Synthesis, DNA-binding, Photocleavage and in vitro Cytotoxicity of Novel Imidazole[4,5-f][1,10]phenanthroline-based Oxovanadium Complexes

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Abstract

Three novel imidazole[4,5-f][1,10]phenanthroline-based oxovanadium complexes [VO(hntdtsc)(HPIP)] (1) (hntdtsc = 2-hydroxy-1-naphthaldehyde thiosemicarbazone, HPIP = 2-(2-hydroxyphenyl)imidazole[4,5-f][1,10]phenanthroline), [VO(hntdtsc)(m-HPIP)] (2) (m-HPIP = 2-(3-hydroxyphenyl)imidazole[4,5-f][1,10]phenanthroline), [VO(hntdtsc)(p-HPIP)] (3) (p-HPIP = 2-(4-hydroxyphenyl)imidazole[4,5-f][1,10]phenanthroline), have been synthesized and characterized by elemental analyses and spectroscopic techniques. Their DNA-binding properties with calf-thymus DNA (CT-DNA) were studied by various methods. The cytotoxicity of these three complexes was evaluated by 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay. The electronic spectral results reveal that three complexes can bind to CT-DNA by intercalation mode. The electrophoresis studies also show that they can efficiently cleave pBR322 DNA. The in vitro antiproliferative activity of complex 1 against human CaSki, SiHa, K562, HepG2, EC9706 and EC109 cell lines is proved to be more effective than both 2 and 3.

Keywords: Oxovanadium complexes; DNA-binding; Photocleavage; Cytotoxicity

Introduction

In recent years, more and more biochemists and pharmacologists have paid their attention to research on other transition metal complexes such as ruthenium (II), copper (II), zinc (II), nickel (II), cobalt (II) complexes compared with traditional platinum compounds in treatment of human cancer due to their potential applications as chemical and stereoselective probes of nucleic acid structures as well as diverse biological activities [1-5]. As a trace bioelement existing in the human body, vanadium has been found to present a variety of physiological activities such as nutritional metabolism [6], insulin mimetic effect [7], as well as closely connected with cell apoptosis and cancer [8]. In the past decades, oxovanadium compounds have been extensively studied for their potential biological and pharmacological activities such as antibacterial [9], biocatalytic oxidation [10], insulin-enhancing effects [11], apoptosis-inducing activity [12], potential capabilities as DNA structural probes and DNA dependent electron transfer likewise [13].

In addition, thiosemicarbazones were often chosen as primary ligands for vanadium owing to their wide range of physiological and pharmacological activities including antifungal [14], antitumor [15], antibacterial [16] and antiviral activities [17]. Oxovanadium complexes incorporating with thiosemicarbazones have demonstrated both in vitro antibacterial and antiproliferative properties [18-19] as well as intense interaction with DNA [20]. On the other hand, 1,10-phenanthroline and its derivatives were usually served as a very efficient class of ligands, which lead to efficient catalysis in cross coupling reaction of aliphatic alcohols and aroyl halides [21], antimicrobial activities [22], efficient DNA-binding and DNA cleavage [23] thus inducing significant cytotoxicity in the cultured cells [24]. Subsequently, research on mixed-ligand metal complexes has been put into practice in order to combine the pharmacological properties of both the ligands and metal thus reduce toxicity simultaneously [25]. So far, a certain attentions have been paid to Ru (II), Cu (II), Co (II) and Ni (II) complexes incorporating imidazole[4,5-f][1,10]phenanthroline derivatives ligands [26-27]. However, there have been relatively few studies on oxovanadium complexes with imidazole derivatives of 1,10-phenanthroline and their mechanism.

Previously, we have reported some oxovanadium (IV) complexes incorporating 1,10-phenanthroline (phen) or bipyridyl (bpy) present high DNA intercalator, photocleavage properties and cytotoxicity in vitro [13,19-20]. Although the molecular basis for the DNA-binding properties of oxovanadium (IV) compound is not completely understood at the present time, it occurs to us that oxovanadium complexes bearing phen moiety possess higher DNA-binding activities and more potent inhibitory effect against cancer cell lines than that of bearing bpy moiety. The previous results provide strong evidence for us to assume that DNA-binding activity of oxovanadium complexes may increase owing to the introduction of larger aromatic ring in auxiliary ligands substituted with 1,10-phenanthroline.

In the present work, we have synthesized three unsymmetrical imidazole[4,5-f][1,10]phenanthroline-based oxovanadium complexes [VO(hntdtsc)(HPIP)] (1) (hntdtsc = 2-hydroxy-1-naphthaldehyde thiosemicarbazone, HPIP = 2-(2-hydroxyphenyl)imidazole[4,5-f][1,10]phenanthroline), [VO(hntdtsc)(m-HPIP)] (2) (m-HPIP = 2-(3-hydroxyphenyl)imidazole[4,5-f][1,10]phenanthroline), [VO(hntdtsc)(p-HPIP)] (3) (p-HPIP = 2-(4-hydroxyphenyl)imidazole[4,5-f][1,10]phenanthroline), and characterized by elemental analysis, IR, molar conductance, ES-MS and 1H NMR. The DNA-binding properties of these three complexes were well studied by UV-Vis titration, fluorescence spectra, viscosity measurements and thermal denaturation studies. Photocleavage reactions with pBR322 supercoiled plasmid DNA were investigated by agarose gel electrophoresis experiments. Their cytotoxicity in vitro against cervical cancer CaSki and SiHa, leukemia K562, HepG2, esophageus carcinoma EC9706 and EC109 cell lines were investigated by 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay. The electronic spectral results reveal that three complexes can bind to CT-DNA by intercalation mode. The electrophoresis studies also show that they can efficiently cleave pBR322 DNA. The in vitro antiproliferative activity of complex 1 against human CaSki, SiHa, K562, HepG2, EC9706 and EC109 cell lines is proved to be more effective than both 2 and 3.
assessed by MTT assay. The compounds employed in this work are shown in Scheme 1.

Experimental

Materials and methods

All starting chemicals used in the synthesis and physical measurements were commercially available reagents and without further purification unless otherwise specified. 1,10-phenanthroline (phen) was obtained from Guangzhou Chemical Reagent Factory. VO(acac)2 were purchased from Alfa Aesar, thiosemicarbazide (TSC) was purchased from TCI, CT-DNA and pBR322 supercoiled plasmid DNA were obtained from Sigma. Cell lines of CaSki, SiHa, K562, HepG2, EC9706 and EC109 were purchased from American Type Culture Collection. Tris buffer A (Tris=tris(hydroxyl-methyl) aminomethane) containing 5 mM Tris-HCl and 50 mM NaCl (pH=7.2) was used for absorption titration and viscosity measurements. Tris buffer B containing 50 mM Tris-HCl and 18 mM NaCl (pH=7.2) was used for gel electrophoresis experiments. Buffer C containing 1.5 mM Na2HPO4, 0.5 mM NaH2PO4 and 0.25 mM Na2H2EDTA (H4EDTA=N,N′-ethylene-1,2-diylbis[N-(carboxymethyl) glycine]) (pH=7.0) was used for thermal denaturation. A solution of CT-DNA in buffer A gave a ratio of UV absorbance at 260 and 280 nm of 1.8-1.9:1, indicating that the DNA was sufficiently free of protein [13,19]. The DNA concentration per thermal denaturation. A solution of CT-DNA in buffer A gave a ratio of UV absorbance at 260 and 280 nm of 1.8-1.9:1, indicating that the DNA was sufficiently free of protein [13,19]. The DNA concentration per

Physical measurements

Microanalysis (C, H, and N) was carried out with a PerkinElmer 240Q elemental analyzer. Electrospray mass spectra (ES-MS) were recorded on an LCQ system (Finnigan MAT, USA) using methanol as mobile phase. 1H NMR spectra were recorded on a Varian-500 spectrometer. All chemical shifts are given relative to tetramethylsilane (TMS). Infrared spectra were recorded on a Bomem FTIR model MB102 instrument using KBr pellets method. UV–Vis spectra were recorded on a Shimadzu UV-3101 PC spectrophotometer at room temperature. Emission spectra were recorded on a Perkin-Elmer Lambda 55 spectrofluorophotometer. Molar conductivities in DMF (10-3M) solution at room temperature were measured using a DDS-307 digital direct reading conductivity meter.

Synthesis of phenanthroline-based ligands: HPIP, m-HPIP and p-HPIP

HPIP was synthesized through modification of a previously reported method [29]. A mixture of salicylaldehyde (0.69 mL, 5 mmol) and ammonium acetate (7.70 g, 0.1 mol) was added into a stirring solution of 1, 10-phenanthroline-5, 6-dione (0.99 g, 5 mmol) in 60 mL of glacial acetic acid, and the mixture was continuously stirred at 60°C for 6 hours. Then the deep red solution was cooled to room temperature and diluted with 100 mL distilled water. A grayish precipitate was obtained by neutralization with ammonium hydroxide. Then the mixture was filtered and washed with water for three times. The crude solid power was purified by chromatography over 60-80 mesh SiO2 using absolute ethanol as eluent. The solvent was removed and the products were collected, and dried in vacuo. m-HPIP and p-HPIP were prepared by a similar procedure as for the compound HPIP, with 3-Hydroxybenzaldehyde (0.69 mL, 5 mmol) and 4-Hydroxybenzaldehyde (0.69 mL, 5 mmol) in place of salicylaldehyde (0.69 mL, 5 mmol) respectively, and a grey-white precipitate were obtained.

HPIP: Yield: 86%. Anal. Found: C, 73.12; H, 3.79; N, 18.03; Calcd for C19H12N4O: C, 73.07; H, 3.87; N, 17.94. 1H NMR (DMSO-d 6, 500 MHz) δ: 12.85 (s, 1H, -NH), 10.07 (s, 1H, -OH), 9.91 (d, 2H, J = 7.4Hz, ArH), 9.85 (d, 2H, J = 7.7Hz, ArH), 8.95 (d, 2H, J = 9.9Hz, ArH), 8.20 (d, 2H, J = 8.2Hz, ArH), 2.90 (s, 4H, 1-CH2).
Synthesis of [VO(hntdtsc)(m-HPIP)] (2)

This complex was obtained by a similar procedure as for the complex 1, with p-HPIP (0.312 g, 1 mmol) in place of HPIP. Yield: 72%. Anal. Found: C, 73.12; H, 3.49; N, 17.83; Calcd for C21H14N4O3SV: C, 72.98; H, 3.49; N, 17.83. 1H-NMR (DMSO-d 6, 500 MHz) δ: 12.35 (s, 1H, -NH), 11.82 (s, 1H, CH=N), 10.30 (s, 1H, -OH), 11.73 (d, 2H, J = 7.6 Hz, ArH). 8.84 (d, 2H, J = 8.0 Hz, ArH), 8.49 (d, 1H, J = 8.2 Hz, ArH), 7.58 (m, 4H, J = 8.0 Hz, ArH), 7.39 (s, 1H, ArH), 6.95-6.94 (m, 4H, J = 8.7 Hz, ArH). Your equation

DNA-binding and photocleavage

The absorption titration of oxovanadium complexes in buffer A were performed at room temperature with a fixed concentration of the oxovanadium complex (20 mM) to which increments of DNA stock solutions were added. The oxovanadium-DNA solutions were incubated at room temperature for 5 min before the absorption spectra were recorded. In order to further elucidate the binding strength of the complex, the intrinsic binding constant Ks with CT-DNA was obtained by monitoring the change in the absorbance of the ligand transfer band with increasing amounts of DNA. Ks was then calculated using the following equation [13,20,28]

\[
\frac{[DNA]}{\varepsilon - \varepsilon_f} = \frac{1}{K_s} + \frac{[DNA]}{\varepsilon_f - \varepsilon}
\]

Where [DNA] is the concentration of DNA in the base pairs, and ε, ε_f and ε refer to the corresponding apparent absorption coefficient A_{app}[Vanadium], the extinction coefficient for the free oxovanadium complex and the extinction coefficient for the oxovanadium complex in the fully bound form, respectively. In plots of [DNA]/(ε - ε_f) versus [DNA], K_s is obtained by the ratio of the slope to the intercept.

Viscosity measurements were carried out with an Ubbelohde viscometer maintained at a constant temperature of (28 ± 0.1) °C in a thermostatic bath. Flow time was measured with a digital stopwatch, and each sample was measured five times to obtain the average flow time. Date are presented as (η/η_0)¹/³ versus binding ratio [30], where η is the viscosity of DNA in the presence of complex while η_0 is the viscosity of DNA alone.

Thermal denaturation studies were carried out with Shimadzu UV-3101 PC spectrophotometer equipped with a Peltier temperature-controlling programmer (± 0.1°C). The melting temperature (T_m) was taken as the mid-point of the hyperchromic transition. The melting curves were obtained by measuring the absorbance at 260 nm for solutions of CT-DNA (80 µM) in the absence and presence of oxovanadium complex [20 µM] as a function of the temperature was scanned from 50 to 90 °C at a speed of 5°C min⁻¹. The data are presented as A/A_0 (A/A_0) versus binding ratio [30], where A is the viscosity of DNA in the presence of complex while A_f is the viscosity of DNA alone.

The cleavage activity of supercoiled pBR322 DNA by the oxovanadium complexes was studied by using agarose gel electrophoresis experiment, pBR322 DNA (0.1 µg) was treated with the complex.
**Cell viability assay**

3-((4,5-Dimethylthiazole)-2,5-diphenyltetrazolium bromide (MTT) dye assay was carried out to evaluate cytotoxicity in human CaSki, SiHa, HeLa, HEp2, HEp3, EC9706 and EC109 cells. Cells were seeded in 96-well microassay culture plates (2 x 10^4 cells per well) and incubated at 37 °C in a 5% CO_2 atmosphere for 48 h. The compounds tested were then added to the wells to achieve final concentration ranging 10^6 to 10^4 mol L^{−1}. Control wells were prepared by addition of culture medium (100 μL). Wells containing culture medium without cells were used as blanks. Upon completion of the incubation, the stock MTT dye solution (20 μL, 5 mg mL^{−1}) was added to each well. After 4 h incubation, a solution containing N, N-dimethylformamide (50%) and sodium dodecyl sulfate (20%) was added to solubilize the MTT formazan. The cell viability of each well was then measured on a Multiskan SSC ent microplate reader at 550 nm. The IC_{50} values for a square-pyramidal geometry [19,28,37] and it also suggested that the DNA-binding affinities of complexes 1, 2 and 3 can be observed in the ultraviolet titration process. The appreciable hypochromism and bathochromism for complex 1 is 56.17% and 3 nm at 272 nm. In contrast, a smaller hypochromism can be observed for complexes 2 and 3, which exhibit 12.53% and 25.23%, respectively, and also bathochromism of 1 nm at 268 nm and 1.5 nm at 270 nm, respectively. According to previously reported results [13,19,24], the UV–Vis spectral characteristics suggest that complexes 1-3 interact with CT-DNA most likely through a mode that involves a stacking interaction between the planar aromatic chromophore of the complexes and the base pairs of DNA usually result in hypochromism and bathochromism [13,20,28]. As can be seen in Figure 1, with the CT-DNA concentration increasing, obvious hypochromism as well as a certain degree of bathochromism for the oxovanadium complex 1-3 can be observed in the ultraviolet titration process. The appreciable hypochromism and bathochromism for complex 1 is 56.17% and 3 nm at 272 nm. In contrast, a slighter hypochromism can be observed for complexes 2 and 3, which exhibit 12.53% and 25.23%, respectively, and also bathochromism of 1 nm at 268 nm and 1.5 nm at 270 nm, respectively. According to previously reported results [13,19,24], the UV–Vis spectral characteristics suggest that complexes 1-3 interact with CT-DNA most likely through a mode that involves a stacking interaction between the aromatic chromophore and the base pairs of DNA.

**Results and Discussion**

**Synthesis of ligands and V–complexes**

The phenanthroline-based derivatives ligands were prepared by the reaction of 1,10-phenanthroline-5,6-dione with hydroxybenzaldehyde and VO(acac)₂. The desired complexes were synthesized by refluxing of corresponding ligand, VO(acac)₂ and phenanthroline-based ligand in absolute methanol. The desired complexes were purified by recrystallization.

In the IR spectra of phenanthroline-based ligands show absorption at ca. 3340 cm⁻¹ (v_{OH}), 3200 cm⁻¹ (imidazole N-H), 3040 cm⁻¹ (v_{CO}), 1448-1618 cm⁻¹ (v_{C=C}), 740 cm⁻¹, 700 cm⁻¹ (δ_{C-H}), respectively. In parallel, the electronic absorption spectra of complexes 1-3 observed at ca. 3440 cm⁻¹ (v_{OH}) and 3340 cm⁻¹ and 3340 cm⁻¹ was assigned to (NH₂), 3030 cm⁻¹ (v_{CO}), 1490-1580 cm⁻¹ (v_{C=C}), 1612-1616 cm⁻¹ (v_{C=O}), 1312 cm⁻¹, 810-313 cm⁻¹ (v_{C-O}, 731-737 cm⁻¹ and v_{OH}), which indicates both two mixed ligands are coordinated to vanadium. The strong (ν V=O) band at 956-974 cm⁻¹ is observed at 956-974 cm⁻¹ observed at oxovanadium complexes, which is not present in the spectrum of free ligands could be clearly identified for the formation of the complex [33-34].

Electronic spectra of complexes 1-3 show an intense band at ca. 268 nm assignable to π−π* transition of aromatic rings of phenanthroline-based ligands [35-36]. The remaining bands in the UV–vis region (320-330 nm) are attributed to intraligand transitions of the Schiff base. The absorption spectra of complexes 1-3, which exhibit 12.53% and 25.23%, respectively, and also bathochromism of 1 nm at 268 nm and 1.5 nm at 270 nm, respectively. According to previously reported results [13,19,24], the UV–Vis spectral characteristics suggest that complexes 1-3 interact with CT-DNA most likely through a mode that involves a stacking interaction between the aromatic chromophore and the base pairs of DNA.

In order to compare quantitatively the DNA-binding strengths of these complexes, the intrinsic binding constant K_{b} was calculated from the changes in absorbance in the ligand change transfer bands with increasing amounts of CT-DNA. The value of K_{b} was calculated as well using the equation, ranging (4.38 ± 0.01) x 10^{5} M^{-1} (1.07 ± 0.01) x 10^{5} M^{-1} and (1.83 ± 0.01) x 10^{5} M^{-1} for complexes 1, 2 and 3, respectively. The K_{b} values indicate that oxovanadium complexes 1-3 interact with CT-DNA by intercalation modes. It is interesting to note that the DNA-binding affinities of complexes 1, 2 and 3 are stronger than that of the reported oxovanadium complexes, such as VO(SAA) (phen) (4.50 x 10^{5} M^{-1}), VO(MOSAA)(phen) (2.95 x 10^{5} M^{-1}) [19] and VO(hntdtsc/phen) (8.2 x 10^{5} M^{-1}) [28], under our experimental conditions. This may due to the introduction of imidazole ring in auxiliary ligands substituted with 1, 10-phenanthroline, which leads to formation of a richer conjugated aromatic structure, thus resulting in stronger insertion into DNA. Meanwhile, the presence of an –OH group on the aromatic moieties, which acts as an electron-donating

| Compounds               | Ω cm⁻¹ mol⁻¹ |
|-------------------------|--------------|
| HPiP                    | 8.6          |
| m-HPiP                  | 9.8          |
| p-HPiP                  | 11.4         |
| VO(hntdtsc)(HPiP)       | 13.6         |
| VO(hntdtsc)(m-HPiP)     | 12.9         |
| VO(hntdtsc)(p-HPiP)     | 18.1         |

**Table 1:** Molar conductivity data of oxovanadium complexes and ligands.
The intrinsic binding constant, $K_b$, increase in the order 2<3<1. Interestingly, from a comparison of the DNA-binding activities of complexes 1, 2 and 3, the complex 1 appeared to be a much stronger DNA intercalator. The differences of their binding strength may due to the changes in the electronic characteristics of substituted group at the different locations introduced on the aromatic ring of phenanthroline-based ligands, which may make differences in the DNA-binding affinities. The results also indicates an –OH group substituted on the ortho-position of the aromatic ring exhibits stronger DNA-binding affinity than substituted on the para-position and meta-position [19,28]. And it thus provides strong evidence for the electronic effect of phenanthroline is one of the factors in determining the binding affinities.

**Fluorescence spectroscopic studies**

The interaction of the complexes (20 μmol L⁻¹) with CT-DNA was investigated using fluorescence emission titration experiment in the Tris buffer A at room temperature. The emission spectra of complexes 1, 2 and 3 in the absence and presence of CT-DNA exhibit luminescence in Tris buffer A, with maxima appearing at 296 nm, 656 nm and 697 nm respectively, which are shown in Figure 2. Emission intensity of complexes 1-3 is found to depend on DNA concentration.

Upon increasing concentrations of CT-DNA, the emission intensities of complexes 1-3 grow to about by around 1.21, 0.38, and 0.64 times larger than those in the absence of CT-DNA and saturates at a [DNA]/[V] ratio of 20 : 1. The enhancement of emission intensity is an indication of binding of the complexes to the hydrophobic pocket of DNA, since the hydrophobic environment inside the DNA helix reduces the accessibility of water molecules to the complex and the complex mobility is restricted at the binding site, leading to decrease of the vibrational modes of relaxation [38]. The fluorescence spectroscopic data shows that 1 interacts with CT-DNA more strongly than both 1 and 2, consistent with the UV-Vis absorption titration spectral results.

**Viscosity measurements**

Viscosity measurements are regarded as the least controversial and the most rigorous means of testing the binding mode of DNA in solution [19,28]. To further clarify the DNA-binding mode of complexes 1-3, viscosity measurements on solutions of CT-DNA incubated with the complexes were performed. It is well-known that a classical intercalation model leads to an apparent increase in viscosity of DNA solution due to base-pairs are pushed apart and hence an increase in overall DNA length. In contrast, a partial, non-classical intercalation of compounds could bend (or kink) the DNA helix and reduce its effective length and, concomitantly, its viscosity [19,20,24].

The effects of complexes 1-3 on the viscosity of CT-DNA are shown...
in Figure 3. As can be seen in Figure 3, with amounts of the complexes increasing, the relative viscosity of DNA increases continuously to some extent. The results thus provide strong evidence for the interaction of complexes 1-3 with CT-DNA by intercalation modes. Moreover, the large increase in the relative viscosity revealed that 1 is a better intercalator than 2 and 3, which is consistent with our foregoing hypothesis that electronic effects of introduction of larger planar aromatic rings on phenanthroline-based ligands play a key role in DNA-binding affinities.

Thermal denaturation studies

DNA melting experiments are generally applied to investigate the extent of intercalation, which were carried out by monitoring the intensity of DNA bases at 260 nm at different temperatures, both in the absence and presence of oxovanadium complex and ligands. Thermal behaviors of DNA in the presence of compounds can give insight into their conformational changes when the temperature is raised and offer information about the interaction strength of complexes with DNA. With the temperature in the solution rising, the double-stranded DNA will gradually dissociate to single strands and generate a hyperchromic effect on absorption spectra of DNA bases. Thus, the melting temperature $T_m$, which is defined as the temperature where half of the total base pairs are unbounded, is usually introduced. Generally, $T_m$ will increase considerable when intercalative binding occurs, since intercalation of the complex into DNA base pairs causes stabilization of base stacking and hence raises the melting temperature of the double-stranded DNA [13,24,28].

The melting curves of CT-DNA in the absence and presence of complexes 1, 2 and 3 are shown in Figure 4. As can be seen from Figure 4, the $T_m$ of CT-DNA in the absence of the complex is $61.6 \pm 0.2 ^\circ C$, meanwhile, when the complexes at a concentration ratio [V]/[DNA] of 1:4, we can observe that $T_m$ values in the presence of the complexes 1-3 are $71.4 \pm 0.2 ^\circ C$, $64.7 \pm 0.2 ^\circ C$ and $67.8 \pm 0.2 ^\circ C$, respectively. A comparison of the $\Delta T_m$ values of the complex and its ligands (9.8$ ^\circ C$, 3.1$ ^\circ C$ and 6.2$ ^\circ C$, respectively) was agreement with those classical intercalators [19,20,24,28], which provided strong evidence for their binding with DNA by intercalation modes.

The experimental data reveals that oxovanadium complexes in collaboration with Schiff base and imidazole[4,5-f][1,10] phenanthroline-based ligands exhibit appreciable DNA intercalative activities, it is quite consistent with their binding abilities with CT-DNA.

Photocleavage studies

The interaction mode between the oxovanadium complex and
The complexes increase from 15 μmol L⁻¹ to 60 μM (lane 3, 4 and 5 in Figures 5 and 6). Furthermore, under the same conditions with lane 4, nicked form II can barely be observed for H₂O₂ concentrations of V-complexes at 37°C for 1 h at dark condition. The cleavage reactions on plasmid DNA induced by oxovanadium complexes were investigated and monitored by gel electrophoresis [39]. The cleavage of pBR322 DNA observed from the formation of form II, indicating that the introduction of imidazole ring on the auxiliary ligands substituted with 1,10-phenanthroline as well as electrochemical characteristics of the existence of substituted electronic-donating group (OH) at the different locations introduced on the aromatic ring of phenanthroline-based ligands makes a big difference in the DNA-binding affinity.

**In vitro cytotoxicity assays**

The cytotoxicity in vitro assay for complexes 1-3 against cervical cancer CaSki and SiHa, leukemia K562, HepG2, esophagus carcinoma EC9706 and EC109 cell lines were evaluated by MTT assay. The inhibitory percentage against growth of cancer cells was determined. The cell viabilities (%) obtained with continuous exposure for 48 h are depicted in Figure 7. Cell viability decreased with increasing concentrations of 1-3. The IC₅₀ values were calculated and are listed in Table 2. As shown in Table 2, three oxovanadium complexes exhibit broad inhibition of tested cancer cell lines, with IC₅₀ values ranging from 0.41 to 89.2 μM, and the cytotoxicity was concentration-dependent.

It is notable that complex 1 possessed the most potent cytotoxicity against the cell lines of CaSki, SiHa, K562, HepG2, EC9706 and EC109. Although complexes 2 and 3 appeared to be less antiproliferative activity towards HepG2, complex 1 still show high inhibitory effect, which IC₅₀ values reaches to 0.73 μM. This is consistent with its binding abilities with CT-DNA, indicating that the antitumor abilities of the oxovanadium complexes may be closely related to their DNA binding mode.
In this paper, three oxovanadium complexes incorporating Schiff base and phenanthroline-based ligands have been synthesized and characterized. Their DNA-binding activities with CT-DNA indicate that they bind to DNA by intercalation modes and the DNA-binding affinity follows the order 1>3>2. These oxovanadium complexes can cause DNA cleavage. The results imply that the interaction with DNA may be closed associated with the introduction of imidazole ring in auxiliary ligands substituted with 1,10-phenanthroline as well as the electronic effects of substituted electronic-donating group (-OH) at the different locations introduced on the aromatic ring of phenanthroline-based ligands makes a big difference in the DNA-binding affinity. Furthermore, they also show highly cytotoxic activities against cervical, leukemia, Hepatoma and esophagus carcinoma cell lines. Complex 1 was found to be the most potent antitumor agent among the three complexes. Further investigation is required to study the possible cytotoxicity mechanisms of these complexes.

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