

Cite this: *RSC Adv.*, 2018, 8, 20295

Studies of the binding properties of the food preservative thiabendazole to DNA by computer simulations and NMR relaxation

Qiaomei Sun,^a Zili Suo,^a Hongyu Pu,^a Peixiao Tang,^a Na Gan,^a Ruixue Gan,^a Yuanming Zhai,^{*b} Xiaohui Ding^a and Hui Li^a

Thiabendazole (TBZ) is a commonly used food preservative and has a wide range of anthelmintic properties. In this study, computer simulations and experiments were conducted to investigate the interaction mechanism of TBZ and herring sperm DNA (hsDNA) at the molecular level. Molecular docking showed that TBZ interacted with DNA in groove mode and bound in A-T and C-G base pair regions. Molecular dynamics (MD) was used to evaluate the stability of the TBZ–DNA complex and found that the three phases in MD and the hydrogen bonds helped maintain the combination. NMR relaxation indicated that TBZ had a certain affinity to hsDNA with a binding constant of 462.43 L mol⁻¹, and the thiazole ring was the main group bound with DNA. Results obtained from fluorescence experiments showed that the binding of TBZ and hsDNA was predominantly driven by enthalpy through a static quenching mechanism. Circular dichroism and viscosity measurements proved the groove binding mode. The FTIR results clarified the conformational changes of DNA, that the DNA helix became shorter and compact, and the DNA structure transformed from B-form to A-form.

Received 30th April 2018
Accepted 24th May 2018

DOI: 10.1039/c8ra03702g

rsc.li/rsc-advances

1. Introduction

Thiabendazole [2-(4-thiazolyl) benzimidazole, TBZ] is a commonly used food preservative that is widely applied in various fruits and vegetables to slow down putridity caused by fungi.¹ The structure of TBZ was shown in Fig. 1. Although the use of TBZ can provide benefits, its unreasonable use or excessive consumption leads to health risks; for example, TBZ can disrupt microtubules and induce anaphase–telophase chromosomal aberrations.^{2,3} Thus, many researchers have focused on the binding properties of food additives and

physiological macromolecules to preliminarily evaluate the safety and physiological properties of food additives.^{4–6}

Deoxyribonucleic acid (DNA) is an important bio-macromolecule that has potential therapeutic activity against many diseases, given that it is the pharmacological target of many drugs.^{7–9} The bindings of ligand and DNA have been indicated to affect DNA duplication and culminating transcription.¹⁰ The non-covalent interaction binding mode between ligands and the double helix of DNA include (i) groove binding where the ligand is located outside of the DNA helix groove with hydrogen bonds and (or) van der Waals interactions, and (ii) intercalation binding between ligand and the stacked base pairs of DNA, which usually have a greater impact on the structure and function of DNA compared with groove binding.^{11,12} The DNA damage and genotoxic and mutagenic effects of some food additives have been recently considered, and some of their adverse effects on DNA have been revealed.¹³ Moreover, there are some studies indicated that the difference in DNA structure had some effect on the binding process of ligand and receptor.^{14,15} Hence, studies on the interaction between food additives and DNA can provide theoretical knowledge for the physiological properties of food additives and are beneficial to promote the development of safe food preservatives.

Receptor–ligand interactions have caught many researchers' attention, especially in the field of chemistry and life sciences. Many methods have developed to investigate these interactions.^{16–18} Among these methods, computer simulations

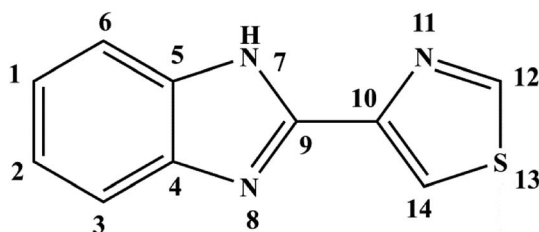


Fig. 1 Structure of thiabendazole (TBZ).

^aSchool of Chemical Engineering, Sichuan University, Chengdu 610065, China. E-mail: tangpeixiao@126.com; Fax: +86 028 85401207; Tel: +86 028 85405220

^bAnalytical & Testing Center, Sichuan University, Chengdu 610065, China. E-mail: yuanmingzhai@scu.edu.cn



including molecule docking and molecular dynamic (MD) simulation, are a powerful tool to determine the orientation and binding site for visualization of the binding properties and to monitor the dynamics behavior of the ligand/receptor structures during combination.^{19,20} Complementarily, nuclear magnetic resonance (NMR) provides reasonable and reliable data to clarify the binding properties of ligand at the molecular level by identifying ligand epitope and structure affinity.^{21,22} Relaxation time technique is an effective method to investigate the binding affinity and dynamic properties on the basis of the quantitative analysis of spin-lattice relaxation rate changes of proton.^{23–25}

Given that the binding mode of TBZ and calf thymus DNA (ctDNA) had been researched by fluorescence, UV absorbance and circular dichroism (CD) spectra,²⁶ the main aim of this work was to further investigate the binding mechanism between TBZ and herring sperm DNA (hsDNA) and probe into the effects on the binding properties for different DNA models by combining theories and experiments. The interaction characteristics of TBZ–hsDNA would be clarified from the point of the superior groups and proton affinity abilities of TBZ. Molecular docking and MD were conducted to estimate the binding mode, binding region, and structural characteristics and to evaluate the dynamics behavior and stability of TBZ–DNA complex. NMR relaxation experiments probed into the binding mechanism (including binding constant and affinity) from the point of ligand molecule. The binding mode and structural feature of TBZ to hsDNA were corroborated by spectroscopy. This study will help us understand the physiological pharmacokinetic behavior of TBZ at the molecular level and will provide interesting insights into the interactions of food additives and DNA.

2. Materials and methods

2.1. Reagents and chemicals

Herring sperm DNA (hs-DNA) was obtained from Solarbio (Beijing, China) and was dissolved in a phosphate buffer solution (PBS, 0.01 M). The purity of the DNA was checked by a ratio of UV absorbance at 260 and 280 nm. The ratio was larger than 1.8, indicating that the hs-DNA was sufficiently free from any protein contamination.²⁷ The stock solution of the DNA was stored in the dark at 4 °C and used within a week.

TBZ was purchased from Aladdin Chemical Reagent (Shanghai, China). All of the other reagents were of analytical grade and used without further purification. Ultrapure water was used throughout the experiment.

2.2. Molecule docking and molecule dynamics simulation

FlexX docking program interfaced within LeadIT was used to examine possible binding modes and active site of TBZ and DNA.²⁰ The 3D structure of TBZ was obtained from PubChem (Pub-Chem CID: 5430), and the crystallographic structure of DNA duplex (PBD ID: 1BNA) was downloaded from PCSB protein data bank with the sequence of (CGCGAATTCGCG)₂. The hydrogen atoms of DNA were added using Receptor Wizard module. The FlexX hybrid algorithm was selected to perform the

docking with a radius of 20 Å binding site definition. The docking binding modes of the top five highest scored docking poses were evaluated, and the optimal binding poses were used for molecular dynamics simulation.

MD simulations were conducted using YASARA v16.7.22 package with the AMBER14 force field, to determine the optimal binding conformation and assess the system stability.¹⁹ The simulation was performed at a NPT ensemble with a constant temperature at 298 K and constant pressure (1 bar) at pH 7.4. Counter ions (Na⁺ or Cl[−]) were added by randomly replacing water molecules to obtain a charge-neutral system. Periodic boundary conditions were applied in the MD process. Simulations were carried out after initial energy minimization procedures were conducted, by using a pre-defined macro (md_run) within the YASARA package. According to the MD trajectories, root mean square deviations (RMSD) were analyzed.

2.3. NMR measurements

All data were acquired using a Bruker Advance 400 MHz NMR spectrometer, operated at 400.13 MHz at 25 °C. The spin-lattice relaxation rates were measured using the standard inversion-recovery $(180^\circ - \tau - 90^\circ - t)_n$ sequence. The τ values used for the experiments were 0.01, 0.02, 0.04, 0.06, 0.1, 0.2, 0.4, 0.8, 1, 3, 5, 7, 20, and 25 s. The delay time t in this case was 25 s. The 180° selective inversion of the proton spin population was obtained through a selective soft Gaussian perturbation pulse (width: 20 ms, power: 60 dB) with an excitation width of about 45 Hz. The addition of HSA did not change the viscosity of the system, and the observed R_1^{se} enhancements were mainly affected by the ligand–protein complex formation. All NMR data processing and analyses were performed using Topspin 2.1 software (Bruker BioSpin, Ltd.).

2.4. Fluorescence spectroscopy

Fluorescence intensities were recorded using a Cary Eclipse fluorophotometer (Varian, USA) equipped with 0.1 cm quartz cells. Each solution was left to stand for 1 hour to achieve equilibrium. The wavelength range from 310 nm to 550 nm with excitation wavelength at 300 nm was selected. The slit widths for both excitation and emission were set to 10.

2.5. Fluorescence lifetime measurements

The fluorescence lifetimes were determined by time-correlated single photon counting (TCSPC) using a Horiba Jobin Yvon FluoroMax-4 spectrofluorometer (HORIBA, FRA). The time-resolved intensity decay of TBZ was measured in the absence and presence of hs-DNA in PBS buffer (pH = 7.4) at an excitation and emission wavelengths of 317 nm and 360 nm. The TBZ concentration was fixed at 2.0×10^{-5} mol L^{−1}, and the hs-DNA concentrations were varied from 4.0×10^{-4} mol L^{−1} to 1.2×10^{-3} mol L^{−1} at room temperature.

2.6. Circular dichroism spectra

Far UV-CD spectra were recorded using a Hitachi-F7000 fluorescence spectrometer with a 1.0 cm path length quartz cell at



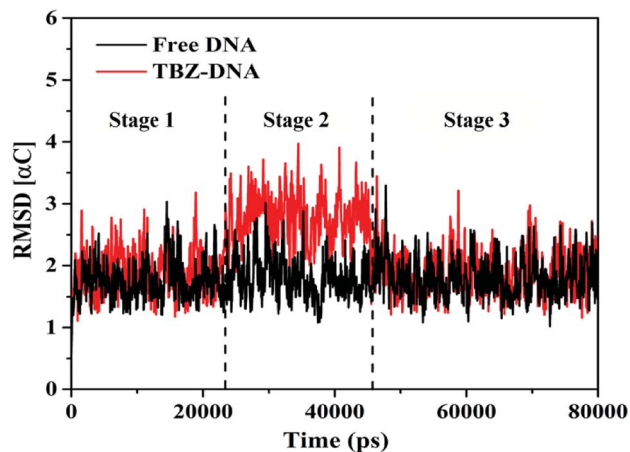


Fig. 3 Root mean square deviations of the complex systems in the MD simulations.

atoms on the pyrimidine ring of adenine and guanine. This phenomenon indicated that adenine and guanine were the main base pairs that participated in the interaction. Given the similar binding characteristics of the top five binding poses, the top one was selected for molecular dynamical simulation to evaluate the stability of the interaction and explore the steady binding conformation.

3.1.2 Analysis of molecular dynamics

3.1.2.1. Analysis of the systematic stabilities using RMSD. In this study, MD simulations were performed under simulated physiological conditions to further understand the stability and dynamics behavior of TBZ–DNA complex in different time-scales. Root mean square deviations, which is deviation statistics on behalf of the structural changes and the atomic fluctuation from the initial position, is a parameter for assessing the stability of simulation system.²⁸ The RMSD values of α -carbon atoms for the free DNA and TBZ–DNA complex systems *versus* simulation time were showed in Fig. 3. The RMSD values for free DNA fluctuated in a balanced position throughout the simulation and thus indicated the stable double helix structure of DNA. Compared with that in blank control system, the RMSD in TBZ–DNA systems displayed an apparent disturbance and eventually reached plateau at approximately 46 ns. Thus, TBZ could steadily bind with DNA and stayed at a stable binding position after a slight structural drift and low atomic fluctuations.

3.1.2.2. Dynamics behavior analysis of trajectory. According to the fluctuation of RMSD in 80 ns, the molecular dynamics process can be divided into three stages for trajectory analysis to evaluate the dynamics behavior of the binding (Fig. 3). The first stage (Stage 1) was from the initial state to 24 ns, the second stage (Stage 2) was from 24 ns to 46 ns, and the third stage (Stage 3) was from 46 ns to 80 ns. A representative trajectory for each stage was selected, specifically, 0 ns, 35 ns, and 80 ns represented Stage 1, Stage 2, and Stage 3, respectively. As shown in Fig. 4, the TBZ molecule moved from the center of the DNA to the edge area in the MD simulation. In the initial conformation (Stage 1, time = 0 ns), the TBZ molecule bound in the border

area of A-T and C-G base pairs of DNA, and formed two hydrogen bonds with adenine and guanine in the distance of 2.99 and 2.95 Å, respectively. In Stage 2 (time = 35 ns), the hydrogen bond between TBZ and DNA fractured, and the TBZ molecule drifted from the initial binding position and was maintained by hydrophobic interaction with DNA. In the final stable state (Stage 3, time = 80 ns), TBZ steadily bound in the terminal region enriched with A-T base pairs by forming hydrogen bonds with cytosine and guanine in the distances of 2.71 and 3.14 Å, respectively. In conclusion, TBZ bound to the groove position of DNA comprising A-T and C-G base pairs. Hydrogen bonds and hydrophobic interaction were the main forces for maintaining this combination.

The hydrogen bond numbers of TBZ–DNA system were analyzed throughout the MD simulation and were shown in Fig. 5. The numbers of hydrogen bond ranged from 0 to 2, and the variation of hydrogen bonds number could be divided into three stages similar to RMSD. In Stages 1 and 3, the numbers of hydrogen bond ranged from 1 to 2, whereas that in Stage 2 ranged from 0 to 1. The results indicated that hydrogen bonds played an important role in the binding and stabilities of TBZ and DNA.

3.2. Analysis of the binding mechanism of TBZ–DNA system

3.2.1. Theory of NMR relaxation study. The proton spin relaxation rate of the small molecules was measured using NMR spectroscopy. The selective spin–lattice relaxation rates (R^{se}) are sensitive to the affinity of ligand with and without receptors. The binding process of the ligand–receptor can be described as follows:



When the ligand–receptor reach equilibration, the thermodynamic equilibrium constant can be written as $K = [ML]/[M][L]$. Under conditions of fast chemical exchange between free and bound environments, the R^{se} can be expressed by the following equation:²⁴

$$\Delta R^{\text{se}} = \frac{KR_b^{\text{se}}}{(1 + K[L])} [M_0] \quad (2)$$

where $\Delta R^{\text{se}} = R_{\text{obs}}^{\text{se}} - R_f^{\text{se}}$, $R_{\text{obs}}^{\text{se}}$ is the experimentally determined selective relaxation rate, R_f^{se} and R_b^{se} are the selective spin–lattice relaxation rates in free and bound states, respectively. The affinity index needs be normalized to the free ligand relaxation rate due to the differences in the portions dynamics of the molecule caused by the different correlation times and modulation of the dipolar interactions between protons at different positions and spin densities on the ligand proton. The normalization of $\Delta R^{\text{se}} = R_{\text{obs}}^{\text{se}} - R_f^{\text{se}}$ to R_f^{se} cancels the effects of the above-mentioned factors on the interaction between ligand and macromolecule and leads to a normalized affinity index:

$$\Delta R_N^{\text{se}} = \frac{KR_b^{\text{se}}}{(1 + K[L])R_f^{\text{se}}} [M_0] \quad (3)$$



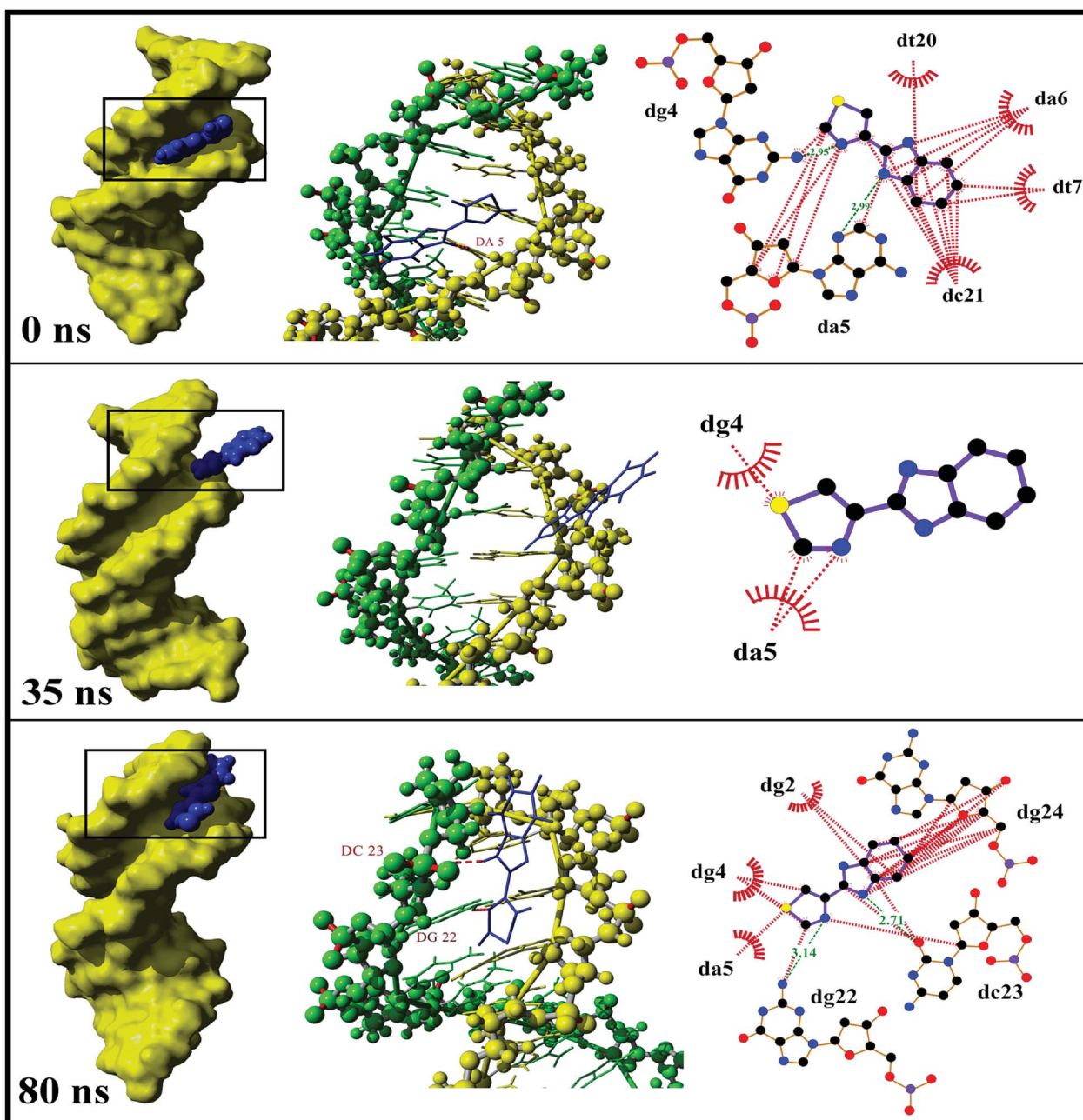


Fig. 4 Trajectory analysis of three stages at 0, 35, and 80 ns. Left images showed the binding region and binding mode of TBZ to DNA, where DNA was showed as molecular surface and TBZ was spherical model. Middle images were the interaction conformations of complex, in which the hydrogen bonds were red dotted line and the base formed hydrogen bond with TBZ was labeled. Right images showed the schematic representation of Ligplot software, wherein hydrophobic interaction was colored red and hydrogen bond colored green with distance.

The normalized relaxation rate ΔR_N^{se} versus the macromolecule concentration $[M_0]$ is characterized by a positive linear correlation. The slope of the straight line passing through the origin of the axes is $[A^N]_L^T$:

$$[A^N]_L^T = \frac{KR_b^{se}}{(1 + K[L])R_f^{se}} \quad (4)$$

which is defined as the “normalized affinity index” ($L \text{ mol}^{-1}$). This value remains constant at a specified temperature and

ligand concentration. Moreover, the normalized relaxation rate ΔR_N^{se} eqn (3) can be shown as:

$$\frac{1}{\Delta R_N^{se}} = \frac{R_f^{se}}{KR_b^{se}[M_0]} + \frac{[L]R_f^{se}}{R_b^{se}[M_0]} \quad (5)$$

Plotting $1/\Delta R_N^{se}$ to the ligand concentration $[L]$, the binding constant K can be calculated.



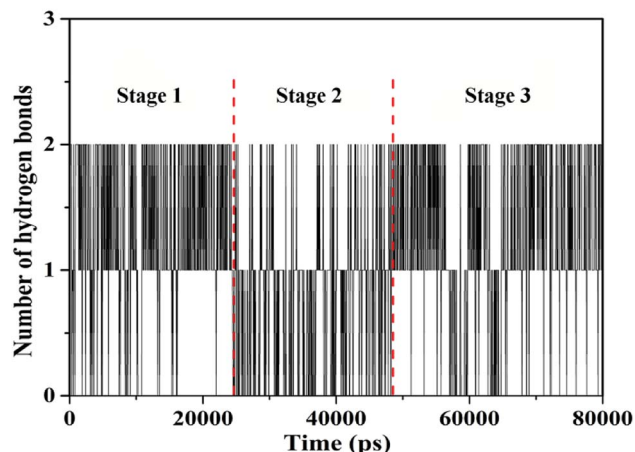


Fig. 5 The number of hydrogen bonds between TBZ and DNA throughout the MD simulations.

3.2.2. Analysis of the affinities and binding constant. The ^1H NMR spectrum of TBZ was shown in Fig. 6A. According to the results of computer simulations, three groups of hydrogen protons from the benzene ring and the thiazole ring (H1/H2, H14, and H12) were selected to compare the binding abilities. The normalized affinity indexes $[A_i^{\text{N}}]_{\text{L}}^{\text{T}}$ were fitted in Fig. 6B to remove the effects of different correlation times and proton densities. The curves showed that the normalized selective

relaxation rate of TBZ increased with the addition of DNA. For the H1/H2, H14, and H12, the values of $[A_i^{\text{N}}]_{\text{L}}^{\text{T}}$ for the TBZ–DNA system were 1396.23 ± 254.19 , $33\,573.03 \pm 4120.80$, and $8498.93 \pm 5.52 \text{ L mol}^{-1}$, respectively. The NMR results confirmed that the thiazole ring was the main group bound with DNA, and the affinity of benzene rings was weak due to the absence of hydrogen protons on the imidazole ring.

In order to ascertain the binding mechanism between TBZ and DNA, the binding constant was calculated according to eqn (5) in Fig. 6C. The binding constant of the TBZ–DNA complex was $462.43 \text{ L mol}^{-1}$ at 298 K, which was lower than that of the reported intercalate binding mode ranging from 10^5 L mol^{-1} to 10^6 L mol^{-1} .²⁹ The low binding affinity of the TBZ and DNA served as an evidence for the non-intercalative binding mode.

3.2.3. Investigation of fluorescence emission spectrum. The emission spectra of TBZ at different concentrations of DNA were shown in the inner picture of Fig. 7. The maximum fluorescence intensity of TBZ decreased with the increase of DNA, indicating that the fluorophore of TBZ was affected with the added DNA. The quenching constant (K_{sv}) and binding constant (K) were calculated by the linear Stern–Volmer equation^{30,31} at four different temperatures (298, 303, 310, and 317 K) and were presented in Table 1. It is well-known that the decrease of K_{sv} with the increased temperatures implies that the fluorescence quenching mechanism is static quenching, whereas a reverse trend is dynamic quenching.^{32,33} Thus, the interaction between TBZ and hsDNA was static quenching.

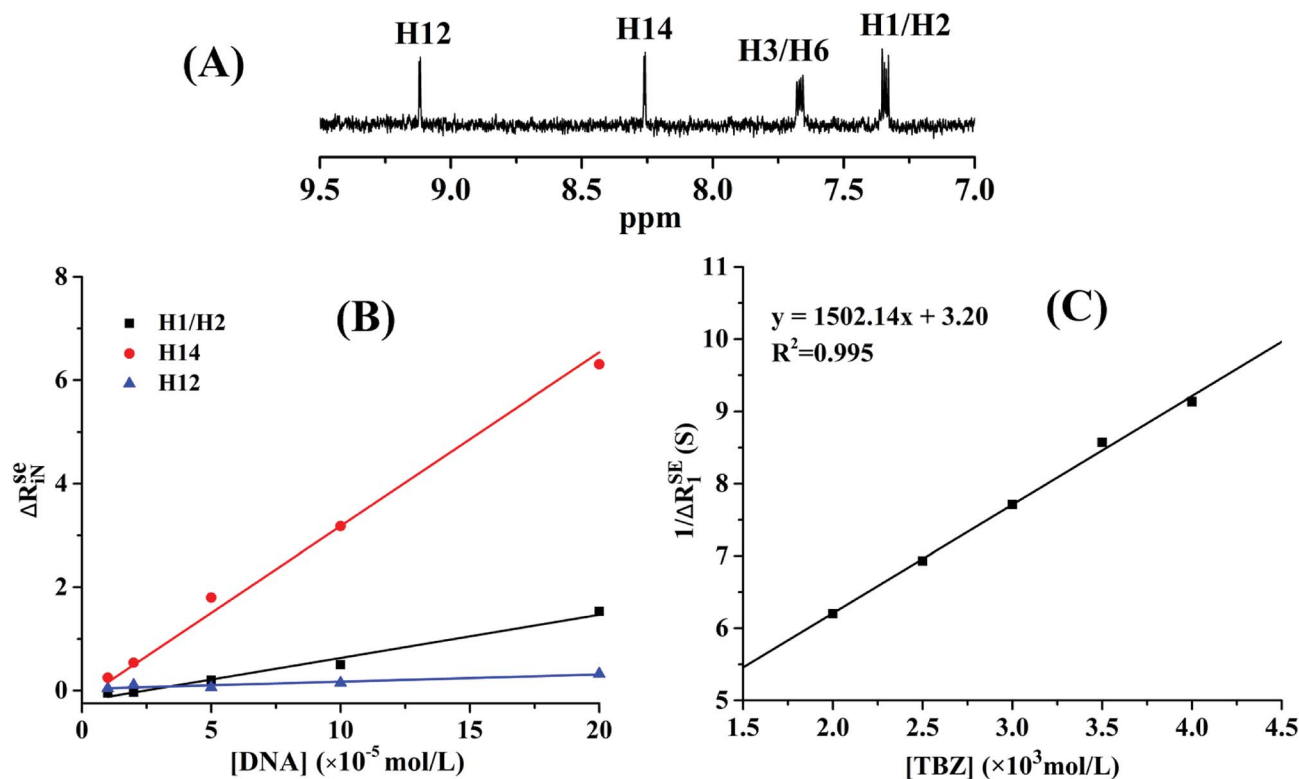


Fig. 6 (A) Proton NMR spectrum of thiabendazole in DMSO- d_6 solution. (B) Plots of normalized selective relaxation rate enhancements of H1/H2, H14, and H12. $[\text{TBZ}] = 2 \times 10^{-3} \text{ mol L}^{-1}$, $[\text{DNA}] = 1, 2, 5, 10, 20 \times 10^{-5} \text{ mol L}^{-1}$, $\text{pH} = 7.4$, $T = 298 \text{ K}$. (C) Equilibrium constant of the TBZ–DNA system. $[\text{DNA}] = 2 \times 10^{-5} \text{ mol L}^{-1}$, $[\text{TBZ}] = 2, 2.5, 3, 3.5, 4 \times 10^{-3} \text{ mol L}^{-1}$, $\text{pH} = 7.4$, $T = 298 \text{ K}$.



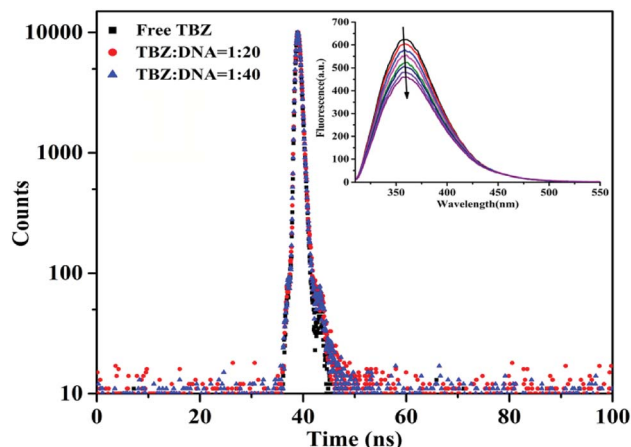


Fig. 7 The fluorescence decay curve of TBZ in the presence and absence of hs-DNA ($\lambda_{\text{ex}} = 310$ nm and $\lambda_{\text{em}} = 360$ nm). The inner figure is emission spectra of TBZ in different concentrations of DNA at 298 K.

As shown in Table 1, the value of K was $471.53 \text{ L mol}^{-1}$ at 298 K, which was similar with NMR results. The values of K decreased with the increase of temperature, possibly indicating that the decomposition of the TBZ–DNA complex with the temperature increased. Moreover, the binding forces (including hydrogen bonds, electrostatic attraction, van der Waals forces, and hydrophobic interaction) were analyzed by the

thermodynamic parameters using the Van't Hoff equation.^{14,34} Hydrogen bonds and van der Waals forces played the main role in the interaction because of the negative ΔH and ΔS . These values showed that the interaction was spontaneous and predominantly driven by enthalpy; these characteristics were consistent with the other non-intercalative binding.³⁵ A previous study²⁶ had shown that the binding was an entropy-driven process and maintained by hydrophobic force, the different thermodynamic behavior compared with this work might be due to the model selection of DNA from different sources.

3.2.4. Time-resolved fluorescence spectrum. Fluorescence lifetime measurement is an important method to distinguish between static and dynamic quenching of acceptor and ligand. In static quenching, the decay time of the free fluorophores does not change, whereas in dynamic quenching, the mean decay time of the entire population in the excited state decreases.³⁶ In Fig. 7, the average lifetime of TBZ remained almost constant with the addition of hsDNA. This finding led to the conclusion of static quenching mechanism between TBZ and hsDNA, and was in good agreement with the fluorescence quenching analysis.

3.3. Investigation of the binding mode

3.3.1. Circular dichroism (CD) spectroscopy. CD is a powerful tool to monitor the structure of DNA, which the characteristic peak of right-handed B form DNA was a positive peak at ~ 276 nm (base stacking) and a negative peak at

Table 1 Quenching and thermodynamic parameters of TBZ–DNA systems at four different temperatures

T (K)	K_{sv} (L mol^{-1})	R^a	K_a (L mol^{-1})	n	R^b	ΔG (kJ mol^{-1})	ΔH (kJ mol^{-1})	ΔS [$\text{J (mol}^{-1} \text{K}^{-1})$]	R^c
298	275.63	0.991	471.56	1.094	0.996	−15.220	−30.779	−52.385	0.975
303	265.23	0.975	385.75	1.063	0.999	−14.906			
310	258.06	0.985	267.44	1.004	0.994	−14.540			
317	249.94	0.994	221.65	0.966	0.995	−14.173			

^a The correlation coefficient for the K_{sv} values. ^b Is the standard deviation for the K values. ^c The correlation coefficient for the van't Hoff plot.

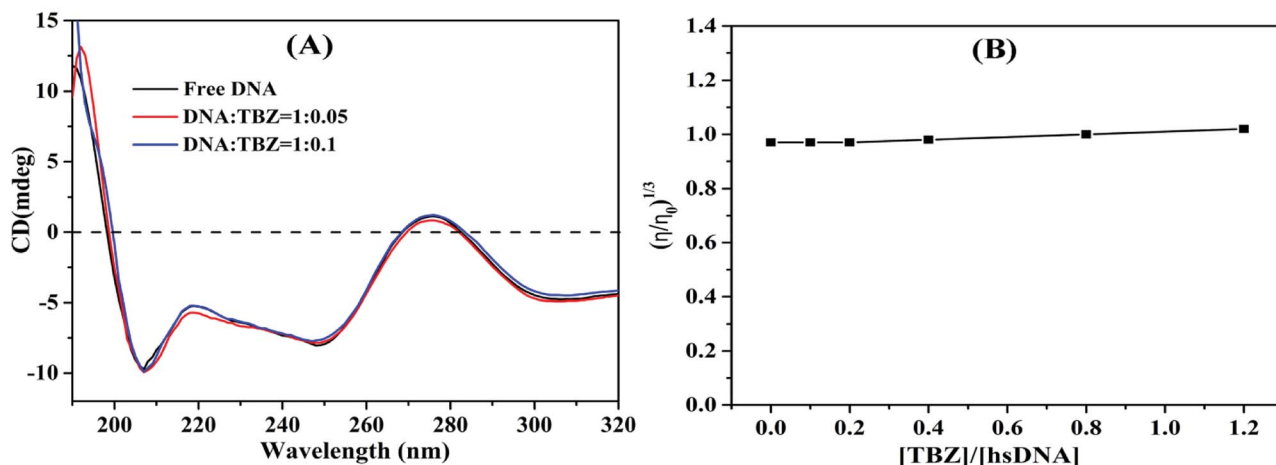


Fig. 8 (A) CD spectra of hsDNA with the increasing concentrations of TBZ. (B) Effect of increasing amounts of TBZ on the relative viscosity of hsDNA at 298 K.



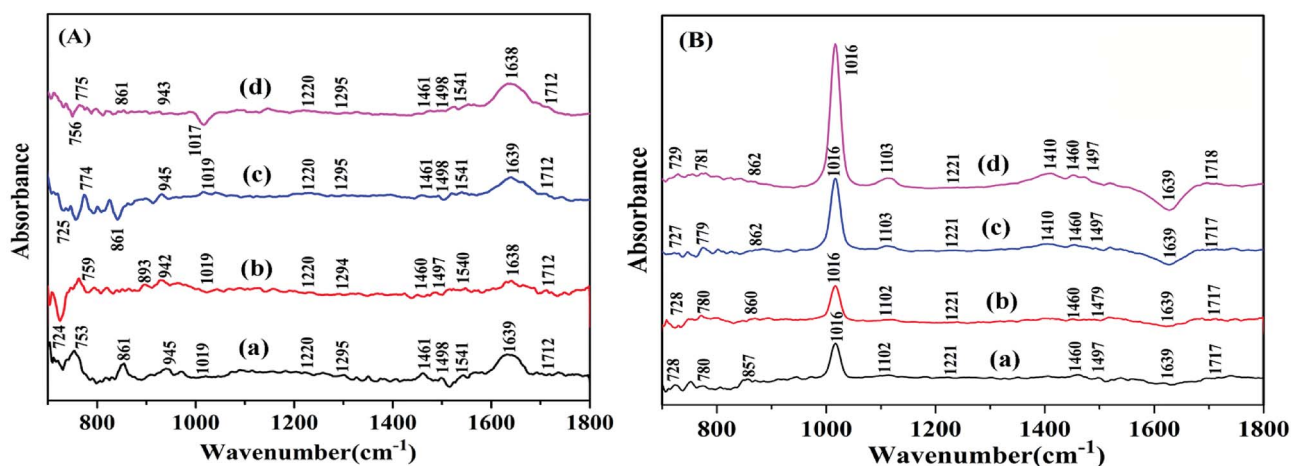


Fig. 9 (A) The difference spectra of free DNA (a) and [(DNA + TBZ) – TBZ]. (B) The difference spectra of free drug (a) and [(DNA + TBZ) – DNA]. The molar ratios ([TBZ] : [hsDNA]) for (b)–(d) were 1 : 20, 1 : 40 and 1 : 80, respectively.

~250 nm (right-handed helicity).³⁷ The changes in the CD spectra are related to the binding mode; that is, intercalation affects both the positive and negative peaks, and groove interaction almost does not change the CD spectra of DNA. In this work, the CD spectra of hsDNA with and without TBZ were shown in Fig. 8A. The most probable binding mode of TBZ with hsDNA was groove binding because the addition of TBZ had minimal effect on the DNA structure.

3.3.2. Viscosity measurements. Viscosity measurement is a useful and effectively method to distinguish the binding mode of ligand–DNA interaction.³⁸ In general, intercalative interaction separates the local base pairs to accommodate the ligand, thus resulting in DNA lengthening and increased viscosity; groove binding mode does not lead to remarkable changes in DNA viscosity.³⁹ Fig. 8B showed the slight increase in DNA viscosity changes with the addition of TBZ. The results indicated that TBZ bound to hsDNA though groove rather than through intercalation mode. Therefore, computer simulations and experiments demonstrated that TBZ bound to hsDNA with groove binding mode, which had no effect on the structure of hsDNA. The results were consistent with the study of Jalali *et al.*²⁶

3.4. Structural changes analysis of DNA

The infrared spectrum (FT-IR) is usually used to monitor the structure changes of biomacromolecules; the spectra of DNA are mainly confined from 1800 cm^{-1} to 700 cm^{-1} region to display protein secondary structure.^{40,41} The infrared spectral features of the free hsDNA and its complex form interacted with TBZ were shown in Fig. 9A. The band at 1712 cm^{-1} was related to guanine, which showed no change in wavenumber and intensity. The infrared band observed at 1639 cm^{-1} (thymine) showed a slight downshift. The infrared bands that appeared at 1541 (guanine and cytosine) and 1498 cm^{-1} (cytosine) all decreased in terms of intensity at all drug/DNA molar ratios. The deoxyribose sugar vibrations caused by C–C stretching were denoted by the infrared bands at 945 cm^{-1} in the spectrum of free hsDNA.⁴² The new negative band at 1017 cm^{-1} confirmed that TBZ reacted with hsDNA.

The difference spectra of [(DNA + TBZ) – DNA] (Fig. 9B) showed the remarkable negative band at 1639 cm^{-1} of thymine with decreasing intensity. The infrared band at 1102 cm^{-1} was primarily due to the phosphate symmetric vibrations; this band was assigned to the vibrations of the phosphodiester bonds (857 cm^{-1}).⁴³ The new band at 1410 cm^{-1} appeared when TBZ was added, which indicated that TBZ bound to hsDNA.

The structure of B-form is the closest with DNA in cell. FT-IR can be used to distinguish the DNA structure between B-form and A-form.⁴⁴ The bands at 1461 and 1220 cm^{-1} were stable, revealing that the DNA conformation tend to be in B-form. In Fig. 9A, the positive band at 893 cm^{-1} appeared, indicating the partial perturbation of DNA conformation. Moreover, the appearance and disappearance of the band at 861 cm^{-1} with the addition of TBZ confirmed that the A-form DNA was unstable. Thus, the interaction of the food additive TBZ with hsDNA did not result in transitions in the DNA double helix conformation. hsDNA was in the B-form with the perturbation of B-form transformed into A-form, which meant that the DNA helix became short and compact.

4. Conclusions

In this study, the binding mechanism and conformation stability of TBZ to hsDNA were characterized by computer simulations and spectroscopic methods. Computer simulations showed that TBZ could combine with the groove region of DNA through hydrogen bonds and hydrophobic interactions. The thiazole and imidazole rings of TBZ, particularly nitrogen atoms, were the main groups that maintain the stability of the binding with DNA. According to RMSD results, MD simulations were divided into three stages. This finding indicated that TBZ moved from the middle to the terminal region of DNA to form a stable conformation. The mechanism of TBZ binding to DNA was static quenching with weak binding. The binding of TBZ induced the slight conformation changes in DNA. Through comparison of the present results with previous studies, the selection of DNA was important for the study of ligand–receptor



interaction because of different binding mechanisms. This research provided an in-depth study of the binding mechanism, facilitated the application and development of computer simulations on the interactions of small molecules and biomacromolecules, and provided information on the selection of DNA model from different sources in interaction studies.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by Sichuan Science and Technology Program (Grant No. 2018JY0188), the National Natural Science Foundation of China (NNSFC Grant 21704069), and the Fundamental Research Funds for the Central Universities (Grant No. 2018SCU12043) the Postdoctoral Foundation of Sichuan University.

References

- M. Blazková, P. Rauch and L. Fukal, *Biosens. Bioelectron.*, 2010, **25**, 2122–2128.
- N. B. Andrioli and M. D. Mudry, *Discrete Appl. Math.*, 2011, **22**, 17–23.
- N. B. Andrioli, S. Soloneski, M. L. Larramendy and M. D. Mudry, *Mutat. Res., Genet. Toxicol. Environ. Mutagen.*, 2014, **772**, 1–5.
- Y. Feng, M. Lv, Y. Lu, K. Liu, L. Liu, Z. He, K. Wu, X. Wang, B. Zhang and X. Wu, *Food Chem.*, 2017, **243**, 118–124.
- V. Ferraro, A. R. Madureira, B. Sarmiento, A. Gomes and M. E. Pintado, *Food Res. Int.*, 2015, **77**, 450–459.
- X. Wang, Y. Liu, L. L. He, B. Liu, S. Y. Zhang, X. Ye, J. J. Jing, J. F. Zhang, M. Gao and X. Wang, *Food Chem. Toxicol.*, 2015, **78**, 42–51.
- A. Opar, *Nat. Rev. Drug Discovery*, 2009, **8**, 437–438.
- Q. Zhang, Y. Huang, L. Guo, C. Chen, D. Guo, Y. Chen and Y. Fu, *New J. Chem.*, 2014, **38**, 4600–4606.
- N. Shahabadi and M. Maghsudi, *Mol. BioSyst.*, 2013, **10**, 338–347.
- A. Boroujeni, Z. Khorasani-Motlagh and M. Noroozifar, *J. Biomol. Struct. Dyn.*, 2016, **34**, 414–426.
- Z. Aramesh Boroujeni, M. Khorasani-Motlagh and M. Noroozifar, *J. Biomol. Struct. Dyn.*, 2016, **34**, 414–426.
- X. Zhou, C. Zhang, G. Zhang and Y. Liao, *RSC Adv.*, 2016, **6**, 5408–5418.
- S. Yilmaz, F. Unal, D. Yüzbaşıoğlu and M. Celik, *Toxicol. Ind. Health*, 2014, **30**, 926–937.
- S. Sharma, M. Yadav, S. P. Gupta, K. Pandav and S. Kumar, *Chem.-Biol. Interact.*, 2016, **260**, 256–262.
- C. G. Reinhardt and T. R. Krugh, *Biochemistry*, 1978, **17**, 4845–4854.
- X. Li and S. Wang, *New J. Chem.*, 2014, **39**, 386–395.
- C. Ràfols, S. Zarza and E. Bosch, *Talanta*, 2014, **130**, 241–250.
- M. Šimšíková, *Arch. Biochem. Biophys.*, 2016, **593**, 69–79.
- K. J. Czogalla, A. Biswas, K. Höning, V. Hornung, K. Liphardt, M. Watzka and J. Oldenburg, *Nat. Struct. Mol. Biol.*, 2016, **24**, 77–85.
- A. Ahmedbelkacem, L. Colliandre, N. Ahnou, Q. Nevers, M. Gelin, Y. Bessin, R. Brillet, O. Cala, D. Douguet and W. Bourguet, *Nat. Commun.*, 2016, **7**, 12777.
- M. P. Williamson, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2013, **73**, 1–16.
- A. Gossert and W. Jahnke, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2016, **97**, 82–125.
- R. Amiri, A. K. Bordbar, M. F. Garcã-A-Mayoral, A. R. Khosropour, I. Mohammadpoor-Baltork, M. MenãNdez and D. V. Laurents, *J. Colloid Interface Sci.*, 2012, **369**, 245–255.
- D. Wu, Y. Zhai, J. Yan, K. Xu, Q. Wang, Y. Li and H. Li, *RSC Adv.*, 2015, **5**, 11036–11042.
- Y. Wang, T. Zhang, J. Xu and W. Du, *Int. J. Biol. Macromol.*, 2011, **48**, 81–86.
- F. Jalali and P. S. Dorraji, *Arabian J. Chem.*, 2013, **77**, S3947–S3954.
- M. Xu, Z. R. Ma, L. Huang, F. J. Chen and Z. Z. Zeng, *Spectrochim. Acta, Part A*, 2011, **78**, 503–511.
- S. Fujiwara and T. Amisaki, *Proteins: Struct., Funct., Bioinf.*, 2006, **64**, 730–739.
- Y. Cao and X. W. He, *Spectrochim. Acta, Part A*, 1998, **54**, 883–892.
- Q. Sun, H. Yang, P. Tang, J. Liu, W. Wang and H. Li, *Food Chem.*, 2017, **243**, 74–81.
- A. Sharma, S. Anandakumar and M. Ilanchelian, *RSC Adv.*, 2014, **4**, 36267–36281.
- L. Fotouhi and M. Zabeti, *Monatsh. Chem.*, 2016, **147**, 837–844.
- B. Barare, M. Yildiz, H. Ünver and K. Aslan, *Tetrahedron Lett.*, 2016, **57**, 537–542.
- Q. Wang, Q. Sun, P. Tang, B. Tang, J. He, X. Ma and H. Li, *RSC Adv.*, 2015, **5**, 81696–81706.
- M. M. V. Ramana, R. Betkar, A. Nimkar, P. Ranade, B. Mundhe and S. Pardeshi, *Spectrochim. Acta, Part A*, 2016, **152**, 165–171.
- H. Yang, P. Tang, B. Tang, Y. Huang, X. Xiong and H. Li, *RSC Adv.*, 2017, **7**, 10242–10251.
- D. Ajloo, M. E. Moghadam, K. Ghadimi, M. Ghadamgahi, A. A. Saboury, A. Divsalar, M. Sheikhmohammadi and K. Yousefi, *Inorg. Chim. Acta*, 2015, **430**, 144–160.
- N. Shahabadi and S. Amiri, *Spectrochim. Acta, Part A*, 2015, **138**, 840–845.
- S. Zhang, X. Sun, R. Kong and M. Xu, *Spectrochim. Acta, Part A*, 2015, **136**, 1666–1670.
- A. A. Ouameur and H. A. Tajmir-Riahi, *J. Biol. Chem.*, 2004, **279**, 42041–42054.
- F. A. Joozdani, F. Yari, P. A. Joozdani and S. Nafisi, *PLoS One*, 2015, **10**, e0127541.
- D. K. Jangir, G. Tyagi, R. Mehrotra and S. Kundu, *J. Mol. Struct.*, 2010, **969**, 126–129.
- G. Tyagi, S. Charak and R. Mehrotra, *J. Photochem. Photobiol., B*, 2012, **108**, 48–52.
- L. Han, Y. Zhou, X. Huang, M. Xiao, L. Zhou, J. Zhou, A. Wang and J. Shen, *Spectrochim. Acta, Part A*, 2014, **123**, 497–502.

