Original Article

Biological investigation of novel metal complexes of 2-amino-4-substituted phenylthiazole Schiff bases

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Abstract

Objective: Invasive microorganisms and free radicals are responsible for the delayed healing of various infections. It is necessary to discovery of novel molecules that are effective against invasive microorganisms and inhibit free radicals. Therefore, a series of metal complexes of 2-amino-4-substituted phenylthiazole Schiff bases were synthesized.

Methods: Structural characterization of the synthesized molecules was performed by elemental analysis, FT/IR, <sup>1</sup>H NMR, UV–Vis spectrophotometry, LC-MS, XRD, and SEM. The antimicrobial activities of all the synthesized molecules were investigated by an agar well diffusion method. An acute oral toxicity study of the synthesized ligands and their metal complexes was conducted according to OECD guidelines. The DPPH assay was used to evaluate the radical-scavenging activities of the compounds.

Results: Results of the oral acute toxicity study revealed that the synthesized analogues are safe up to a dose of 2000 mg/kg body weight. The complexes bis[{4-((4-bromo-3-methylphenyl)diazenyl)-2-((4-phenylthiazol-2-ylimino)methyl)phenoxy}cobalt (<sup>6a</sup>) and bis[{4-((4-bromo-3-methylphenyl)diazenyl)-2-((4-chlorophenyl)thiazol-2-ylimino)methyl)phenoxy}cobalt (<sup>6d</sup>) exhibited significant antibacterial activities against drug-resistant bacterial strains as well as potent radical-scavenging properties.

Conclusion: The results justify that the chelation of metals with Schiff base ligands enhances their biological activities against drug-resistant microbial strains.

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Introduction

Azo-linked Schiff bases exhibit diverse biological properties. Compounds having an azomethine (−CH≡N−) functional group are known as Schiff bases; they function as organic synthons for the synthesis of new molecules. They exhibit antitumor, anti-tubercular, antiviral, anticonvulsant, and antimicrobial activities. They are flexible and structurally similar to natural biological substances due to the presence of an imine (−N═CH−) group.

The azomethine group is a good electron donor, and forms stable complexes with transition metal ions. Both azo and azomethine groups in azo-linked Schiff bases are capable of forming coordination bonds with metal ions; however, metal ions interact preferentially with the azomethine group, leaving the azo group free. Coordination of an organic compound with a metal drastically changes its biological properties. Metal complexes of Schiff bases exhibit antitumor, antimicrobial, and antioxidant activities as well as decreased cytotoxicities compared to individual metal ions and Schiff bases. The treatment of infections is challenged by multidrug resistance in pathogenic organisms and oxidative stress. Hence, the development of newer molecules for the management of infections and oxidative stress is warranted. Metal complexation alters the therapeutic efficiency of azo-linked ligands. Azo-linked Schiff bases contain both azomethine and azo groups. Coordination of metals to such groups is responsible for their versatile biological properties. Schiff bases derived from salicylaldehydes are known polydentate ligands which can coordinate to metals in both their deprotonated and neutral forms. Salicylaldehyde-bearing molecules exhibit, antimicrobial, antiviral, cytotoxic, DNA-binding, and antioxidant activities.

The objective of the study was to synthesize a series of metal complexes retaining both azo and azomethine groups and identify molecules that are effective against drug-resistant microbial strains and possess significant antioxidant properties, thereby acting as efficient anti-infective agents. Antimicrobial activities were evaluated using drug-resistant bacterial strains. In this study, different transitional metals were conjugated with Schiff base intermediates of 5-[(4-bromo-3-methylphenyl)diazene]-2-hydroxybenzaldehyde, and their biological properties were evaluated.

Materials and Methods

All the chemicals (Merck specialties Ltd., Mumbai, India) used were of synthesis grade. Melting points were determined by the open capillary method (Elmeco), and were uncorrected. An FT/IR spectrophotometer was used to record the IR spectra of the synthesized molecules (JASCO FT/IR 4100, JASCO, JAPAN). A mass spectrophotometer with a Cs column (150 mm × 4.6 mm, 5 μm) was used to determine the molecular mass (Shimadzu). H NMR spectra were recorded using tetramethylsilane as an internal standard (Bruker H NMR, 400 MHz), and chemical shifts (δ) were reported in ppm. UV and elemental analyses were performed using a JASCO V-630 spectrophotometer and a Perkin Elmer 2400 CHNS/O analyser, respectively. XRD analysis was performed using a Cu Kα X-ray source; step = 0.02(20), run 20 = 2−80°, scanning speed = 2°/min (Shimadzu XRD 7000). Structure elucidation was performed using the Origin data analysis software. A scanning electron microscope (EGOMA 15 ZEISS) was used to study the structural environment. Magnetic susceptibility was analysed by a Faraday balance.

The synthesis of 5-[(4-bromo-3-methylphenyl)diazene]-2-hydroxybenzaldehyde (4e) was carried out as previously reported.

Synthesis of Schiff base ligands 5 (a,b)

An ethanolic solution (50 mL) of 2-amino-4-substituted phenylthiazole 4 (a,b) was mixed with an aqueous solution (100 mL) of 4e in equimolar concentrations, and the mixture was refluxed in the presence of glacial acetic acid for 2 h at 70 °C to obtain Schiff base ligands 5 (a,b). The precipitates obtained were repeatedly washed with a mixture of ethanol and diethyl ether.

Synthesis of complexes 6 (a−f)

Each Schiff base ligand (2 mmol) was mixed with an equal proportion of respective metal chlorides (Co2+; Cu2+, and Ni2+; 1 mmol) and refluxed with ethanol for 2 h at 70 °C, followed by recrystallization in ethanol. The progression of the reaction was monitored by TLC with a solvent system containing ethyl acetate and cyclohexane (1:3).

Antimicrobial activity

Antimicrobial activities of the synthesized molecules were studied by an agar well diffusion method using nutrient agar and Sabouraud dextrose agar media for bacterial and fungal pathogens, respectively. Freshly subcultured microbial strains of Klebsiella pneumoniae (MTCC 109) and Candida albicans (MTCC 3017) were procured from the Institute of Microbial Technology and Gene bank (IMTECH), Chandigarh, India. Escherichia coli (res) (resistant to norfloxacin, ofloxacin, ampicillin, cefixime, and nitrofurantoin) and Staphylococcus aureus (res) (resistant to norfloxacin, ofloxacin, ampicillin, and cefotaxime) were isolated by the Pharmaceutical Biotechnology Division of the University Department of Pharmaceutical Sciences, Utkal University, Bhubaneswar,
Incorporating the stock solution of test and reference compounds prepared with a two-fold dilution method and loaded into agar wells. Before the test was used for analysis of the data.21

**Statistical analysis**

Results

Schiff base ligands, 4-[4-bromo-3-methylphenyl]diazenyl]-2-[4-substituted phenylthiazol-2-ylamino]methyl]phenol, 5 (a–b), along with their metal complexes 6 (a–f) were synthesized as stated in Scheme I.

The physical properties of the synthesized analogues are reported in Table 1.

The far and near IR spectra of the newly synthesized compounds were studied in the frequency (v) range of 4000–4000 cm⁻¹. A band at v 1660 cm⁻¹ indicated the presence of the carbonyl stretching of salicylaldehyde in 4e. A weak absorption band appeared at v 1481–1477 cm⁻¹ in the spectra of all the compounds due to the presence of an (−N=N−) group. Broad, medium-intensity bands at v 3155 and 3159 cm⁻¹ were observed in the FT/IR spectra of Schiff bases 5 (a–b), indicating intra-molecular hydrogen bonding between hydroxyl groups and the nitrogen of azomethine. The FT/IR spectra of Schiff base ligands 5a(Lig) are reported in Figure 1. The spectra of the intermediate Schiff bases, 5a(Lig) and 5b(Lig), showed peaks indicating the azomethine group at v 1615 and 1610 cm⁻¹, respectively, whereas the spectra of the metal complexes 6 (a–f), derived from 5a(Lig) and 5b(Lig), showed moderate changes in the azomethine frequency. The phenolic (C–O) stretching vibrations of Schiff base ligands at v 1132 and 1142 cm⁻¹ were shifted in all the complexes towards the lower frequencies v 1104 and 1115 cm⁻¹. The IR spectra of the metal complexes showed bands at v 493–455 and 597–535 cm⁻¹ assigned to (M−N) and (M–O), respectively (Table 2).

The ¹H NMR spectra of 5a(Lig) and 6a are reported in Figure 2 and Figure 3, respectively. Singlet signals appeared at δ 11.34, 11.29, and 11.33 ppm in the spectra of 5a(Lig) and 5b(Lig), showed moderate changes in the azomethine frequency. The interaction of the nitrogen atom of the (−CH=N−) group with the corresponding metal chlorides in all the complexes 6 (a–f) was responsible for the sharp singlet signals in the range of δ 8.17–8.23 ppm. The sharp singlet signals observed in the spectra of all the compounds, 4e, 5 (a–b), and 6(a–f), in the range of δ 2.39–2.45 ppm corresponded to methyl protons (Table 2).

The λmax of all the synthesized molecules was studied on a UV–Vis spectrophotometer at concentrations of 10⁻³–10⁻⁶ M in ethanol. The Schiff base ligands and their metal complexes showed absorption bands in the range of 345–411 nm due to n→π* transitions indicating the presence of −N=N− in all the molecules (Table 2). The solvatochromic effects of compounds 4e and 5a(Lig), and metal complexes 6(a–c) are illustrated in Figure 4. The predicted molecular weights of the synthesized compounds were confirmed by LC-MS (Table 1). Compound 5a(Lig) was predicted to have the molecular formula C₂₃H₁₇BrN₆O₅ due to the molecular ion peak at 477.3 m/z (Figure 5). Elemental analysis of the synthesized molecules verified their molecular formulas.

Structures of the Schiff base 5b(Lig) and its cobalt complex (6d) based on XRD analysis are shown in Figure 6, which reveals two strong and three moderate reflections shown by 5b(Lig) in the 2θ range of 30–80° (Figure 6a), whereas 6d showed five strong and three moderate
Scheme I: Synthesis of metal complexes of Schiff bases derived from 2-amino-4-substituted phenylthiazole. i. NaNO₂/HCl, 0–5 °C, diazotization; ii. 10% NaOH, coupling reaction; iii. Ethanol reflux, 2 h, 70 °C, Schiff base; iv. MCl₂/x H₂O, ethanol (M = Co²⁺, Cu²⁺, or Ni²⁺).

X = H or Cl
| Compounds | Chemical name                                                                 | M. formula       | LC-MS (RT, % area) | m/z       | Rf | M. P. (°C) | Colour     | Yield (%) |
|-----------|-------------------------------------------------------------------------------|------------------|--------------------|-----------|----|-----------|------------|-----------|
| 4e        | 5-[(4-Bromo-3-methylphenyl)diazenyl]-2-hydroxybenzaldehyde                   | C_{14}H_{11}BrN_{2}O_{2} | 3.053, 95.57       | 319.01 (M + 1) | 0.8 | 180–90    | Dark red (powder) | 85        |
| 5a(Lig)   | 4-[(4-bromo-3-methylphenyl)diazenyl]-2-[(4-phenylthiazol-2-ylimino)methyl]phenol | C_{23}H_{17}BrN_{4}OS | 1.581, 99.93       | 477.3 (M - 1)  | 0.6 | 235–37    | Brown (crystal)  | 74        |
| 5b(Lig)   | 4-[(4-bromo-3-methylphenyl)diazenyl]-2-[(4-(4-chlorophenyl)thiazol-2-ylimino)methyl]phenol | C_{23}H_{16}BrClN_{4}OS | 1.861, 89.73       | 512.3 (M + 1)  | 0.5 | 242–45    | Brown (crystal)  | 78        |
| 6a        | Bis[4-((4-bromo-3-methylphenyl)diazenyl)-2-((4-phenylthiazol-2-ylimino)methyl)phenoxy]cobalt | C_{46}H_{32}Br_{2}CoN_{8}O_{2}S_{2} | 1.761, 91.73       | 1009.3 (M + 1) | 0.6 | 258–60    | Brown (powder)  | 43        |
| 6b        | Bis[4-((4-bromo-3-methylphenyl)diazenyl)-2-((4-phenylthiazol-2-ylimino)methyl)phenoxy]copper | C_{46}H_{32}Br_{2}CuN_{8}O_{2}S_{2} | 1.351, 97.43       | 1013 (M + 1)   | 0.7 | 262–65    | Brown (powder)  | 49        |
| 6c        | Bis[4-((4-bromo-3-methylphenyl)diazenyl)-2-((4-phenylthiazol-2-ylimino)methyl)phenoxy]nickel | C_{46}H_{32}Br_{2}NiN_{8}O_{2}S_{2} | 1.661, 87.73       | 1008.79 (M - 1) | 0.8 | 253–58    | Brown (powder)  | 51        |
| 6d        | Bis[4-((4-bromo-3-methylphenyl)diazenyl)-2-((4-(4-chlorophenyl)thiazol-2-ylimino)methyl)phenoxy]cobalt | C_{46}H_{30}Br_{2}Cl_{2}CoN_{8}O_{2}S_{2} | 1.561, 88.73       | 1079.1 (M + 1) | 0.7 | 238–40    | Brown (powder)  | 53        |
| 6e        | Bis[4-((4-bromo-3-methylphenyl)diazenyl)-2-((4-(4-chlorophenyl)thiazol-2-ylimino)methyl)phenoxy]copper | C_{46}H_{30}Br_{2}Cl_{2}CuN_{8}O_{2}S_{2} | 1.356, 81.73       | 1085.31 (M + 1) | 0.8 | 246–50    | Brown (powder)  | 73        |
| 6f        | Bis[4-((4-bromo-3-methylphenyl)diazenyl)-2-((4-(4-chlorophenyl)thiazol-2-ylimino)methyl)phenoxy]nickel | C_{46}H_{30}Br_{2}Cl_{2}NiN_{8}O_{2}S_{2} | 1.531, 79.73       | 1081.10 (M + 2) | 0.6 | 253–55    | Brown (powder)  | 67        |

**Figure 1:** FT/IR spectra of 4-[(4-bromo-3-methylphenyl)diazenyl]-2-[(4-phenylthiazol-2-ylimino)methyl]phenol: 5a(Lig).

Schiff base analogues as potent biomolecules
Table 2: Spectral characterization of novel metal complexes of 2-amino-4-substituted phenylthiazole Schiff bases.

| Compound | UV–Vis λ_{max} (nm) | IR (cm^{-1}) | ^1H NMR (δ, ppm) | Elemental Analysis Calculated (Found) % |
|----------|----------------------|--------------|------------------|-----------------------------------------|
|          | (Ethanol)            |              |                  |                                         |
| 4e       | 346                  | 3444 (O–H str.), 2857, 2743 (CH str. of aldehyde), 1660 (C=O str.), 1615 (C=C str.), 1478 (N=N=N=), 1305, 1175 (SO$_2$ str.,SO$_3$H), 1137 (C=O str.); (CDCl$_3$, 400 MHz): 2.50 (s, 3H, CH$_3$), 7.26 (d, 1H, salicylaldehyde H-5), 7.69–7.78 (d, 2H, Ar H), 7.78 (s, 1H, Ar H), 8.20 (s, 1H, salicylaldehyde H-2), 8.17 (d, 1H, salicylaldehyde H-6), 10.38 (s, 1H, CHO), 11.34 (s, 1H, OH); | 52.69 | 3.47 | 8.78 | – |
| 5a(Lig)  | 403                  | 3155 (O–H str. intra mol. H.), 1615 (C=N str.), 1481 (N=N=N=), 1284 (OH bend), 1132 (C=O str.), 845 (C=Br str.); (CDCl$_3$, 400 MHz): 2.49 (s, 3H, CH$_3$), 7.25 (d, 1H, salicylaldehyde H-6), 7.43–7.59 (m, 5H, ArH), 7.67 (d, 1H, diazenylphenyl H-5), 7.76 (s, 1H, diazenylphenyl H-2), 8.17 (s, 1H, –CH=N), 8.18 (s, 1H, Thiazolyl H-5), 8.72 (d, 1H, salicylaldehyde H-5), 9.42 (s, 1H, salicylaldehyde H-3), 11.33 (s, 1H, OH); | 57.87 | 3.59 | 11.74 | 6.72 |
| 5b(Lig)  | 411                  | 3159 (O–H str. intra mol. H.), 1610 (C=N str.), 1477 (N=N=N=), 1265 (OH bend), 1142 (C=O str.), 841 (C=Br str.), 835 (1,4 disubst.); (CDCl$_3$, 400 MHz): 2.39 (s, 3H, CH$_3$), 7.24 (d, 1H, salicylaldehyde H-6), 7.53–7.98 (d, 4H, ArH), 7.69 (d, 1H, diazenylphenyl H-5), 7.71 (d, 1H, diazenylphenyl H-6), 7.76 (s, 1H, diazenylphenyl H-2), 8.13 (s, 1H, –CH=N), 8.16 (s, 1H, Thiazolyl H-5), 8.73 (d, 1H, salicylaldehyde H-5), 9.13 (s, 1H, salicylaldehyde H-3), 11.29 (s, 1H, OH); | 53.97 | 3.15 | 10.95 | 6.26 |
| 6a       | 375                  | 1614 (C=N str.), 1470 (N=N=N=), 1104 (C=O str.), 815 (C=Br str.), 457 (Co–N), 593(Co–O); (DMSO-d$_6$, 400 MHz): 2.50 (s, 3H, CH$_3$), 7.25 (d, 1H, salicylaldehyde H-6), 7.44–7.60 (m, 10H, ArH), 7.66 (d, 1H, diazenylphenyl H-5), 7.77 (s, 1H, diazenylphenyl H-2), 8.17 (s, 1H, –CH=N), 8.19 (s, 1H, Thiazolyl H-5), 8.72 (d, 1H, salicylaldehyde H-5), 9.45 (s, 1H, salicylaldehyde H-3); | 54.61 | 3.19 | 11.08 | 6.34 |
| 6b       | 392                  | 1612 (C=N str.), 1465 (N=N=N=), 1115 (C=O str.), 814 (C=Br str.), 455 (Cu–N), 555 (Cu–O) cm$^{-1}$; (DMSO-d$_6$, 400 MHz): 2.50 (s, 3H, CH$_3$), 7.26 (d, 1H, salicylaldehyde H-6), 7.39–7.66 (m, 10H, ArH), 7.67 (d, 1H, diazenylphenyl H-5), 7.77 (s, 1H, diazenylphenyl H-2), 8.17 (s, 1H, –CH=N), 8.20 (s, 1H, Thiazolyl H-5), 8.79 (d, 1H, salicylaldehyde H-5), 9.63 (s, 1H, salicylaldehyde H-3); | 54.36 | 3.17 | 11.03 | 6.31 |
| 6c       | 350                  | 1611 (C=N str.), 1482 (N=N=N=), 1105 (C=O str.), 816 (C=Br str.), 456 (Ni–N), 593 (Ni–O) cm$^{-1}$; (DMSO-d$_6$, 400 MHz): 2.50 (s, 3H, CH$_3$), 7.26 (d, 1H, salicylaldehyde H-6), 7.38–7.67 (m, 10H, ArH), 7.69 (d, 1H, diazenylphenyl H-5), 7.77 (s, 1H, diazenylphenyl H-2), 8.17 (s, 1H, –CH=N), 8.19 (s, 1H, Thiazolyl H-5), 8.79 (d, 1H, salicylaldehyde H-5), 9.46 (s, 1H, salicylaldehyde H-3); | 54.62 | 3.19 | 11.08 | 6.34 |
| 6d       | 345                  | 1613 (C=N str.), 1472 (N=N=N=), 1109 (C=O str.), 837 (C=Br str.), 463 (Co–N), 597 (Co–O); (DMSO-d$_6$, 400 MHz): 2.45 (s, 3H, CH$_3$), 7.47 (d, 1H, salicylaldehyde H-6), 7.51–7.71 (m, 10H, ArH), 7.73 (d, 1H, diazenylphenyl H-5), 7.79 (s, 1H, diazenylphenyl H-2), 8.21 (s, 1H, –CH=N), 8.28 (s, 1H, Thiazolyl H-5), 8.89 (d, 1H, salicylaldehyde H-5), 9.51 (s, 1H, salicylaldehyde H-3); | 51.13 | 2.80 | 10.37 | 5.93 |

(continued on next page)

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reflections in the 2θ range of 18–80° (Figure 6b) corresponding to their inter-planar distances d = 7.03 and 2.82 Å, respectively.

SEM analysis of 5b(Lig) and its cobalt complex 6d suggested that the particle size of 6d significantly differed from the particle size of 5b(Lig) due to the presence of an auxiliary molecule. The particle diameter of 5b(Lig) was 355.2 nm at an external angle of 360° (Figure 7a), whereas it was 366 nm at an external angle of 355.2° for its cobalt complex (Figure 7b).

The magnetic susceptibilities of the Co²⁺ complex were 5.09 and 5.11 BM, indicating an octahedral symmetry.¹³ The Ni²⁺ complex showed magnetic susceptibilities of 2.91 and 2.99 BM, whereas the Cu²⁺ complex showed magnetic susceptibilities of 2.01 and 1.93 BM, suggesting that they have an octahedral geometry. Compounds 5a(Lig) and 5b(Lig) act as bidentate ligands by chelating with transitional metal ions through the azomethine bearing nitrogen atom. Based on magnetic susceptibilities, the presumed structures of the complexes are shown in Scheme I.
All the compounds showed good antimicrobial activities (Table 3). Compound 4e, Schiff base 5a(Lig), and metal complexes 6a, 6b, and 6d showed significant antimicrobial activities against *E. coli* (res), *K. pneumoniae*, and *S. aureus* (res) in comparison to both amoxicillin (RA) and ampicillin (RA1). The cobalt–conjugated complex 6a showed significant (P < 0.05) antimicrobial activities against *E. coli* (res), *K. pneumoniae*, and *S. aureus* (res) with mean zones of inhibition as 15.5 ± 3.21, 16.17 ± 2.99, and 16.83 ± 2.71 mm, respectively in comparison to both amoxicillin (RA) and ampicillin (RA1). The antibiograms of the synthesized molecules are shown in Figure 8. The antimicrobial activities of the synthesized molecules against *E. coli* (res), *S. aureus* (res), *C. neoformans*, and *C. albicans* in comparison to that of amoxicillin (RA) are graphically presented in Figure 9.

Molecules 4e, 5a(Lig), 5b(Lig) 6a, 6b, and 6d inhibited the growth of *E. coli* (res), *K. pneumoniae*, and *S. aureus* (res) at an MIC of 31.25 µg/mL (Table 4). All the complexes showed potential antimicrobial activities against *K. pneumoniae* at 31.25 µg/mL, whereas Schiff base ligands 5a(Lig) and 5b(Lig) were active only at 125 µg/mL.

The results of the oral acute toxicity study revealed no mortality, toxicity, and gross behavioural changes in the experimental rats. All the compounds are found to be safe up to a dose of 2000 mg/kg body weight.

The radical scavenging activities of the newly synthesized molecules are reported in Table 5. All the test compounds (4e–6f) showed 50% inhibition at concentrations of 72 ± 1.4, 56 ± 0.78, 52 ± 1.13, 36 ± 0.56, 48 ± 0.89, 40 ± 0.59, 38 ± 1.4, 46 ± 0.98, 37.3 ± 0.02, and 31 ± 0.70 µg/mL, respectively. Molecules 6a, 6c, and 6d showed significant radical scavenging activities (>61%) at a concentration of 50 µg/mL in comparison to standard BHT. However, the cobalt–conjugated complexes 6a and 6d showed significantly low IC50 values of 36 and 38 µg/mL, respectively. The antioxidant activities of the synthesized molecules showing % inhibition at different concentrations are graphically presented in Figure 10.
Table 3: Antimicrobial activities of novel metal complexes of 2-amino-4-substituted phenylthiazole Schiff bases against different microbial strains (mean ± S.D.).

| Compounds | Bacterial strain |   |   | Fungal strain |
|-----------|------------------|---|---|---------------|
|           | E. coli (res)    | K. pneumoniae | S. aureus (res) | C. albicans | C. neoformans |
| 4e        | 15.83 ± 1.47*    | 15.17 ± 2.14 | 16.17 ± 2.04*    | 10.17 ± 1.84 | 12.5 ± 1.52   |
| 5a(Lig)   | 13.67 ± 2.34     | 12.67 ± 2.25 | 15.83 ± 2.79*    | 12.17 ± 2.23 | 13.33 ± 1.37  |
| 5b(Lig)   | 15.33 ± 2.07     | 13.5 ± 2.43  | 11.5 ± 1.87      | 12 ± 1.27    | 12.33 ± 1.75  |
| 6a        | 16.67 ± 2.34*    | 16.17 ± 1.94* | 20.83 ± 1.84*    | 12.83 ± 2.23 | 13.67 ± 2.94  |
| 6b        | 15.83 ± 1.84*    | 16.3 ± 2.37*  | 18.33 ± 1.75*    | 13.33 ± 2.34 | 12.67 ± 1.51  |
| 6c        | –                | 14.67 ± 3.14  | 14.83 ± 1.84     | 11.33 ± 1.51 | 15.83 ± 1.84  |
| 6d        | 15.5 ± 3.21*     | 16.17 ± 2.99* | 16.83 ± 2.71*    | 12.67 ± 2.34 | 12.17 ± 2.71  |
| 6e        | 14.83 ± 2.56     | 15.83 ± 2.23  | 14.5 ± 1.52      | 11.17 ± 6.21 | 10.5 ± 1.52   |
| 6f        | –                | 16.67 ± 1.37* | 15.17 ± 1.94     | 12.5 ± 1.52  | 10.33 ± 1.75  |
| RA        | 12.67 ± 1.51     | 15.33 ± 1.97  | 13 ± 1.67        | 19.33 ± 4.68 | 24.17 ± 1.94  |
| RA¹       | –                | 15.09 ± 1.13  | –               | –            | –             |

Results expressed in Mean ± S.D. (n = 6). The data were analysed by One Way ANOVA followed by Dunnett’s Post Hoc test, (statistical significance at *p < 0.05 in comparison to reference antibiotic (RA): amoxicillin (antibacterial), fluconazole (antifungal) and RA¹: ampicillin (antibacterial); -, no zone of inhibition.)

Figure 5: LCMS of 4-[(4-bromo-3-methylphenyl)diazenyl]-2-[(4-phenylthiazol-2-ylimino)methyl]phenol: 5a(Lig).

Figure 6: X-ray Diffraction pattern of (a): 4-[(4-bromo-3-methylphenyl)diazenyl]-2-[(4-(4-chlorophenyl)thiazol-2-ylimino)methyl]phenol 5b(Lig): and (b): bis[4-[(4-bromo-3-methylphenyl)diazenyl]-2-[(4-(4-chlorophenyl)thiazol-2-ylimino)methyl]phenoxy]cobalt (6d).
Discussion

Weak absorption bands at $n$ 1481-1477 cm$^{-1}$ in the FT/IR spectra of 5a and 5b confirm the existence of an $-N=N-$ group; the azomethine group preferentially forms a coordination bond with the metal ion, whereas the azo group is left free, as indicated by an unaltered $-N=N-$ band.8 The absence of a carbonyl stretching band of 4e at $n$ 1660 cm$^{-1}$ in the spectra of 5a and 5b confirmed the formation of an azomethine moiety. The spectra of the metal complexes 6a–f derived from 5a and 5b showed moderately altered frequencies for the azomethine group in the range of $n$ 1614–1609 cm$^{-1}$. Broad medium-intensity bands at 3155 and 3159 cm$^{-1}$ observed in the spectra of Schiff bases 5a and 5b, respectively, indicate intramolecular hydrogen bonding between hydroxyl groups and the azomethine nitrogen.

Singlet peaks due to the phenolic $-OH$ proton of salicylaldehyde were observed in the NMR spectra of Schiff bases 5a and 5b at $\delta$ 11.29 and 11.33 ppm, respectively; they were absent in the spectra of all the metal complexes 6a–f due to the deprotonation of the hydroxyl proton by the interaction with metal chlorides. Because the nitrogen atom of the $-CH=N-$ group formed coordinate bonds with metal ions, all the metal complexes 6a–f showed sharp singlets in the range of $\delta$ 8.17–8.23 ppm corresponding to the azomethine proton.

XRD analysis of cobalt complex 6d revealed three moderate reflections at 20°, 27°, and 40°, which were not shown by 5b, verifying the conjugation of the metal (cobalt) with the Schiff base.

Complexes 6a and 6d exhibited better antibacterial activities against E. coli (res), K. pneumoniae, and S. aureus.

Figure 7: SEM micrograph of (a): 4-[(4-bromo-3-methylphenyl)diazenyl]-2-[(4-(4-chlorophenyl)thiazol-2-ylimino)methyl]phenol 5b and (b): bis-4-[(4-bromo-3-methylphenyl)diazenyl]-2-[(4-(4-chlorophenyl)thiazol-2-ylimino)methyl]phenoxy]cobalt (6d).

Figure 8: Antibiogram pattern of 2-amino-4-substituted phenylthiazole based Schiff base bearing metal complexes against S. aureus (res) (a) and C. neoformans (b).
Figure 9: Error gram showing the statistical interpretation of antimicrobial activity of 2-amino-4-substituted phenylthiazole based Schiff base bearing metal complexes against (a) *E. coli* (res), (b) *S. aureus* (res), (c) *C. albicans*, and (d) *C. neoformans*. 
than their respective Schiff bases due to the insertion of the Co$^{2+}$ ion to the intermediate Schiff bases 5a(Lig) and 5b(Lig), respectively. The coordination of metals with ligands reduces the polarity and enhances the lipophilicity of the resulting molecules, increasing their permeation across the lipoidal cell membranes of microorganisms, causing increased inhibition of microbial growth.14

The variation in the DPPH scavenging capacity of the different molecules may be attributed to the substitution of the various metals to the Schiff base ligands, which increases the capacity to stabilize unpaired electrons, thereby scavenging free radicals.11 Our findings are consistent with previous reports that metal complexes of Schiff bases show improved antimicrobial and antioxidant activities. This study is a continuation of our earlier reported study.6

Table 4: Minimum inhibitory concentration (MIC, µg/mL) of novel metal complexes of 2-amino-4-substituted phenylthiazole Schiff bases against different microbial strains.

| Compounds | Bacterial strain | Fungal strain |
|-----------|-----------------|--------------|
|           | E. coli (res) | K. pneumoniae | S. aureus (res) | C. albicans | C. neoformans |
| 4e        | 31.25 | 31.25 | 31.25 | 500 | 250 |
| 5a(Lig)   | 125  | 125  | 31.25 | 250 | 250 |
| 5b(Lig)   | 31.25 | 125  | 250  | 250 | 250 |
| 6a        | 31.25 | 31.25 | 31.25 | 125 | 250 |
| 6b        | 31.25 | 31.25 | 31.25 | 250 | 250 |
| 6c        | 62.5 | 62.5 | 250  | 250 | 250 |
| 6d        | 62.5 | 31.25 | 250  | 250 | 250 |
| 6e        | 31.25 | 31.25 | 31.25 | 125 | 250 |
| 6f        | --  | 31.25 | 31.25 | 250 | 250 |

—, no zone of inhibition; E. coli(res), Escherichia coli resistant; S. aureus(res), Staphylococcus aureus resistant; C. albicans, Candida albicans; C. neoformans, Cryptococcus neoformans.

Table 5: DPPH radical scavenging activities of novel metal complexes of 2-amino-4-substituted phenylthiazole Schiff bases.

| Conc. (µg/mL) | 10 | 30 | 50 | 70 | IC$_{50}$ |
|--------------|----|----|----|----|---------|
| Compound     | % Inhibition | % Inhibition | % Inhibition | % Inhibition | % Inhibition |
| 4e           | 17.96 ± 0.03 | 29.54 ± 0.05 | 33.64 ± 0.09 | 56.39 ± 0.08 | 72 ± 1.4 |
| 5a(Lig)      | 27.99 ± 0.04 | 39.34 ± 0.02 | 49.09 ± 0.07 | 63.9 ± 0.05 | 56 ± 0.78 |
| 5b(Lig)      | 19.26 ± 0.04 | 38.53 ± 0.09 | 48.33 ± 0.07 | 79.89 ± 0.06 | 52 ± 1.13 |
| 6a           | 29.72 ± 0.07 | 43.32 ± 0.11 | 63.44 ± 0.07* | 83.02 ± 0.06 | 36 ± 0.56 |
| 6b           | 28.43 ± 0.07 | 41.83 ± 0.02 | 51.41 ± 0.05 | 79.97 ± 0.06 | 48 ± 0.89 |
| 6c           | 25.43 ± 0.08 | 39.65 ± 0.09 | 61.41 ± 0.04* | 83.11 ± 0.04 | 40 ± 0.59 |
| 6d           | 36.53 ± 0.02* | 43.03 ± 0.04 | 61.97 ± 0.08* | 73.78 ± 0.07 | 38 ± 1.4 |
| 6e           | 31.24 ± 0.06 | 41.31 ± 0.02 | 51.77 ± 0.08 | 79.89 ± 0.04 | 46 ± 0.98 |
| 6f           | 39.73 ± 0.02* | 53.38 ± 0.03* | 63.38 ± 0.01* | 79.53 ± 0.02 | 37 ± 0.02 |
| BHT          | 32.32 ± 0.01 | 49.54 ± 0.01 | 59.99 ± 0.10 | 87.74 ± 0.06 | 31 ± 0.70 |

All values are expressed as Mean ± S.D., (n = 3), The data were analysed by one-way ANOVA followed by Dunnett’s post hoc test, (statistical significance at *p < 0.05 in comparison to standard).
Conclusions

We previously showed that the complex bis[2-{(E)-4-methoxyphenylimino)methyl}-4-{(E)-3-nitrophenyl}diazonyl phenoxycobalt], 5a([Co L1]), exhibits significant antimicrobial activity.6 In the present study, we consistently observed that the complexes bis[4-([(4-bromo-3-methyl-phenyldiazenyl)])-2-[(4-phenylthiazol-2-ylmino)methyl]phenoxycobalt, 6a, and bis[4-[(4-bromo-3-methylphenyldiazenyl)]-2-[(4-chlorophenyl)thiazol-2-ylmino)methyl]phenoxycobalt, 6d, exhibit significant biological activities. We reported earlier that the complex, 5a([Co L2]), exhibited significant antibacterial activity against K. pneumoniae. In the present study, the complexes 6a and 6d exhibited significant antibacterial activities against drug-resistant pathogens, E. coli (res), K. pneumonia, and S. aureus (res). In the previous study, all the complexes showed significant antifungal activities; however, none of the complexes in the present study exhibited a significant antifungal activity. Comparing the antimicrobial activities of the complexes in our earlier report with those of the complexes in the present study, we found that complexes bearing the 3-nitrophenylazo salicylalddehyde nucleus exhibit excellent antifungal activities, whereas those with the 4-bromo-3-methylphenylazo salicylalddehyde nucleus showed significant antibacterial activities against drug-resistant bacterial strains, despite both the Schiff base complexes bearing the azosalicylalddehyde nucleus. However, the molecules described in the present study also showed significant antioxidant activities. Based on our findings on the biological activities of previously and recently synthesized complexes, it can be concluded that chelating the transitional metal Co2+ with Schiff base ligands bearing an azosalicylaldehyde nucleus enhances their biological activities. Because the novel synthesized metal complexes of 2-amino-4-substituted phenylthiazole Schiff bases showed potential antimicrobial activities against drug-resistant pathogens and exhibited improved radical scavenging activities, they may be effective in the treatment of infections.

Conflict of interest

The authors have no conflict of interest.

Author contribution

The concept of the work was fabricated by SKP. The experimental work and the drafting of the manuscript was performed by JS. Both the authors equally contributed to this research.

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Appendix A. Supplementary data

Supplementary data related to this article can be found at https://doi.org/10.1016/j.jtumed.2017.10.007.

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