Experimental and Theoretical Studies on the Anti-Amyloidogenic and Destabilizing Effects of Pyrogallol Against Human Insulin Protein

Parastoo Shouhani  
Lorestan University Faculty of Basic Science

Seifollah Bahramikia (✉ bahramikia.s@lu.ac.ir)  
Lorestan University  https://orcid.org/0000-0002-9482-6828

Seyed Hesam Hejazi  
Lorestan University Faculty of Basic Science

Research Article

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Abstract

One of the major problems caused by repeated subcutaneous insulin injections in diabetic patients is insulin amyloidosis. Understanding the molecular mechanism of amyloid fibril formation of insulin and finding effective compounds to inhibit or eliminate is very important and extensive research has been done on it. In this study, the anti-amyloidogenic and destabilizing effects of the pyrogallol, as a phenolic compound, on human insulin protein were investigated by CR absorbance, ThT and ANS fluorescence, FTIR spectroscopy and atomic force microscopy. According to the obtained results, the formation of amyloid fibrils at pH 2.0 and 50°C was confirmed by CR, ThT, ANS, FTIR assays. Microscopic images also showed the twisted and long structures of amyloid fibrils. Simultaneous incubation of the protein with pyrogallol at different concentrations reduced the intensities of CR, ThT and ANS in a dose-dependent manner and no trace of fibrillar structures was observed in the microscopic images. FTIR spectroscopy also showed that the position of amide I band in the spectrum of samples containing pyrogallol was shifted. Based on the findings in this study, it can be concluded that pyrogallol can be effective in preventing and suppressing human insulin amyloid fibrils.

Introduction

Diseases related to protein misfolding are disorders caused by various factors such as genetic mutation, heat shock and malfunction of monitoring systems. As a result of these processes, the correct process of protein folding is disrupted and leads to its opening and exposure to hydrophobic surfaces. Following the change in protein structure, incorrect interactions occur between adjacent protein molecules, and eventually, very orderly and insoluble structures called amyloid fibrils are formed (Chiti and Dobson 2006; Invernizzi et al. 2012; Knowles et al. 2014; Rochet 2007). Many amyloidogenic proteins such as amyloid β, α-synuclein, islet amyloid polypeptide, and human lysozyme have been associated with protein misfolding diseases (Cheng et al. 2013). Extracellular aggregation and deposition of amyloid fibrils lead to cell death and in many cases cytotoxicity (Rochet 2007).

Insulin hormone is a small globular protein with 51 amino acid residues (Hua and Weiss 2004). This hormone plays an important role in regulating glucose metabolism and is used in the treatment of diabetes (Groenning et al. 2009). Insulin fibrillation in the body may occur after continuous and subcutaneous injection in diabetic patients and lead to local deposition of insulin amyloid fibrils at the injection site (Okamura et al. 2013; Shikama et al. 2010). On the other hand, fibrillation is also one of the problems that occur during the production and storage of insulin as well as the agitation of insulin solutions during transport (Groenning et al. 2009; Ahmad et al. 2003). In vitro factors such as temperature, pH, protein concentration, ionic strength, the presence of organic solvents and denaturants such as urea and guanidine chloride are effective in changing the structure and formation of insulin amyloid fibrils (Tartaglia et al. 2008; Selivanova and Galzitskaya 2012).

In recent years, finding a strategy for the treatment of amyloid diseases has been considered by many researchers. Important treatment strategies proven in most of these studies are to prevent the formation
of amyloid fibrils and their disaggregation. It has been shown that small organic molecules, including polyphenols, due to having aromatic rings, possess the ability to interact and bind to proteins and show significant anti-amyloidogenic effects. In addition, the antioxidant properties of polyphenols have made these compounds very effective in clearing free radicals and thus reducing cancer, as well as in the treatment of chronic and degenerative diseases (Sgarbossa 2012; Porat et al. 2006; Žerovnik 2002). Reduction of amyloid-induced cell death and inhibition of self-association of amyloidogenic proteins in vitro are other beneficial functions of polyphenols (Shaham-Niv et al. 2018).

In the structure of polyphenols, there is at least one aromatic ring with hydroxyl groups attached to it. Due to the number and location of these phenolic rings, as well as the presence of other constituent elements, various structures of polyphenols have emerged (Ozcan et al. 2014; Pandey and Rizvi 2009). Phenolic compounds are found in large quantities mainly in natural foods such as vegetables, fruits, seeds, nuts, as well as in flowers. Due to their natural origin and presence in the human diet, they are given more attention than other inhibitory molecules (Stefani and Rigacci 2013; Ozturk Sarikaya 2015).

1,2,3-Trihydroxybenzene, commonly known as pyrogallol (Py), is a phenolic compound like other natural polyphenols that is present in human food sources, including fruits and vegetables, and a variety of edible and medicinal plants (Ozturk Sarikaya 2015). Among the plants that have been mentioned as sources of pyrogallol are: Emblica officinalis (Amla or Indian Gosseberry), avocado, apricot, Terminaliachebula (Black Myrobalan and Harad), Entada abyssinica tree, tea, coffee, etc. So far, many beneficial activities for pyrogallol have been reported in various studies, which can be mentioned as anticancer, antioxidant, antibacterial, antiviral, antifungal and other activities (Gupta et al. 2021). Pyrogallol, through hydroxyl groups attached to its aromatic ring, has a high tendency to form non-covalent interactions such as hydrogen bonds and hydrophobic with a variety of biological macromolecules, including proteins (Shin et al. 2019).

In this study, from the phenolic compound of pyrogallol was used with aim to investigate its anti-amyloidogenic and destabilizing effects on the process of fibrillation and defibrillation of human insulin (HI) protein in vitro by several techniques such as CR absorbance, ThT and ANS fluorescence, FTIR spectroscopy and atomic force microscopy.

Materials And Methods

Protein and chemicals

Human insulin, Thioflavin T (ThT), 1-anilino naphthalene 8-sulfonic acid (ANS), Congo red (CR) were purchased from Sigma-Aldrich (St. Louis, MO, USA). Pyrogallol and all salts and organic solvents were obtained from Merck (Darmstadt, Germany).

Preparation of insulin solution for fibril formation
HI fibrils were prepared according to Amini et al. (2013) method. HI (4 mg/ml) was dissolved in a 50 mM glycine – HCl buffer (pH 2.0) containing 0.02% NaN₃. Then, it was incubated in the presence and absence of different concentrations of Py (in ratios of 0.1, 0.5, 1, and 5 Py to 1 protein in terms of mM) at 50°C without agitation.

To investigate the destabilizing effects of Py on HI amyloid fibrils, preformed fibrils in the presence and absence of different concentrations of Py (in ratios of 0.1, 0.5, 1, and 5 Py to 1 protein in terms of mM) were incubated at 37°C for 3 days.

The formation of HI fibrils was monitored by CR and FTIR spectroscopy, ThT and ANS fluorescence intensity, and atomic force microscopy.

**Congo red (CR) binding assay**

CR binding assay is mainly used to identify amyloids *in vitro*. CR was dissolved in potassium phosphate buffer (5 mM, 0.15 M NaCl, 0.02% NaN₃, pH 7.4) and filtered. This stock solution (2 mM) was diluted to a final concentration of 12.5 µM and used as a working solution. 5 µl of each sample solution was added to 195 µl of CR working solution and incubated at room temperature and in the dark for 30 minutes. The spectrum of CR was measured at a wavelength of 400–700 nm using an Epoch microplate reader (BioTek, USA).

**Thioflavin T (ThT) fluorescence assay**

ThT fluorescence was also used as a reliable method in identifying HI amyloid fibrils and also evaluating the effect of Py on HI fibrillation. After binding of ThT to fibrillar structures, dye fluorescence intensity significantly increases (Biancalana and Koide 2010).

ThT was dissolved in sodium phosphate buffer (10 mM, 0.15 M NaCl, 0.02% NaN₃, pH 7.0) and filtered. This stock solution (2 mM) was diluted to a final concentration of 20 µM and used as a working solution. 15 µl of each sample solution was added to 585 µl of ThT working solution and incubated at room temperature for 2 minutes. Then, the ThT fluorescence intensity was measured with excitation at 440 nm and emission at 488 nm and the slit widths for excitation and emission 5 nm and 10 nm, respectively, using the Cary-Eclipse fluorescence spectrophotometer (Agilent, USA).

To obtain the sigmoid curve, the data collected from ThT fluorescence measurements at 485 nm were fitted by the following equation (Esmaeili et al. 2020):

\[
F = \frac{F_{\text{max}}}{1 + e^{-\left(\frac{t-t_0}{\tau}\right)}}
\]

Where F is the fluorescence intensity at time t, t is the incubation time, \(F_{\text{max}}\) is the fluorescence intensity at the final time and \(t_0\) is the time to reach 50% of the maximum fluorescence. The value of \(\tau\) (the time
constant of fibril growth) was obtained by nonlinear regression. Kapp (the apparent growth rate constant of fibril) and lag phase time were obtained from \(1 / \tau\) and \(t_0 - 2\tau\), respectively.

The inhibition percentage was calculated by the following equation (Phan et al. 2019):

\[
\text{Inhibition percentage} = \frac{I_{\text{control}} - I}{I_{\text{control}}} \times 100
\]

Where \(I\) is the final ThT fluorescence intensity.

**Anilinonaphthalene-8-sulfonic acid (ANS) fluorescence assay**

ANS is generally used to identify partially folded intermediates and to investigate the surface hydrophobicity of proteins. The fluorescence intensity of this dye is significantly increased due to its binding to hydrophobic parts in proteins (Sulatsky et al. 2020). Therefore, ANS fluorescence is a very suitable method to identify amyloid structures due to their high surface hydrophobicity.

ANS was dissolved in sodium phosphate buffer (10 mM, 0.15 M NaCl, 0.02% NaN₃, pH 7.0) and filtered. This stock solution (2 mM) was diluted to a final concentration of 20 µM and used as a working solution. 15 µl of each sample solution was added to 585 µl of ANS working solution, mixed, and incubated at room temperature for 1 minute. The intensity of ANS fluorescence was measured with excitation at 380 nm and emission at 420–600 nm and the slit widths for excitation and emission 5 nm and 10 nm, respectively, using the Cary-Eclipse fluorescence spectrophotometer (Agilent, USA).

**Fourier transform infrared (FTIR) spectroscopic assay**

To investigate changes in the secondary structure of the protein, according to Kawasaki et al. (2014) method, Protein samples were mixed with completely dry KBr and compressed into a thin disc using a tablet press. Then all spectra were recorded from 400 cm\(^{-1}\) to 4000 cm\(^{-1}\) at a resolution of 4 cm\(^{-1}\) with 16 scans and by the 8400S spectrometer (Shimadzu, Japan).

**Atomic force microscopy (AFM)**

To investigate the effects of Py on the morphology of HI amyloid fibrils, 5 µl of each sample diluted with deionized water was dried on a mica plate in the presence of air. After drying, the samples were imaged in the non-contact AFM (NC-AFM) imaging mode at a scan frequency of 0.5 Hz using Full Plus AFM (Ara Pajoohesh, Iran).

**Docking studies**

For molecular docking analysis, the structure of HI was obtained from the protein data bank (http://rcsb.org/) (PDB id code: 1GUJ) and the SDF file of Py was obtained from PubChem database (http://pubchem.ncbi.nlm.nih.gov/) (PubChem CID: 2519). Protein docking was performed using Autodock v1.5.6 software. Discovery studio and VMD v1.9.3 program software were also used to prepare
the 2D and 3D schematic diagrams of the docking model, as well as to exhibit different orientations between ligand and the protein.

**Statistical analysis**

Results were presented as mean ± standard deviation (SD). All experiments were performed at least three times and one-way analysis of variance (ANOVA) was used. *P*-value < 0.05 was considered significant. GraphPad Prism software version 9.2.0 was used for statistical analysis.

**Results**

**Investigation of the effects of Py on the process of HI fibrillation**

Effect of Py on the kinetics of HI fibril formation

To investigate the effect of Py on the kinetics of HI fibril formation, ThT fluorescence of HI incubated in the absence and in the presence of different concentrations of Py was measured at different times. The sigmoid curve obtained in this study showed that HI fibrillation follows the nucleation-dependent elongation mechanism (Nielsen et al. 2001).

According to the results presented in Fig. 1 and Table 1; the lag phase time of HI incubated in the absence of Py is about 47.29 hours, after which the intensity of ThT fluorescence increases strongly and enters the fibril growth phase. Incubation of insulin in the presence of different concentrations of Py resulted in a reduction of the final ThT fluorescence intensity (increasing the inhibition percentage) in a dose-dependent manner (Table 1). The high dose of Py (HI + Py (1:5)), in addition to having the highest effect on reducing fibril formation, was somewhat more effective in prolonging the lag phase time of this process than other concentrations. In the presence of Py, no effect was observed on reducing the rate of the growth of fibril.

<table>
<thead>
<tr>
<th></th>
<th>Lag time (h)</th>
<th>Inhibition (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HI + Py (1:0)</td>
<td>47.29</td>
<td>-</td>
</tr>
<tr>
<td>HI + Py (1:0.1)</td>
<td>48.33</td>
<td>23.79</td>
</tr>
<tr>
<td>HI + Py (1:0.5)</td>
<td>50.64</td>
<td>36.97</td>
</tr>
<tr>
<td>HI + Py (1:1)</td>
<td>52.70</td>
<td>52.40</td>
</tr>
<tr>
<td>HI + Py (1:5)</td>
<td>56.28</td>
<td>67.44</td>
</tr>
</tbody>
</table>
CR, ThT, ANS and FTIR spectroscopy

According to the results presented in Fig. 2A; the CR absorbance intensity of incubated HI in the absence of Py increased and its maximum optical absorption has red-shifted from the wavelength of 490 nm to the wavelength of 500 nm. When insulin was incubated in the presence of different concentrations of Py, a decrease in the CR absorbance intensity was observed, dose-dependently. In the ratios of 1:1 and 1:5 HI to Py, in addition to a significant decrease in the absorbance intensity, a blue shift (shift maximum absorption from 500 nm to 490 nm) was also observed. Gallic acid (Ga) is a polyphenol composed of trihydroxy benzene (Py) and a carboxyl group. In order to compare the inhibitory effect of Ga with Py due to the structural similarity of these two compounds and to understand whether the carboxyl group has an effect on inhibiting the fibrillation process, a ratio of 1:5 HI to Ga was prepared and its effect on the fibrillation process was investigated. As shown in Fig. 2A; it was observed that Ga reduced the CR absorbance intensity of incubated HI by approximately as much as the ratio of 1:5 HI to Py.

According to the results presented in Fig. 2B; the ThT fluorescence intensity of incubated HI in the absence of Py at 485 nm compared to pre-incubation HI (native HI) increased strongly. This result confirms the results of the CR assay. In the presence of different concentrations of Py, the ThT fluorescence intensity of incubated HI decreased in a dose-dependent manner. In addition, The ThT fluorescence intensity of HI samples incubated with Ga at the same concentration decreased and confirms the CR assay results.

According to the results presented in Fig. 2C; the ANS fluorescence intensity of pre-incubation HI is low due to the hydrophobic regions buried inside the protein structure and the lack of ANS access to them. When HI is incubated under acidic conditions at 50°C the ANS fluorescence intensity was significantly increased. In the investigating of samples containing Py, it was observed that with increasing the concentration of this compound, its effectiveness in reducing the ANS fluorescence intensity of incubated HI increases. The same decrease in ANS fluorescence intensity of incubated HI by Py and Ga also indicates a similar effect of these two compounds in reducing exposure of hydrophobic regions on the protein surface.

According to the results presented in Fig. 2D; The FTIR spectrum of pre-incubation HI showed an amide I band in region 1655 cm⁻¹. After incubation, displacement was observed in the amide I band position and the FTIR spectrum of HI (HI fibril) showed the amide I band in the 1630 cm⁻¹ region. According to previous reports (Shivu et al. 2013), the range 1653 – 1660 cm⁻¹ is attributed to the α-helix conformation and the range 1615 – 1643 cm⁻¹ is attributed to the β-sheet conformation. It was also reported that the β structure of amyloid fibrils has been observed in the range of 1628–1632 cm⁻¹. Therefore, FTIR measurement confirms the change in HI conformation from α-helix to β-sheet and thus the presence of fibrillar structure in the sample after incubation. The FTIR spectrum of HI + Py in ratios of 1:1 and 1:5 showed the position of amide I band in region 1626 cm⁻¹ which is in the range of β structure of amorphous aggregates (1624–1630 cm⁻¹) (Shivu et al. 2013).
Investigation of destabilizing effects of HI amyloid fibrils by Py

Pre-formed HI fibrils were incubated in the absence and in the presence of different concentrations of Py for 3 days at 37°C and then their structural changes were evaluated by CR, ThT, ANS and FTIR spectroscopy.

According to the results presented in Fig. 3A and Fig. 3B; The CR absorbance and ThT intensity of Py-free fibrillar samples increased. In contrast, samples containing Py showed a reduction in the CR absorbance and ThT intensity, dose-dependently. In addition, in the study of CR absorbance, the ratios of 1:1 and 1:5 HI fibril to Py showed a blue shift (shift maximum absorption from 500 nm to 490 nm).

According to the results presented in Fig. 3C; in the investigating of the surface hydrophobicity of pre-formed HI fibrils by ANS fluorescence, it was found that all Py concentrations were effective in reducing the ANS intensity of fibrils and this effect was in a dose-dependent manner.

According to the results presented in Fig. 3D; Compared with the FTIR spectrum of HI fibril which showed the amide I band in the 1630 cm$^{-1}$ region, the FTIR spectrum of the HI fibril + Py in the ratios of 1:1 and 1:5 showed the position of amide I band in region 1624 cm$^{-1}$ which is in the range of β structure of amorphous aggregates (1624–1630 cm$^{-1}$) (Shivu et al. 2013).

**AFM scanning**

AFM was used to investigate the effects of Py on the structural morphology of HI in fibrillation and defibrillation processes.

According to the results presented in Fig. 4A; after exposure to acidic pH and incubation at 50°C, twisted and long amyloid fibrils were observed in Py-free samples, however, as shown in Fig. 4B, in the presence of Py, no fibrillar structure was observed and the images showed granular structures instead of fibrillar morphology. It is possible that Py redirects the protein fibril formation pathway toward the creation of amorphous aggregation because it is observed that the amorphous aggregation in the electron microscope images has a granular appearance (Invernizzi et al. 2012). These results and the data of other tests performed in this study confirm the anti-amyloidogenic properties of Py.

As shown in Fig. 4C, Py also successfully acted in the elimination of amyloid fibrils and leads to a change in the fibrillar structure of the protein and probably the creation of amorphous aggregation.

**Docking studies**

AutoDock software was used to better understand the interactions between HI and Py as well as to determine possible connection locations.
According to the results presented in Fig. 5; the amino acid residues of Gly1A, Ile2A, Val3A, Tyr19A, Tyr26B, and Thr27B interact with the ligand through hydrophobic interactions, and Thr27B also forms a hydrogen bond with the ligand (Length of hydrogen bond 2.93 Å).

Since the GFFYTPKT (Residues 23–30) peptide fragment of the insulin B chain helps to create aggregation (Das and Bhattacharyya 2017), Py reacted with the two amino acid residues in this fragment (Tyr26B and Thr27B), whereby It can cause disruption in the process of fibril formation. Despite the lack of interaction of Py with the LVEALYL (Residues 11–17) peptide fragment of the insulin B chain which has a very high tendency to form amyloid (Tartaglia et al. 2008; Sawaya et al. 2007), it was shown that Py has a significant effect in preventing the formation of HI amyloid aggregation. In the study of molecular docking of ascorbic acid (Alam et al. 2017) and peptides P4 and P5 (Siddiqi et al. 2018) also no interaction with this amyloidogenic fragment of HI was observed and the researchers concluded that the reaction of other amino acid residues with inhibitors leads to the prevention of their participation in the interactions that cause the creation of amyloid structures.

**Discussion**

Targeting protein aggregates by small organic molecules such as polyphenols is one of the most desirable and effective strategies to prevent and improve amyloid disease, which has received much attention in recent years. So far, the anti-amyloidogenic activity of many polyphenols, including quercetin (Wang et al. 2011), myricetin (Zelus et al. 2012), silibinin (Cheng et al. 2012), curcumin (Ono et al. 2004a; Pandey et al. 2008), tannic acid (Ono et al. 2004b), resveratrol (Feng et al. 2009), etc. have been investigated. Phenolic compounds in interaction with peptides and proteins, modify their structural properties and prevent amyloid aggregates and their toxicity. In addition, the antioxidant properties of these compounds play an important role in inhibiting amyloid fibrils (Sgarbossa 2012; Porat et al. 2006; Stefani and Rigacci 2013). It is said that the number of hydroxyl groups and the type of side-chain attached to the aromatic ring in the structure of polyphenols play an important role in the amount of possible binding to proteins so that with increasing the number of hydroxyl groups, the tendency of polyphenols to interact with proteins will be higher (Ozdal et al. 2018). In addition, reports indicate that prevention of π-π interaction and inhibition of fibrillation process is due to interactions between the phenolic rings of polyphenols and the aromatic residues of amyloidogenic proteins (Ngoungoure et al. 2015). Among the non-covalent interactions that may occur between polyphenols and proteins, hydrophobic interactions and hydrogen bonds are known as the main interactions and are of great importance (Ozdal et al. 2018). The results of molecular docking simulation in this study showed that Py was able to interact with HI through hydrophobic interactions and hydrogen bonding, thereby resulting in the inhibition of HI fibrillation. According to the results, it can be concluded that the presence of an aromatic ring and its attached hydroxyl groups in the Py structure is the reason for the potential of this phenolic compound to interact with the protein.

In this study, HI protein was exposed to pH 2.0 and a temperature of 50°C. These conditions lead to the opening of the folded structure of the protein and the formation of amyloid aggregates. The acidic pH
makes the protein highly vulnerable and heat is the main factor in the dissociation of the dimer and the production of monomeric protein (Whittingham et al. 2002). It breaks hydrogen bonds, salt bridges, and disulfide bonds that play an important role in maintaining the native structure of the protein (Arora et al. 2004) and thus leads to exposure of hydrophobic regions. On the other hand, heating, by increasing the surface hydrophobicity of the protein, provides more binding sites for polyphenol, in which the hydrophobic bonds contribute significantly. It is said that the most interaction between protein and polyphenol is the hydrophobic interactions (Siebert et al. 1996).

Results of our experimental studies (CR absorbance, ThT and ANS fluorescence intensity) showed that Py has significant inhibitory potential against amyloid fibrils formation and is very effective in reducing the exposure of hydrophobic regions and thus reducing the tendency of the protein to form amyloid fibrils. This effect, like the inhibitory effect of many polyphenols, including gallic acid (Jayamani and Shanmugam 2014), silibinin (Katebi et al. 2018) and curcumin (Rabiee et al. 2013) was observed in a dose-dependent manner. In addition, our findings in the study of the effect of Py destabilizing on the process of HI defibrillation indicated the elimination of fibrillar structures and reduction of surface hydrophobicity of protein is due to the disappearance of hydrophobic regions. Also, in this process, with increasing Py concentration, the rate of defibrillation increased.

The inhibitory effect of Py on other amyloidogenic proteins has also been investigated. According to Di Giovanni et al. (2010), Py is effective in preventing the aggregation of α-synuclein and inhibiting the conversion of β-amyloid protofibrils to mature fibrils.

It has been reported that main factor in the inhibitory properties of polyphenols with similar chemical structure is the phenolic hydroxyl group (Liu et al. 2013). Our results confirmed this claim. The samples containing HI + Ga(1:5) and the sample containing HI + Py(1:5) showed the same inhibitory effect, so the structural difference between the two compounds (carboxyl group) has no effect on inhibiting the HI fibrillation process. These results support the Jayamani and Shanmugam (2014) hypothesis that the carboxyl group is not effective in inhibiting fibril formation. In fact, the trihydroxy benzene group in Ga is the major component of the inhibitory effect in this compound. In a study by Konar et al. (2018), the inhibitory effect of several phenolic compounds including benzoic acid (BA), 4-hydroxy benzoic acid (4-HBA), 3,4-dihydroxy benzoic acid (3,4-DHBA), 3,5-dihydroxy benzoic acid (3,5-DHBA), and Py, which their structures are close to the structure of Ga, were investigated on HEWL fibrillation. It was reported that Ga and Py had the greatest inhibitory effect compared to other compounds. The presence of more hydroxyl groups in Ga and Py is the most significant reason for the high inhibitory effect of these two compounds, compared to other phenolic compounds. On the other hand, the similar inhibitory effect of Ga and Py, as well as the ineffectiveness of benzoic acid (which has only one carboxyl group in its structure) in the process of preventing the formation of HEWL fibrils, were other results obtained in this study which led to the conclusion that the carboxyl group had no significant effect on the inhibitory properties of polyphenols.
In another study, the effect of hydroxyl groups of three polyphenolic compounds, myricetin, morin, and flavone, on the elimination of amyloid fibrils was investigated (Gargari et al. 2018). These three compounds have a similar structure and the only difference is in the number of hydroxyl groups. The results of this study demonstrated that myricetin, with six hydroxyl groups, and morin, with five hydroxyl groups, were more effective at destabilizing amyloid fibrils than flavone, which lacked the hydroxyl group. According to Ono et al. (2006), the number of hydroxyl groups is directly related to the anti-amyloidogenic effects of polyphenolic compounds.

Investigation of microscopic images of the anti-amyloidogenic effects of small organic molecules on amyloid aggregates in many studies showed that some compounds belonging to this group of inhibitors, in addition to preventing fibril formation, can also redirect amyloidogenic proteins toward the creation of non-fibrillar aggregates. Wang et al. (2012) in the study of the effect of Epigallocatechin-3-gallate (EGCG) on human insulin fibrillation found that EGCG in physiological conditions, in addition to having high inhibitory effects, leads to a change in the path of aggregation and the formation of globular aggregates. In a study on inhibitory and destabilizing effects of quercetin against bovine insulin fibrillation (at acidic pH and high temperature) and its defibrillation it was shown that this compound leads to the formation of amorphous aggregates both during inhibition of fibrillation and in the process of elimination of amyloid fibrils (Wang et al. 2011). In other studies, the effect of silibinin (Katebi et al. 2018), curcumin (Rabiee et al. 2013) and ferulic acid (Jayamani et al. 2014) on the formation of bovine insulin fibril also indicated the presence of amorphous species instead of fibrillar morphology in samples containing these compounds. Our observations in this study are consistent with the above reports, as in the microscopic images obtained, the presence of non-fibrillar species and similar to amorphous aggregates in both the inhibitory and destabilizing effects of Py were clearly observed. The results of FTIR spectroscopy also indicate a high probability of amorphous aggregation formation during inhibition and treatment of HI amyloid fibrils by Py.

Totally, based on various studies it can be concluded that polyphenols prevent the formation of amyloid species in different ways. For example, a group of polyphenols, despite being successful in inhibiting the formation of oligomers, lead to the formation of fibrils. Another group prevents the formation of fibrils but has no effect on inhibiting the formation of oligomers. There are also polyphenols that are capable of preventing the formation of both oligomeric and fibrillar forms. Many of them cause a change of the direction of fibrillation and the creation of non-fibrillar oligomers instead of fibrillar forms (Nguoungoure et al. 2015). Regarding these facts, the structure of the compound is very important in determining how it affects the process of amyloid formation.

**Conclusion**

In this study, it was found that Py has a very acceptable function both as an inhibitor in preventing the formation of amyloid fibrils and as a destabilizer in eliminating pre-formed amyloid fibrils. The phenolic structure of Py seems to play an important role in creating effective interactions with the protein. Since various studies have shown that natural polyphenols have special properties to prevent amyloid disease,
the present study could be useful in advancing the design purposes of new anti-amyloid drugs in the future. In addition, more studies are needed to investigate the toxicity of Py in cellular conditions and then to investigate the anti-amyloidogenic properties of Py in animal models in order to prove its medicinal potential.

Declarations

**Ethical approval** Not applicable.

**Consent to participate** Not applicable.

**Consent to Publish** Not applicable.

**Author Contributions** Perform experiments, analyze data and write the paper, P.S.; project designer, Editing and Supervision, S.B.; project adviser, S.H.H. The authors declare that all data were generated in-house and that no paper mill was used.

**Funding** This study was supported by Lorestan University (Lorestan, Iran).

**Competing interests** The authors declare no competing interests.

**Availability of data and materials** This study's data are included in the article, and the corresponding author can provide the primary data.

References


Figures

![Figure 1](image-url)
Effects of pyrogallol on the fibrillation kinetics of human insulin. ThT fluorescence intensity of human insulin was recorded at 485 nm in the absence and in the presence of different concentrations of pyrogallol at pH 2.0 and 50 °C at different times.

Figure 2

Evaluation of pyrogallol effects on the human insulin fibrillation. Protein samples were tested in the absence and in the presence of different concentrations of pyrogallol after the incubation at pH 2.0 and at 50 °C. (A) CR test, (B) ThT fluorescence test, (C) ANS fluorescence test and (D) FTIR spectroscopic test.
In statistical analysis (B,C), pre-incubation human insulin (native human insulin) was considered as a control to compare other groups with it. In this comparison, \( P<0.0001 \) was for all groups. The results are the average of three independent experiments.

**Figure 3**

Evaluation of pyrogallol effects on the human insulin debrillation. Samples containing pre-formed human insulin fibrils were tested in the absence and in the presence of different concentrations of pyrogallol after incubation for 3 days at 37 °C. (A) CR test, (B) ThT fluorescence test, (C) ANS fluorescence test and (D) FTIR spectroscopic test. In statistical analysis (B,C), human insulin fibril in the
absence of pyrogallol was considered as a control to compare other groups with it. In this comparison, $P<0.0001$ was for all groups. The results are the average of three independent experiments.

**Figure 4**

Evaluation of pyrogallol effects on the human insulin morphology using AFM. (A) 2D image and (B) 3D image of pyrogallol-free human insulin fibrils in the fibrillation process (after incubation at pH 2.0 and 50 °C), (C) 2D image and (D) 3D image of human insulin in the presence of pyrogallol (HI+Py (1: 5)) in the
fibrillation process (after incubation at pH 2.0 and 50 °C), (E) 2D image and (F) 3D image of pre-formed human insulin fibrils in the presence of pyrogallol (HI fibril +Py (1: 5)) in defibrillation process (after incubation at 37 °C for 3 days)

Figure 5

Evaluation of possible interactions between insulin and pyrogallol using molecular docking. (A, B) Schematic structure of the binding state of the insulin-pyrogallol complex, (C) An accurate view of the state of the insulin-pyrogallol complex

Supplementary Files

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- Graphicalabstract.jpg
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