Article

Association Mechanism and Conformational Changes in Trypsin on Its Interaction with Atrazine: A Multi- Spectroscopic and Biochemical Study with Computational Approach

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Abstract: Atrazine (ATR) is a herbicide globally used to eliminate undesired weeds. Herbicide usage leads to various adverse effects on human health and the environment. The primary source of herbicides in humans is the food laced with the herbicides. The ATR binding to trypsin (TYP) was investigated in this study to explore its binding potential and toxicity. In vitro interaction of ATR with TYP was studied using multi-spectroscopic methods, molecular docking, and enzyme kinetics to explore the mechanism of binding for the TYP-ATR system. The TYP-ATR complex revealed binding constants (10^3 M^{-1}), suggesting a moderate binding. The free energy for the TYP-ATR complexes was negative, suggesting a spontaneous interaction. Thermodynamic parameters enthalpy (\( \Delta H \)) and entropy (\( \Delta S \)) obtained positive values for the TYP-ATR system suggesting hydrophobic interactions in the binding process. Micro-environmental and conformational changes in TYP molecules were induced on interaction with ATR. Reduced catalytic activity of TYP was observed after interaction with ATR owing to the changes in the secondary structure of the TYP.

Keywords: atrazine; trypsin; activity; fluorescence; quenching; docking

1. Introduction

Herbicide usage is widely practiced in crops to eliminate unwanted plants and weeds, thus essential in food production sustainability [1,2]. However, herbicides are highly toxic chemicals to all organisms, resulting in adverse effects on humans and the environment since they persist in nature for more extended periods [3–6]. Chemically atrazine (ATR) is 2-chloro-4-ethylamine-6-isopropylamine-1,3,5-triazine (Figure 1) and belongs to the triazine family. An estimated 5.6 billion pounds of herbicides are used worldwide annually [7]. The ATR usage increases production by controlling broadleaf and grassy weeds [8,9]. ATR is highly persistent in the environment and has a half-life of 100 days in surface water. It is commonly detected in contaminated groundwater and waterways [10,11]. Therefore, long-term exposure to ATR can cause adverse effects on human health [12]. Several animal studies confirm that ATZ has endocrine-disrupting ability across multiple animal classes [13,14]. However, few studies reported that ATR inhibits the specific binding of various types of protein receptors (estrogens and progesterone) and therefore interferes with the reproductive health of animals [15,16]. ATZ exposure is also reported to cause epigenetic alterations in DNA methylation and microRNA activity [17,18].
In recent years, proteins have been the main investigational target to study the toxicity of environmental pollutants, especially pesticides, which induce changes in the protein’s structural (conformation) and biological function [19–21]. Proteases are essential proteins that play various roles (catalytic ability during physiological and pathological processes) in different biologically relevant processes [22–25]. Trypsin (TYP), a 23.3 kDa protein containing 223 amino acids, 4-tryptophan, 10-tyrosine, and 6-phenylalanine, is a vital protease (serine-endopeptidase) in the digestive system (pancreas of all vertebrates). Trypsin is essential for multifarious biological functions, including digestion and deconstruction of food proteins and other essential physiological processes [22–24]. The catalytic activity and conformational changes of TYP are affected by exogenous environmental pollutants that cause pathological alterations in humans [26,27]. Since no study has been conducted on TYP interaction with ATR, we investigated the interaction mechanism between them and evaluated the conformational changes in TYP. This study provided an insight into the toxicity mechanism of ATR at the molecular level. The influence of ATR on the conformations and catalytic activities of TYP was also explored in this study. Spectroscopic techniques, molecular modeling, and TYP enzyme activity experiments were conducted to comprehend the toxicity mechanism of ATR on the TYP.

2. Materials and Methods

2.1. Materials

Trypsin (bovine pancreas) ATR, N-α-Benzoyl-D, L-arginine-P-nitroanilide hydrochloride in (BApNA) were all obtained from Sigma-Aldric Co. (St. Louis, MO, USA). Analytical grade solvent and chemicals purchased locally were used in the study.
2.2. Sample Preparation

The TYP solution (0.25 mM) was prepared in phosphate-buffered saline (0.1 M containing dibasic (Na$_2$HPO$_4$), monobasic (NaH$_2$PO$_4$), and NaCl) stock solution of ATR (0.5 mM) in anhydrous methanol and BApNA (3 mM) was prepared in DMSO. All the reagents were stored at 2–8 °C until use. Buffers were prepared in ultra-pure water.

2.3. Instrumentations

The UV-vis absorption spectra were recorded with a UV-1800 double beam spectrophotometer (Shimadzu, Kyoto, Japan). The fluorescence measurements were recorded with an RF-5301PC spectrofluorometer (Shimadzu, Kyoto, Japan). The circular dichroism (CD) measurements were recorded on a JASCO J-1500CD spectrophotometer. FT-IR measurements were carried out on Avatar 360 E.S.P. FT-IR spectrometer (Thermo, Montral-Est, QC, Canada)

2.4. Methods

2.4.1. UV-Vis Spectral Measurements

The UV-vis spectra of TYP were recorded in the presence of ATR (0–30 µM). Experiments were performed at fixed TYP concentration (10 µM).

2.4.2. Steady-State Fluorescence Spectroscopy

Fluorescence emission spectra of TYP (10 µM) titrated against ATR (0–45 µM) at 298, 303, and 308 K were recorded. The emission wavelength range was set to 300–500 nm after excitation at 280 nm.

2.4.3. Synchronous Fluorescence Spectroscopy (SFS) Experiments

Synchronous fluorescence spectra measurements of TYP (10 µM) titrated with increasing concentrations of ATR (0–45 µM) were recorded in separate experiments by setting wavelength interval (Δλ) at 15 nm (for tyrosine residues (Y)) and 60 nm (for tryptophan residues (w)) in the same experimental conditions as the steady-state fluorescence emission spectra.

2.4.4. Circular Dichroism (CD) Spectroscopy Measurements

The CD spectra of TYP (5 µM) with ATR (0 and 10 µM) were recorded from 190–260 nm with a scan speed of 20 nm min$^{-1}$ and 1 nm bandwidth. The recorded spectrum was analyzed by the web-based software package tool, CAPITO-CD Analysis and Plotting Tool (uni-jena.de, accessed on 10 April 2022), to evaluate different secondary structures of TYP after being treated with ATR.

2.4.5. Fourier Transform Infrared Spectroscopic (FTIR) Analysis

The FTIR spectra for TYP were recorded with ATR. The FTIR spectroscopy of free TYP (20 µM) and TYP (20 µM)-ATR (100 µM) were recorded in the range of 1800–1400 cm$^{-1}$. The corresponding absorbance of buffer and ATR solutions recorded under similar conditions was subtracted from the spectra.

2.4.6. Computational Analysis

The molecular structure of TYP (PDB ID: 2ZQ1) was obtained from Protein Data Bank (RCSB), http://www.rcsb.org/pdb, accessed on 3 March 2022). Molecular modeling of the ATR–TYP complexation was performed using AutoDock Vina. For docking purposes, water, metal ions, and the initially bound ligand were removed, and hydrogen atoms were added during protein preparation. The pdb structure of ATR was obtained from Pubchem (ID: 2256), and its energy-minimized conformation was analyzed using Chem3D Pro14.0. During the docking study, a grid box defined to enclose the entire binding site of trypsin with dimensions of 40 Å × 40 Å × 40 Å and a grid spacing of 0.375 Å was used. The grid
center of the TYP-ATR system was set at $-26.319, 1.973, \text{and } 16.726$. The docked structure of the TYP-ATR system was analyzed with Discovery studio 4.5.

2.4.7. Measurement of Trypsin Activity

The activity of TYP was measured using BApNA as a substrate according to the reported literature [28–30]. The absorbance at 400 nm was recorded to evaluate the TYP activity. The relative activity of TYP was expressed as the ratio of activity with and without ATR. The kinetic parameters were determined using a fixed concentration of TYP (5 µM), and the ratios for TYP: ATR were (1:0, 1:5, 1:10, 1:15, and 1:20), BApNA (0–3000 µM), The Michaelis-Menten equation was used in the determination of enzyme reaction velocity [29].

\[
V = V_{\text{max}} \frac{[S]}{K_m + [S]}
\]

\[
K_{\text{cat}} = \frac{V_{\text{max}}}{[E]}
\]

The above equations, $V$ and $V_{\text{max}}$, denote the initial and final reaction velocities, $K_m$ is Michaelis Constant, and $K_{\text{cat}}$ is the catalytic rate constant. Whereas $[E]$ represents enzyme concentration and $[S]$: substrate concentration (BApNA). Lineweaver-Burk plots for $1/V$ vs. $1/[S]$ at different BApNA concentrations in the absence and presence of ATR.

3. Results and Discussion

3.1. UV-Vis Absorption Spectroscopy Investigations

UV-vis absorption spectroscopy is a simple and versatile technique for studying the conformational and structural changes in a protein and also explores the protein-ligand complex formation [31]. The UV-vis absorption spectra of TYP and TYP-ATR are given in Figure 2. The TYP usually shows two absorption bands, at around 205 nm, and 275 nm. The peak at 205 nm corresponds to peptide bond absorption (secondary structures), and the one at 275 nm represents aromatic amino acids, respectively [32]. A sharp decline in the peak at 205 nm suggests that ATR and TYP interaction might influence the conformation of TYP. The peak at (205 nm) is mainly due to $\pi-\pi^*$ transitions and corresponds to the polypeptide backbone structure (C=O) of the TYP [33]. Further, a shift in the peak at 205 nm suggests a new complex formation between TYP and ATR. A decline in the absorption at 275 nm also suggests a complex formation occurred between TYP and ATR.

Moreover, the absorbance intensity at 275 nm also decreased, which confirms that the polarity of the microenvironment around the aromatic amino acid residues was affected by the ATR presence [33]. Therefore, the change in absorption peaks and red-shift suggest a conformation change in TYP and complex formation between TYP-ATR and hence a change in the protein structure.
Moreover, the absorbance intensity at 275 nm also decreased, which confirms that the polarity of the microenvironment around the aromatic amino acid residues was affected by the ATR presence [33]. Therefore, the change in absorption peaks and red-shift suggest a conformation change in TYP and complex formation between TYP-ATR and hence a change in the protein structure.

3.2. Fluorescence Emission Spectroscopy of TYP-ATR Complex

Fluorescence emission spectroscopy is a technique to study the protein-ligand interactions and conformational alterations [34,35]. In the steady-state fluorescence measurements (Figure 3A), TYP exhibited a strong emission peak at 337 nm after excitation at 280 nm. The intrinsic fluorescence of TYP reduced progressively with the addition and increase in ATR concentration. This perception demonstrated a strong binding in the TYP-ATR system and microenvironmental changes in TYP on treatment with ATR [33].

The fluorescence quenching mechanisms of protein are classified into three mechanisms: static, dynamic, and mixed (static and dynamic) quenching [36]. The quenching mechanism is identified based on their temperature dependence. The fluorescence spectra for the TYP-ATR system were obtained at 295, 305, and 315 K, and the fluorescence data were analyzed with Stern-Volmer equation [37,38]:

\[ F_0/F = 1 + K_{SV}[Q] = 1 + kQ\tau_0[Q] \]

where \( F_0 \) represents the steady-state fluorescence intensity of TYP, \( F \) fluorescence intensity of TYP in the presence of ATR, \( K_{SV} \) is the Stern-Volmer quenching constant, and \([Q]\) is the quencher concentration.

The Stern-Volmer plot (Figure 3B) for the TYP-ATR system at (295, 305, and 315 K) shows a decline in the \( K_{SV} \) values with temperature rise. This behavior of the \( K_{SV} \) values with respect to temperature represents the static quenching mechanism for the TYP-ATR system [37,39].

Figure 2. UV-vis absorbance spectra of TYP (10 µM) in presence of increasing concentrations of ATR (0–30 µM).

Figure 3. (A) Fluorescence emission spectra of TYP (10 µM) in the presence ATR (0–45 µM). (B) Stern-Volmer plot for TYP with ATR (0–45 µM) at different temperatures. (C) The double-logarithmic plot of \([\log (F_0 - F)/F]\) versus \(\log[C]\) for TYP-ATR system. (D) van’t Hoff plots for TYP-ATR system at different temperatures.
3.2.1. Fluorescence Quenching Mechanism

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\[ \frac{F_0}{F} = 1 + K_{SV}[Q] = 1 + k_q \tau_0 [Q] \]

where \( F_0 \) represents the steady-state fluorescence intensity of TYP, \( F \) fluorescence intensity of TYP in the presence of ATR, \( K_{SV} \) is the Stern-Volmer quenching constant, and \([Q]\) is the quencher concentration.

The Stern-Volmer plot (Figure 3B) for the TYP-ATR system at (295, 305, and 315 K) shows a decline in the (Table 1) for the TYP-ATR system with temperature rise. This behavior of the \( K_{SV} \) values with respect to temperature represents the static quenching mechanism for the TYP-ATR system [37,39].

Table 1. The values of Stern-Volmer constant and quenching rate constant for the TYP-ATR system.

| pH  | Temp (K) | \( K_{SV} (\times 10^3 \text{ M}^{-1}) \) | \( k_q (\times 10^{11} \text{ M}^{-1} \text{ s}^{-1}) \) | \( R^2 \) |
|-----|---------|---------------------------------|----------------|------|
| 7.4 | 295     | 4.24                            | 4.24           | 0.998|
|     | 305     | 3.28                            | 3.28           | 0.996|
|     | 315     | 2.49                            | 2.89           | 0.990|

The bimolecular quenching constant (\( k_q \)) also helps to confirm the quenching mechanism. The \( k_q \) was determined by [37,40]:

\[ k_q = K_{SV} / \tau_0 \]

where \( K_{SV} \) is the Stern-Volmer constant; \( \tau_0 \) represents the lifetime (average) of fluorophore without the quencher and has a value of \( 10^{-8} \text{ s} \) [37]. The \( k_q \) for the TYP-ATR system at different temperatures is presented in Table 1. The values of \( k_q \) were found higher than the value of the collision quenching constant \( (2 \times 10^{10} \text{ M}^{-1} \text{ s}^{-1}) \), suggestive of a static quenching mechanism in the TYP-ATR system [41,42].

3.2.2. Binding Constant Evaluation

The binding parameters of the TYP-ATR complex, binding constant (\( K_b \)), and binding stoichiometry (\( n \)) were evaluated by the equation:

\[ \log \left( \frac{F_0 - F}{F} \right) = \log K_b + n \log [Q] \]

where \( F_0 \) and \( F \) are the fluorescence intensities of TYP in the absence and presence of ATR, respectively. The \( K_b \) and \( n \) were determined from the plot between the \( \log (F_0 - F)/F \) versus \( \log [Q] \) (Figure 3C, Table 2). The binding stoichiometry \( n = 1 \) suggests a single class of available binding sites. The binding constants of \( >10^3 \) present a moderate binding between TYP and ATR. Further, the binding constants decreased with a temperature rise.

Table 2. The values of the binding constant and the number of binding sites for the interaction of ATR with TYP.

| pH | Temp (K) | \( K_b (\times 10^3 \text{ M}^{-1}) \) | \( n \) | \( R^2 \) |
|----|---------|---------------------------------|------|------|
| 7.4| 295     | 2.192                           | 0.93 | 0.995|
|    | 305     | 1.974                           | 0.95 | 0.980|
|    | 315     | 1.887                           | 0.98 | 0.996|
3.2.3. Determination of the Binding Forces between TYP and ATR

The intermolecular acting forces, such as hydrophobic, electrostatic interactions, hydrogen bonds, and van der Waals interactions, exist between ligands and macromolecules [43]. The thermodynamic analysis provides evidence about the binding force involved in the interaction. The enthalpy ($\Delta H^o$) and entropy ($\Delta S^o$) are obtained from the slope and intercept of the van't Hoff plot (Figure 3D) using van't Hoff Equation [39]:

$$\ln K_b = -\frac{\Delta H^o}{RT} + \frac{\Delta S^o}{R}$$

$$\Delta G^o = \Delta H^o - T\Delta S^o$$

where $\Delta G^o$ is the binding free energy, R is the gas constant (8.314 J mol$^{-1}$ K$^{-1}$), T represents different temperatures in °K (295, 305, and 315 K), and $K_b$ is the binding constant. The calculated values for $\Delta H^o$, $\Delta S^o$, and $\Delta G^o$ of the TYP-ATR system are given in Table 3. Further, the negative value for $\Delta G^o$ confirms the interaction to be spontaneous, and the positive $\Delta H^o$ and $\Delta S^o$ values indicate hydrophobic interactions exist in the ATR-TYP system [43,44].

Table 3. Various thermodynamic parameters for TYP-ATR complex formation at various temperatures.

| Temp (K) | $\Delta H$ (kJ mol$^{-1}$) | $\Delta S$ (J mol$^{-1}$ K$^{-1}$) | $\Delta G$ (kJ mol$^{-1}$) | $R^2$ |
|---------|-----------------|-----------------|-----------------|------|
| 295     | 5.8             | 44.21           | -7.2            | 0.9954 |
| 305     | -7.6            | 0.9988          |
| 315     | -8.1            | 0.9986          |

3.2.4. Synchronous Fluorescence Spectroscopy (SFS) Experiment

Synchronous fluorescence spectroscopy (SFS) is used to monitor alteration in the microenvironment of fluorophore amino acid residues [45]. The synchronous fluorescence spectra were obtained at $\Delta \lambda = 15$ nm and $\Delta \lambda = 60$ nm [46]. The SFS emission spectra of TYP (10 µM) in the presence of ATR (0–45 µM) are given in Figure 4A,B for tyrosine and tryptophan residues, respectively. The intensity of fluorescence of TYP (around tyrosine residue) decreased in the presence of ATR, and no shift in emission wavelength, confirming no change in the microenvironment around tyrosine residue (Figure 4A). Further, the fluorescence intensity of TYP (around tryptophan residues) decreased significantly in the presence of ATR with a red-shift of 1 nm in the emission wavelength. The redshift is suggestive of an alteration in the microenvironment of tryptophan residues. It is also apparent from Figure 4C that the synchronous fluorescence spectra for both tyrosine and tryptophan are significantly different. Further, the ratios of synchronous fluorescence quenching (RSFQ) were estimated by Equation:

$$RSFQ(\%) = 1 - \frac{F}{F_0}$$

where $F_0$ and $F$ are fluorescence intensities of TYP in the absence and presence of ATR, respectively. To confirm the binding position of the ATR on the surface of TYP, the RSFQ ratio for the TYP-ATR system at tyrosine ($\Delta \lambda = 15$ nm) and tryptophan ($\Delta \lambda = 60$ nm), respectively, were determined according to the above equation (Figure 4D). As a result, the synchronous fluorescence of tryptophan residue is strongly quenched (82% reduction), whereas, in the case of tyrosine residue, the quenching was about 63% in the presence of ATR. Thus, SFS results suggest that ATR affects the microenvironment of the tryptophan residue more than tyrosine on interaction with TYP [47,48].
Figure 4. Synchronous fluorescence spectra of TYP (10 µM) with ATR (0–45 µM) at (A) Δλ = 15 nm, (B) Δλ = 60 nm, pH = 7.4. (C) The quenching of TYP synchronous fluorescence of TYP-ATR system. (D) Comparison of the effect of ATR on the RSFQ of TYP.

3.2.5. Circular Dichroism (CD) Spectra Changes in TYP upon ATR Binding

The CD is a sensitive and powerful technique that explores the protein’s secondary structure, stability, and conformation. It provided information about perturbations in the secondary structure of TYP after treatment with ATR [49]. The spectra of TYP with or without ATR at room temperature (Figure 5A) show a negative peak at 208 nm (π → π* transition), which represents the α-helical structure in TYP [50]. Moreover, addition of ATR decreased the ellipticity of TYP, indicating loss of α-helical contents. The CAPITO-CD analysis software calculated the relative percentages of the secondary structural elements of TYP. The relative percentage of secondary structural elements of TYP (α-helix (21%), β-sheet (11%) and random coil (62%)) and TYP-ART system α-helix (7%), β-sheet (33%) and random coil (56%) (Figure 5B). Therefore, ATR induced changes in the secondary structure of TYP, which may cause a change in the physiological function of TYP-like enzyme activity [51].
Figure 5. (A) The CD spectra of TYP in the presence ATR, and the molar ratio of ATR to TYP were 0:1, 1:1, TYP (10 µM), ATR (10 µM), respectively. (B) Relative percentages of the secondary structure of TYP-ATR system.

3.2.6. FT-IR Spectroscopy

FT-IR spectroscopy is an efficient technique used in detecting changes in the secondary structure of the protein [46]. Proteins exhibit several amide bands (Amide I band (1700–1600 cm\(^{-1}\) for C=O) and (Amide II band (1510–1580 cm\(^{-1}\) for C-N/N-H vibration)), which represents the secondary structure of TYP [46]. The FTIR spectra of TYP in the absence and presence of ATR (Figure 6) show that the peak position of the amide band shifted slightly from 1654 to 1653 cm\(^{-1}\) (Amide I) and 1544 to 1543 cm\(^{-1}\) (Amide II) in the ATR-TYP complexes, respectively. Thus, TYP interaction with amide I and II groups disturb the naturally occurring conformation and secondary structure of TYP [52]. Although the shift in the peak position is minimal, an alteration in the secondary structure of the protein cannot be ruled out.

Figure 6. FTIR spectra of TYP (20 µM)-ATR system. The ATR concentrations were 0 and 100 µM, respectively, for the two spectra.
3.2.7. Computational Modeling of the TYP-ATR Complex

Molecular docking is a computational approach used to determine the mechanism of protein-ligand interactions [53]. The docking between ATR and TYP was performed by using Autodock Vina to determine the binding sites and the binding mode of ATR on TYP [54]. The docked structure of the TYP-ATR system is given in Figure 7A–C. The ATR binds with TYP with the free binding energy of $-4.1$ Kcal mol$^{-1}$. In the binding pocket of TYP, the ATR molecule was surrounded by amino acid residues (Figure 8A,B) Ala-221, Ser-190, Trp-215, Ser-214, Asp-189, Gly-226, Ser-195, Val-227, Gly-216, and Ser-217. The ATR also interacted with CYS-220 and Gly-219 with hydrogen bonds. Further, the His-57 and Cys-191 also interacted with ATR through pi-alkyl interaction. Moreover, the conformation changes caused by ATR binding with binding pockets interact directly with catalytic amino acid residues of the TYP. The results obtained from the docking agreed with our conclusion in the experiments.

Figure 7. (A) Molecular models of TYP complex with ATR. (B,C) provides a detailed view of the docking poses of the TYP-ATR complex.
3.2.8. Effect of ATR on TYP Activity

The spectroscopic and docking experiment results suggest an interaction between ATR and TYP. The relative activity (RA%) of the TYP was evaluated at different concentrations of ATR, and it was observed that the TYP activity decreased with the addition of ATR (Figure 9A). The TYP has three primary catalytic triads (His-57, Asp-102, and Ser-195) and a substrate-binding pocket (Asp-194, Gly-217, and Gly-227) which play a vital role in the binding of substrate to TYP [55,56]. The molecular docking confirmed that ATR interacts with the catalytic triad (His-57) and hence affects the enzymatic activity TYP. Furthermore, these results affirm that ATR interaction may lead to conformational changes in the structure of TYP, reducing its activity.

However, Michaelis-Menten curves (Figure 9B) of TYP suggest a decrease in the enzymatic activity of TYP in the presence of ATR. Additionally, the activity declined further with an increase in ATR concentration. The decreased enzymatic activity is attributed to conformational structure change and the catalytic ability of TYP [57]. The Lineweaver-Burk plots of TYP and TYP-ATR also confirmed the inhibition mode of ATR (Figure 9C). Moreover, it is clear from Table 4 that the $V_{\text{max}}$ value decreased while the $K_m$ values almost remained unchanged on interaction with higher concentrations of ATR. Further, the TYP-ATR complex formation shows a non-competitive inhibition. In addition, the catalytic constant $k_{\text{cat}}$ and $k_{\text{cat}}/K_m$ values decreased gradually at higher concentrations of ATR [57]. Thus, the results suggest that ATR inhibits the enzymatic activity of TYP in a concentration-dependent manner.

**Figure 8.** (A) The two-dimensional image of the TYP-ATR system and the amino acid residue surrounding the binding pocket. (B) Three-dimensional structure of interaction of TYP with ATR.
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Figure 9. (A) Relative activity of TYP in the presence of ATR. (B) Michaelis-Menten curves of TYP with different concentration of ATR. (C) Double-reciprocal plots of TYP with different concentrations of ATR.

However, Michaelis-Menten curves (Figure 9B) of TYP suggest a decrease in the enzymatic activity of TYP in the presence of ATR. Additionally, the activity declined further with an increase in ATR concentration. The decreased enzymatic activity is attributed to conformational structure change and the catalytic ability of TYP [57]. The Lineweaver-Burk plots of TYP and TYP-ATR also confirmed the inhibition mode of ATR (Figure 9C). Moreover, it is clear from Table 4 that the Vmax value decreased while the Km values almost remained unchanged on interaction with higher concentrations of ATR. Further, the TYP-ATR complex formation shows a non-competitive inhibition. In addition, the catalytic constant kcat and kcat/Km values decreased gradually at higher concentrations of ATR [57].

### Table 4. Michaelis-Menten kinetic parameters of TYP in the presence of ATR concentrations.

| TYP/ATR | Vmax (µM s⁻¹) | Km (µM) | Kcat (s⁻¹) | Kcat/Km (µM⁻¹ s⁻¹) |
|---------|---------------|---------|------------|---------------------|
| 1:0     | 5.44 × 10⁻⁴   | 3.89 × 10⁻³ | 3.62 × 10⁻⁵ | 9.33 × 10⁻³         |
| 1:5     | 4.05 × 10⁻⁴   | 3.02 × 10⁻³ | 2.70 × 10⁻⁵ | 8.93 × 10⁻³         |
| 1:10    | 4.14 × 10⁻⁴   | 4.71 × 10⁻³ | 2.76 × 10⁻⁵ | 5.84 × 10⁻³         |
| 1:15    | 3.08 × 10⁻⁴   | 3.81 × 10⁻³ | 2.05 × 10⁻⁵ | 5.40 × 10⁻³         |
| 1:20    | 2.59 × 10⁻⁴   | 4.36 × 10⁻³ | 1.72 × 10⁻⁵ | 3.96 × 10⁻³         |

4. Conclusions

This study provides a detailed binding mechanism of the TYP-ATR system and its effects on the structure conformations and the influence of these conformational changes on the enzymatic activities of the TYP. The results suggest a moderate binding interaction between TYP and ATR, and the binding strength was temperature-dependent. The molecular docking and thermodynamic results suggest the involvement of hydrophobic interaction and hydrogen bonds between TYP and ATR and a single binding site for ATR on TYP. In the TYP-ATR system, the TYP activity was inhibited with ATR in a concentration-dependent manner since ATR was bound to the catalytic site. This study provides a deeper understanding of the mechanism of binding of ATR with the digestive proteases from the point of view of toxicological effects.
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