Optimum conditions for enzymatic degradation of some oilseed proteins

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INTRODUCTION

The enzymatic hydrolysis of proteins from different sources to improve their solubility, digestibility and other functional properties has been well established (1-3). The improvement of the functional properties enables the protein to be successfully incorporated into many food systems. Proteolytic enzymes from animal plant, and microbial origin (1-5) have been utilized on different substrates, such as soybean (6-9), cottonseed (1), peanut (3), sesame (10,11), and sunflower (8).

Protein hydrolysates used in nutritional formulations are generally categorized into two broad categories, partially hydrolyzed and extensively hydrolyzed proteins. Each category possesses different properties that influence their utilization in the final product. Presently there are no precise, widely accepted specifications to differentiate the protein hydrolysates on chemical basis. Extensively hydrolyzed proteins have substantially reduced immunological reactivities and are primarily used in hypoallergenic formulas, because of the need to almost completely eliminate their allergenicity to avoid sensitization of individuals consuming them (12). These protein hydrolysates are usually comprised of amino acids and very short peptides (di and tri peptides).

The aim of the present investigation was to elucidate the optimum conditions for the enzymatic hydrolysis of soybean, sesame seed and rice bran meal protein, with two enzymes one of plant origin papain, and bromelain. The numerical correlation of enzymatic behaviour for different substrates was also elucidated. The results of this investigation will be the key to further work on partial degradation of these proteins.

MATERIALS AND METHODS

2. Materials

2.1. Oilseed meals

Soybean (glycine max) and sesame (Sesamum indicum) were supplied by the Research Institute of Field Crops, Ministry of Agriculture, Cairo, Egypt.
Rice bran (Oryza sativa) was obtained from a local rice mill.

Soybeans and sesame seeds were dehulled and ground then subjected to several extractions with n-hexane to extract the oil. After two extractions, the meats are reground and extraction continued until residual oil in the meals did not exceed 1%. Rice bran meal was prepared in the same manner but excluding the dehulling step. The defatted meals were spread to dry at room temperature and then ground and sieved to pass an 80 mesh screen.

2.1.2. Enzymes

- Papain: DIFCO LABORATORIES (MICHIGAN USA), activity of papain while carrying experiments was 8.9 units/mg, unit as µg tyrosine released from casein/min/mg.
- Bromelain: A product of SIGMA (Missouri USA), activity of bromelain while carrying the experiments was 27.6 unit /mg, unit as µg tyrosine released from casein/ min/ mg. All reagents were of analytical grade.

2.2. Methods

2.2.1. Determination of relative activity

To a 250 ml beaker that was placed in a thermostatic water bath, were added 100 ml water and a weight of meal to containing 5g protein. The mixture was continuously stirred with an electric stirrer and the pH and temperature adjusted to desired values. The reaction was carried on for 30 minutes while stirring and the pH and temperature maintained at the at previously adjusted values. The reaction was stopped by the addition of 0.5M TCA, and the hydrolysate filtered through Whatman no. 4, the residue washed with distilled water until an approximate volume of 250ml was collected. Aliquots of the filtrate were analyzed for total protein using the kjeldahl method and the relative activity was calculated as follows:

Relative activity (RA) = \( \frac{C_p - H_p}{C_p} \times 100 \)

Where: \( H_p \) is total protein obtained in enzyme assay
\( C_p \) is total protein in original meal (control).

2.2.2. Determination of the optimum conditions for the proteolytic activity of the enzymes

This was accomplished through a series of experiments.

In the first set of experiments the enzyme papain was used together with sodium sulphite (which acts as an activator for papain) on the three oilseed meals. The relative activity was determined by kjeldahl as mentioned above.

The first investigated criteria was the E/S ratio which is the ratio of concentration of enzyme to concentration of substrate. The E/S investigated ranged between 0.01 to 1.83. The temperature and pH were those recommended by the manufacturer, and the experiment was carried as for determination of relative activity.

The pH was the second investigated criteria, the E/S ratio was the best resulting from the former experiment, the temperature was that of the manufacture. The investigated pH were 6.8, 7.0, 7.2, 7.4, 7.6, the experiment proceeded as described for the relative activity determination.

In the third experiment the temperature was the variable ranging from 40 to 80°C, best pH and best E/S as previously elucidated, and experiment proceeded as for determination of relative activity.

In the second set of experiments the enzyme bromelain was used on the three oilseed substrates. The sequence of experiments followed the same route as those carried with papain. The E/S investigated were 0.021 to 0.078, pH 5.8, 6.0, 6.2, 6.4, 6.6, and 6.8, the temperature 35 to 50°C.

2.3. Analysis

Moisture, oil, ash, fiber and nitrogen were determined according to AOCS (13) standard methods. Protein calculated as N x 6.25 for sesame and rice bran, and N x 5.7 for soybean.

3. RESULTS AND DISCUSSION

In order to carry the enzymatic degradation of the proteins in the most effective manner, optimum conditions of E/S ratio, pH and temperature of the reaction were examined. Two enzymes papain and bromelain were investigated and three substrates including soybean, sesame seed and rice bran meal proteins.

Table I gives the chemical composition of the three investigated oilseed meals. All values in table are given on moisture free basis.

3.1. Optimum conditions for the two enzymes studied

The optimum conditions for the enzymes are expressed in terms of highest relative activity of the enzyme at different values of E/S ratios, pH values, and temperatures. The time of reaction was fixed at 30 minutes to be investigated in a continuation of this study. The optimum conditions for each enzyme was determined on each substrate separately.
3.1.1. Papain

Figures 1-3 represent the relation between the relative activity of the enzyme (papain) and the investigated parameters including E/S ratio, pH, temperature.

- Soybean meal: optimum E/S was 0.06, optimum pH 7.2 and optimum temperature 50°C all resulting in highest relative activity value of 107.8.
- Sesame meal: optimum E/S ratio 0.29, pH 7.0, temperature 50°C, giving rise to relative activities of 175.8, 81.8, and 54.4, respectively.
- Rice bran meal: optimum conditions were E/S ratio 0.19, pH 7.0, and temperature 50°C, for highest relative activities 165.8, 155.5 and 164.2, respectively.

3.1.2. Bromelain

Figures 4-6 demonstrates the relation between the relative activity of the enzyme (bromelain) and the investigated parameters (E/S ratio, pH, temperature).

- Soybean meal: E/S resulting in highest relative activity of 77.8 was 0.067,
- Sesame meal: Best E/S ratio, pH and temperature were 0.058, 6.0, and 45°C, respectively.
- Rice bran meal: pH 6.0 and temperature 45°C gave 84.0 and 77.8 highest relative activities, respectively.

Table I

<table>
<thead>
<tr>
<th>Analysis</th>
<th>Soybean</th>
<th>Sesame</th>
<th>Rice bran</th>
</tr>
</thead>
<tbody>
<tr>
<td>Oil%</td>
<td>0.9</td>
<td>0.7</td>
<td>0.5</td>
</tr>
<tr>
<td>Protein%</td>
<td>43.9</td>
<td>53.4</td>
<td>12.7</td>
</tr>
<tr>
<td>Ash%</td>
<td>6.6</td>
<td>6.4</td>
<td>5.9</td>
</tr>
<tr>
<td>Fibre%</td>
<td>2.1</td>
<td>1.9</td>
<td>13</td>
</tr>
<tr>
<td>NFE%</td>
<td>46.5</td>
<td>37.6</td>
<td>68</td>
</tr>
</tbody>
</table>

NFE: Nitrogen free extract
All values are given on moisture free basis.
respectively, resulting in highest relative activities 87.5 for all the parameters.

- Rice bran meal: Relative activities 162.5, 162.2, and 162.4 were attained with E/S 0.21, pH 6.0, and temperature 45°C, respectively.

Hermanson et al. (14), hydrolysed rapeseed protein concentrate with papain using the following conditions: pH 6.9, temperature 65°C and an enzyme concentration of 0.30%. Childs (15), reported optimum pH for papain to be 7.5, while Sekul and Ory (3), working with peanut flour protein and papain recommended a temperature of 45°C and enzyme concentration 0.5%. Arzu et al. (1) determining the relative activity of ten commercially available proteolytic enzymes reported relative activity of papain concentrate to be 129.1 at pH 7.2, and the relative activity of two bacterial proteinase to be 49.5 and 58.1 at pH 6.8.

3.2. Numerical correlation of enzymatic behavior for different substrates

3.2.1. Papain

Empirical formulae, which correlate the influencing parameters with the relative activity of papain are obtained by plotting E/S and temperature against the relative activity on ordinary scale and conducting of regressions, the following were the results.

The relative activity is directly proportional to the pH of the reaction media, and is proportional exponentially with approximately 2.8 E/S, in case of soybean and sesame while with approximately – 0.4 E/S in case of rice bran, and is also proportional exponentially with approximately – 0.03 temperature.

Therefore in case of soybean

$$RA = 16.3 \times e^{1.57 \times (pH - 0.03T \times e^{-2.84E/S})}$$  

(1)

Correlation (1) is applicable within the following range:

E/S = 0.01 - 0.06
pH = 7.2 – 7.6
Temperature: 50 - 80°C

In case of rice bran

$$RA = 50.78 \times e^{0.693 \times (pH - 0.03T \times e^{-0.41E/S})}$$  

(2)

And it is applicable within the following range:

E/S = 0.01 – 1.9
pH = 7.0 – 7.4
Temperature: 50 - 80°C

In case of sesame

$$RA = 2278.23 \times e^{-3.772 \times (pH - 0.03T \times e^{-2.82E/S})}$$  

(3)

Which is applicable within the range of:

E/S = 0.01-0.3
pH = 7.0 – 7.6
Temperature: 40 - 80°C

$$RA = 40.79 \times e^{-2.089 \times (pH - 0.03T \times e^{-2.82E/S})}$$  

(4)

Which is applicable within the range of:

E/S = 0.3 – 0.5
pH = 7.0 – 7.6
Temperature: 40 - 80°C

3.2.2. Bromelain

Empirical formulae, which correlate the influencing parameters with the relative activity of bromelain were obtained for each substrate at certain specific ranges of those parameters.

Plotting of E/S, pH and temperature against relative activity on ordinary scale, and by conducting linear regressions, one can find that the relative activity of bromelain for each substrate (soybean, sesame an rice bran) is directly proportional to E/S, pH, and temperature.

Therefore in case of soybean

$$RA = 31.776 \times e^{-0.05 \times (E/S \times pH \times T)}$$  

(5)

In the ranges of
E/S = 0.022 – 0.067  
\[ \text{pH} = 6.2 \text{ – 6.8} \]  
Temperature = 35 -45°C  
And  
\[ \text{RA} = 266.35 - 10.446 \times (E/S \times \text{pH} \times T) \]  
(6)

Correlation (6) is applicable within the following range:  
\[ E/S = 0.067 - 0.078 \]  
\[ \text{pH} = 6.2 \]  
Temperature: 45 -50°C

In case of rice bran  
\[ \text{RA} = 383 - 4.38 \times (E/S \times \text{pH} \times T) \]  
(7)

Correlation (7) is applicable within the following range:  
\[ E/S = 0.21 \]  
\[ \text{pH} = 5.8 \text{ – 6.8} \]  
Temperature = 35 - 45°C  
And  
\[ \text{RA} = 372.62 - 4.17 \times (E/S \times \text{pH} \times T) \]  
(8)

Correlation (8) is applicable within the following range:  
\[ E/S = 0.21 \text{ – 0.25} \]  
\[ \text{pH} = 6.0 \]  
Temperature = 45 -50°C

In case of sesame  
\[ \text{RA} = 6.22 - 5.845 \times (E/S \times \text{pH} \times T) \]  
(9)

Correlation (9) is applicable within the following range:  
\[ E/S = 0.02 \text{ – 0.058} \]  
\[ \text{pH} = 5.8 \text{ – 6.0} \]  
Temperature = 40°C  
\[ \text{RA} = 234.78 - 10.35 \times (E/S \times \text{pH} \times T) \]  
(10)

Correlation (10) is applicable within the following range:  
\[ E/S = 0.058 \]  
\[ \text{pH} = 6.0 \text{ – 6.8} \]  
Temperature = 40 -45°C

The relative activity of papain and bromelain calculated from the above empirical formulae were in fair agreement with those obtained from experimental work, and the mean standard errors for each enzyme were within the ± 10% as shown in (Figures 7 and 8).

**REFERENCES**


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