table 4. V_g and V_d in foams prepared using SPIHs at pH 8.0 for different protein concentrations

SPIH	[protein]: 1.0 mg/ml		[protein]: 2.0 mg/ml		[protein]: 5.0 mg/ml	
	V_g (ml)	V_d (ml)	V_g (ml)	V_d (ml)	V_g (ml)	V_d (ml)
HT HDeg 0%	0.90 ± 0.02[a]	0.10 ± 0.01[b]	0.99 ± 0.02[c]	0.01 ± 0.01[d]	1.00 ± 0.00[c]	0.00
HT HDeg 1.8%	0.89 ± 0.02[a]	0.11 ± 0.01[b]	0.97 ± 0.02[c]	0.03 ± 0.01[d]	1.00 ± 0.00[c]	0.00
HT HDeg 2.5%	0.92 ± 0.02[a]	0.08 ± 0.01[b]	0.94 ± 0.03[a,c]	0.06 ± 0.01[b,d]	0.98 ± 0.01[c]	0.02 ± 0.01[d]
HT HDeg 6.0%	0.90 ± 0.02[a]	0.10 ± 0.01[b]	0.95 ± 0.02[c]	0.05 ± 0.01[d]	0.99 ± 0.01[c]	0.01 ± 0.01[d]
HpH HDeg 0%	0.90 ± 0.02[a]	0.10 ± 0.01[b]	0.87 ± 0.02[a]	0.13 ± 0.01[b]	0.84 ± 0.02[a]	0.16 ± 0.01[b]
HpH HDeg 1.8%	0.87 ± 0.02[a]	0.13 ± 0.01[b]	0.87 ± 0.02[a]	0.13 ± 0.01[b]	0.85 ± 0.02[a]	0.15 ± 0.01[b]
HpH HDeg 2.5%	0.85 ± 0.02[a]	0.15 ± 0.01[b]	0.86 ± 0.02[a]	0.14 ± 0.01[b]	0.86 ± 0.02[a]	0.14 ± 0.01[b]
HpH HDeg 6.0%	0.89 ± 0.02[a]	0.11 ± 0.01[b]	0.88 ± 0.02[a]	0.12 ± 0.01b[b]	0.84 ± 0.02[a]	0.16 ± 0.01[b]

Mean values of triplicate measurements are shown
Mean values identified by the same letter do not differ in significant amounts ($\alpha = 0.05$)

The relative amounts of liquid drained by gravitational drainage and disproportionation respectively do not only depend on the bubble size but also on the resistance of the protein film at the interface. In conditions as used for the above-described assays, liquid drainage was predominantly driven by gravitation rather than disproportionation.

Worth noting is the mutual dependence between V_g and V_d values, unlike the case of k_g and k_d, as the former are expressed as proportions.

Unlike k_g, k_d values were found to depend on the protein concentration in aqueous dissolution as used for the above-described assays, resulting in a reduction of k_d with increasing concentration, and, in turn, a lesser extent of drainage due to disproportionation, and thus a higher proportion of the total drained liquid volume due to gravitational drainage (Table 3). The rate of gravitational drainage may be described by means of the Reynolds equation [36]:

$$V = -dh/dt = (2h^3/3\mu R^2) \Delta P \qquad (2)$$

where h is the thickness of the lamella, t time, μ the dynamic viscosity, R the bubble radius and ΔP the pressure gradient according to Laplace's law. The rate of gravitational drainage clearly decreases with decreasing thickness of the lamella. A similar prediction may be made for drainage through Plateau borders [34]. The volume of liquid drained by gravitation will approach a constant value with time; a value which was found not to vary significantly with increasing protein concentration, consistent with insignificant differences found in k_g or V_{max} values for different concentration levels. Therefore, the decrease in k_d did not lead to a reduction of the volume of drained liquid due to disproportionation, yet, it did result in an increase of the time required to attain complete drainage of liquid incorporated in the foam.

Whereas the difference in k_g values between HT and HpH foams lessened with increasing protein concentration, V_g values for HT foams were significantly higher than for HpH

foams at protein concentrations of 2 mg/ml and 5 mg/ml. In HT foams, while an increase in protein concentration led to retardation in the rates of both gravitational drainage and disproportionation processes, the latter was found to occur to a less significant extent.

4. Conclusions

According to the results reported here, a low hydrolysis degree (HDeg 1.8 and 2.5%) did not result in the enhancement of the foaming properties of hydrolysates obtained by freezing. For hydrolysates of this type, only a high hydrolysis degree (HT of HDeg 6%) affected the foaming properties adversely. The loss of foaming capacity of HT of HDeg 6% was due to a greater bubble size and a lower V_{max} value, and the lower stability found for foams of this hydrolysate was reflected in a higher k_g value.

The adverse effect of an increasing hydrolysis degree on the behaviour of foams prepared with HTs was attributed to a reduction in polypeptide molecular size, leading to the formation of a film layer lacking adequate viscoelasticity and cohesion. Such adverse behaviour may be ascribed to the effect of hydrolysis on β-conglycinin -rather than on glycinin. Whereas improved foaming properties may be attributed to glycinin, an excess number of small-sized polypeptides - particularly those resulting from the hydrolysis of β-conglycinin- were found to compete with the larger-sized proteins for adsorption at the interface, thus leading to the formation of less cohesive, viscoelastic and resistant film.

HpHs showed improved foaming properties with respect to the untreated isolates, consistently with smaller-sized bubbles and greater v_i, V_{max} and k_g values found for foams prepared with the former. The positive effect of pH treatment on interfacial properties was attributed to the dissociation and denaturation of soybean protein leading to favourable viscoelastic and cohesive properties of the film layer at the interface.

No adverse effect of a high hydrolysis degree (HDeg 6%)

was found on the foaming properties of hydrolysates subject to pH treatment. Whereas low-pH treatment leads to dissociation of β-conglycinin, a great extent of hydrolysis adversely affected the foaming properties of HpHs of HDeg 6%. Therefore, the improvement of film properties leading to enhanced functional properties found for HpHs may be largely attributed to the dissociation of glycinin into subunits AB and to their denaturation.

Acknowledgements

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