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1. Introduction

Mild steel is one of the most used steels due to its relative lowcost and good mechanical strength.¹⁻³ Mild steel is widely used in the petroleum industry as transport pipes for oil in petroleum production and refining, chemical processing industries, engineering materials in transportation, pipeline industries, mining and marine applications, nuclear power plants, construction and metal processing equipment.^{4,5} Mild steel is, however,

Synthesis and structures of divalent Co, Ni, Zn and Cd complexes of mixed dichalcogen and dipnictogen ligands with corrosion inhibition properties: experimental and computational studies[†]

Tunde L. Yusuf,^a Taiwo W. Quadri, ^b Gideon F. Tolufashe, ^c Lukman O. Olasunkanmi, ^b Cno E. Ebenso ^b and Werner E. van Zyl

The structural and corrosion inhibition properties of four different transition-metal complexes of heteroleptic S-donor atom dithiophosphonate and N-donor atom phenanthroline ligands are reported. Full structural characterization of the Co, Ni, Zn and Cd complexes was achieved with the aid of singlecrystal X-ray crystallography. Structural elucidation revealed the formation of a 4-coordinate Zn(II) complex, and 6-coordinate Ni(II) and Cd(II), as well as a novel dithiophosphonato Co(II) complex. The ability of the complexes with this ligand type to act as inhibitors of mild steel corrosion in 1 M HCl solution is reported for the first time. Corrosion inhibition potentials of the complexes were assessed using potentiodynamic polarization, electrochemical impedance spectroscopy (EIS), and density functional theory (DFT). The open circuit potential (OCP) time profile showed the system achieved a steady-state potential before the first 600 s after submerging the working electrode in the corrosive medium. The studied metal complexes are good inhibitors of mild steel corrosion in 1 M HCl and were found to retard the corrosion rate by forming an adsorbed pseudocapacitive film on the steel surface. The order of inhibition efficiencies was in the order Ni (94.14%) > Cd (92.28%) > Zn (91.14%) > Co (72.53%).

> susceptible to corrosion, especially in acidic conditions and can readily be exposed to acid during acid cleaning, acid pickling and acid descaling, which are famous industrial procedures.⁶⁻¹⁰ The use of corrosion inhibitors in different media has become an efficient and common way of protecting metallic surfaces from corrosion¹¹⁻¹³ because it is easy to apply and relatively inexpensive.^{4,13,14}

> Organic compounds with heteroatoms like O, S, P and N often display potentials to inhibit metal corrosion in different electrolytic media.15,16 The inhibition properties of these classes of compounds are attributed to their strong polarizability and low electronegativity. The presence of heteroatoms in corrosion inhibitors facilitates transfer of electrons to empty atomic orbitals in the metal, thereby promotes adsorption of the inhibitor molecule on the metallic surface.17 Reports showed that nitrogen-containing organic compounds are excellent anticorrosion agents for metals in hydrochloric acid solution, while sulfur-containing compounds have the tendency to perform better in inhibiting metal corrosion in sulphuric acid solution.¹⁸ Furthermore, compounds containing N- and S- atoms usually exhibit substantial inhibition potentials in both hydrochloric and sulphuric acid solutions.19 The efficiency of inhibitors with different heteroatoms usually increases with decreasing

^aSchool of Chemistry and Physics, University of KwaZulu-Natal, Westville Campus, Chiltern Hills, Private Bag X54001, Durban, 4000, South Africa. E-mail: vanzylw@ ukzn.ac.za; Tel: +27 31 260 3188

^bDepartment of Chemistry, School of Physical and Chemical Sciences, Materials Science Innovation and Modelling (MaSIM) Research Focus Area, Faculty of Natural and Agricultural Sciences, North-West University (Mafikeng Campus), Private Bag X2046, Mmabatho 2735, South Africa

^cDepartment of Chemistry and Biochemistry, Faculty of Sciences, University of Porto, 4169-007 Porto, Portugal

^dDepartment of Chemistry, Faculty of Science, Obafemi Awolowo University, Ile-Ife 220005, Nigeria

^{*}Institute for Nanotechnology and Water Sustainability, College of Science, Engineering and Technology, University of South Africa, Johannesburg, South Africa † Electronic supplementary information (ESI) available. CCDC 2008849–2008851, 2008799, 2008800. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d0ra07770d

electronegativity, such that the trend of inhibition efficiencies with respect to heteroatoms is in the order O < N < S < P.²⁰

Metal complexes have also been reported to show excellent corrosion inhibition activities.^{21,22} The interaction of transition metal complexes with mild steel is significantly affected by the reactivity and nature of the ligand and the standard electrode potential of the metal cation.²² The standard electrode potential of divalent cations considered in the present study follows the order: Ni(π)/Ni(-0.23 V) > Co(π)/Co (-0.277 V) > Cd(π)/Cd (-0.40 V) > Zn(π)/Zn (-0.76 V)²³ and the incorporation of metal into a heteroatom-rich ligand should improve its corrosion inhibition potentials.

Electrochemical techniques like potentiodynamic polarization (PDP) and electrochemical impedance spectroscopy (EIS) are powerful techniques that are widely used to research electrochemical corrosion. Inhibition efficiencies of corrosion inhibitors, and organic and inorganic coatings are usually estimated from PDP and EIS measurements.^{24–27} Computational studies are often used to provide corroborative explanations to the experimental findings.²⁸ Quantum chemical calculations are employed to investigate possible correlations between molecular quantum chemical descriptors and specific reactivities of a molecular system, including adsorption behaviour, which translates to corrosion inhibition^{14,29–31}

Inspired by the application of zinc dialkyldithiophosphates (ZDDPs) as an antioxidant in the oil industry,³² we synthesized and characterized four new transition-metal complexes (Co^{2+} , Ni^{2+} , Zn^{2+} and Cd^{2+}) of dithiophosphonate and N-donor (2,2-bipyridine, 1,10-phenanthroline) ligands, and assessed their corrosion inhibition properties for mild steel in 1 M HCl. Additionally, single crystal X-ray crystallography revealed new structures and were further supported by elemental analysis, magnetic susceptibility measurements, multi-nuclear magnetic resonance, Fourier transform infrared and electronic spectroscopies, and mass spectrometry.

2. Experimental

2.1 Materials and methods

All reactions were performed using standard Schlenk line techniques under a nitrogen atmosphere. All chemicals were purchased from commercial sources. Tetrahydrofuran (THF) and dichloromethane (DCM) was dried using Innovative Technology PURE-SOLV solvent purification system, USA. NMR spectra were recorded on a Bruker AV-400 instrument operating at 400 MHz while ¹H and ³¹P was recorded at 162 MHz. For the ¹H residual solvent proton a standard reference was used $(\delta, \text{ppm}, \text{CDCl}_3, 7.26)$. For ³¹P NMR, H₃PO₄ (85%) was used as an external reference. The melting points were measured on an Electrothermal 9100 melting point apparatus and are uncorrected. The electronic absorption spectra were recorded on a Shimadzu UV-3600 UV-VIS-NIR spectrophotometer using quartz cuvettes having a path length of 1 cm in the range 200-400 nm for UV and 400-900 nm visible regions. Electrospray ionization mass spectra (ESI-MS) for the complexes were recorded using a Waters Micromass LCT Premier TOF-MS. The LC/MS spectra of the ligand and some complexes were obtained

from Shimadzu LCMS-2020 using ESI methods. The molar conductance of 10^{-3} M complexes in DMSO solutions was obtained at room temperature on an Edge HI2003 conductivity meter.

2.2 Synthesis of ligand

2.2.1 Synthesis of $[NH_4][S_2P{1,4-C_6H_4OMe}{OCH(CH_3)_2}]$ (L1). An oven dried Schlenk flask was charged with Lawesson's reagent (1.0 g, 2.47 mmol) and iso-propanol (0.4 mL, 4.94 mmol) in a dry THF mixture (30 mL) and stirred for 30 minutes until the solution was clear. The solution was cooled to 0 °C in an ice-bath and anhydrous NH₃ gas was bubbled into the clear solution to afford a white precipitate. The mixture was concentrated and further stirred in dry hexane (30 mL). A colourless free flowing powder was obtained upon removal of the solvent under reduced pressure. Yield: 1.31 g (96%) melting point: 112–114 °C ¹H NMR (400 MHz, DMSO- d_6) δ 7.91 (dd, J =12.8, 8.6 Hz, 2H, Ar-H), 7.24 (bs, 4H, NH_4), 6.84 (dd, J = 8.8, 2.4 Hz, 2H, Ar-H), 4.59-4.40 (m, 1H, OCH), 3.76 (s, 3H, Ar- OCH_3 , 1.04 (d, J = 6.2 Hz, 6H, CH_3). ¹³C NMR (101 MHz, DMSO d_6) δ 160.01, 139.21, 138.11, 131.96, 131.83, 113.57, 112.39, 112.26, 66.42, 66.36, 55.51, 24.66, 24.63. LC/MS MS: m/z 263 ([M + 2H]⁺) IR(ATR, cm⁻¹) 2838(s), 1594(s), 1571(s), 1496(s), 1464(s), 1439(s), 1406(s), 1303(s), 1291(s). 1241(s), 1217(s), 1178(s), 1045(s), 1019(s), 925(s), 862(s), 824(s).

2.2.2 Synthesis of $[NH_4][S_2P{1,4-C_6H_4OMe}{OCH_2CH_2-CH_3}]$ (L2). The ligand was prepared similar to ligand L1 above. Yield: 1.35 g (98%) melting point: 118–119 °C. ¹H NMR (400 MHz, DMSO-*d*₆) δ 7.90 (dd, *J* = 12.9, 8.6 Hz, 2H, Ar–H), 6.86 (dd, *J* = 8.8, 2.4 Hz, 2H, Ar–H), 6.81 (bs, 4H, NH₄), 3.75 (s, 3H, Ar–OCH₃), 3.64–3.56 (m, 2H, OCH₂), 1.46 (h, *J* = 7.2 Hz, 2H, CH₂), 0.79 (t, *J* = 7.4 Hz, 3H, CH₃). ¹³C NMR (101 MHz, DMSO-*d*₆) δ 160.20, 160.17, 138.12, 137.02, 131.91, 131.78, 112.62, 112.48, 64.49, 64.42, 55.58, 23.87, 23.78, 10.95. LC/MS MS: *m/z* 263 ([M + 2H]⁺), 523 (2[M]⁺). IR(ATR, cm⁻¹) 2836(s), 1586(s), 1570(s), 1491(s), 1458(s), 1440(s), 1313(s), 1295(s). 1238(s), 1219(s), 1179(s), 1028(s), 924(s), 860(s), 826(s).

2.3 Synthesis of metal complexes

2.3.1 Synthesis of $[Co{S_2P(1,4-C_6H_4OMe)(OCH(CH_3)_2)}_2{-phen}]$ (1). A mixture of $CoCl_2 \cdot 6H_2O$ (238 mg, 1 mmol) and L1 (559 mg, 2 mmol) in ethanol solution (25 mL) was stirred in a beaker for 25 minutes. To the resulting blue mixture was added 1,10-phenanthroline monohydrate (198 mg, 1 mmol) and stirred for 1 hour. The resulting brown precipitate was filtered and washed with water/ethanol mixture (50 : 50). Single crystals suitable for X-ray diffraction were obtained by vapour diffusion of hexane into DCM. Yield: 640 mg (84%). Melting point: 186–188 °C LC-MS(m/z) (cal.) 761(761) for ($[M]^+$; 500 (500) ($[M-{S_2P(4-C_6H_4OCH_3)(OC_3H_7)}]^+$. $\mu_{eff} = 4.82$ B.M. Anal. calc. for C, 50.46; H, 4.76; N, 3.68, found: C, 49.98; H, 4.91; N, 3.78.

2.3.2 Synthesis of $[Ni{S_2P(1,4-C_6H_4OMe)(OCH_2CH_2-CH_3)}_{2}{phen}]$ (2). To a stirring methanol solution (30 mL) of L2 (559 mg, 2 mmol) was added a solution of NiCl₂·6H₂O (238 mg, 1 mmol) in ethanol in a dropwise manner. To the resulting violet solution was added 1,10-phenanthroline (198 mg, 1

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mmol) with constant stirring resulting in a brown coloured precipitate. The precipitate was filtered *in vacuo* and washed with methanol and water (50 : 50)% to afford a light green powder. Single crystals suitable for X-ray diffraction were obtained as light green blocks from vapour diffusion of hexane into a concentrated DCM solution. Yield: 568 mg (75%). Melting point: 147–149 °C. $\mu_{\rm eff} = 2.83$ B.M. Anal. calc. for C₃₂H₃₆N₂-NiO₄P₂S₄: C, 50.47; H, 4.77; N, 3.68, found: C, 50.34; H, 4.95; N, 3.54.

2.3.3 Synthesis of [Zn{S₂P(1,4-C₆H₄OMe)(OCH₂CH₂CH₃)} {phen}] (3). An ethanolic solution of $ZnNO_3 \cdot 6H_2O$ (297 mg, 1 mmol) was added to a stirring solution of L2 (559 mg, 2 mmol) in ethanol. The solution was stirred for 20 min and 1,10-phenanthroline monohydrate (198 mg, 1 mmol) was added to the mixture and further allowed to stir for 1 hour, the resulting white precipitated was filtered and washed with ethanol/water. Recrystallization from hexane layered over DCM vielded colourless block crystals in two weeks. Yield: 607 mg (79%). Melting point 211 °C (dec.). ¹³C NMR (151 MHz, chloroform-*d*) δ 161.50, 149.75, 140.79, 139.08, 131.92, 131.82, 128.85, 126.83, 125.26, 113.06, 112.96, 66.80, 55.26, 23.61, 23.55, 10.30. ³¹P NMR (243 MHz, chloroformd) δ 103.50, 100.21. ESI-MS(m/z) (cal.) 505.0329 (505.0152) for ([M- $\{S_2P(4-C_6H_4OMe)(OCH_2CH_2CH_3)\}]^+$, 100%). Anal. calc. for C_{32} -H₃₆N₂O₄P₂S₄Zn: C, 50.03; H, 4.72; N, 3.65, found: C, 49.92; H, 4.80; N, 3.78 IR(ATR, cm⁻¹) 2970(s), 2939(s), 2882(s), 1623(s), 1594(s), 1570(s), 1516(s), 1499(s), 1461(s), 1439(s), 1423(s), 1373(s), 1321(s), 1294(s), 1183(s).

2.3.4 Synthesis of $[Cd{S_2P(1,4-C_6H_4OMe)(OCH_2CH_2 CH_3$ }{phen}] (4). In a beaker, a $Cd(NO_3)_2 \cdot 4H_2O$ (308 mg, 1 mmol) methanol solution (20 mL) was added dropwise to a stirring solution of 1,10-phenanthroline (198 mg, 1 mmol) and L2 (559 mg, 2 mmol) in 40 mL methanol. A white precipitate immediately formed and was further stirred for 1 hour at room temperature. The precipitate was collected by filtration and washed with MeOH : H_2O (50 : 50) solvent. The resulting colourless solid was dried in air. Slow evaporation from acetone yielded colourless crystals suitable for X-ray diffraction measurement. Yield: 717 mg (88%). Melting point: 256 °C (dec.). ¹H NMR (400 MHz, chloroform-*d*) δ 9.52 (s, 2H), 8.37 (dd, J = 8.2, 1.6 Hz, 2H), 7.90–7.80 (m, 6H), 7.76 (dd, J = 8.2, 4.7 Hz, 2H), 6.76 (dd, J = 8.8, 2.9 Hz, 4H), 4.13 (dt, J = 9.1, 6.7 Hz, 4H), 3.72 (s, 6H), 1.66 (dt, J = 14.2, 7.1 Hz, 4H), 0.86 (t, J = 7.4 Hz, 6H). ¹³C NMR (101 MHz, chloroform-d) & 161.51, 149.74, 138.26, 131.84, 131.70, 129.09, 126.89, 124.68, 113.29, 113.13, 66.81, 55.31, 23.71, 23.63, 10.40 anal. calc. for C₃₂H₃₆CdN₂O₄P₂S₄: C, 47.15; H, 4.45; N, 3.44; S, 15.73, found: C, 47.01; H, 4.56; N, 3.30; S, 15.13.

2.4 X-ray crystallography

Crystal evaluation and data collection of **1–4** were done on a Bruker Smart APEXII diffractometer equipped with an Oxford Cryostream low-temperature apparatus operating at 100 K, with Mo-K α radiation (I = 0.71073 Å). Reflections were collected at different starting angles and the APEXII program suite was used to index the reflections.³³ Data reduction was performed using SAINT³⁴ software, and the scaling and absorption corrections were applied using the SADABS³⁵ multi-scan technique. The structure was solved by direct methods using the SHELXS program and refined using SHELXL program.³⁶ Graphics of the crystal structures were drawn using Mercury software.³⁷ Nonhydrogen atoms were first refined isotropically and then by anisotropic refinement with the full-matrix least square method based on F^2 using SHELXL. All hydrogen atoms were positioned geometrically, allowed to ride on their parent atoms and refined isotropically.

2.5 Electrochemical measurements

To explore the effect of the synthesized complexes on the corrosion of mild steel in 1 M HCl, impedance and polarization tests were conducted in the absence and presence of the synthesized complexes. The mild steel sheets purchased from commercial supplier with a chemical composition of (in wt%): Fe (99.621), C(0.076), P (0.012), Cr (0.050), Al (0.023), Si (0.026), Mn (0.192) were cut into 1 cm \times 1 cm dimension and embedded in epoxy resin leaving an exposed area of 1 cm². Mechanical abrasion of the exposed surface was done with the aid of a Struers LaboPol-1 machine to eliminate traces of epoxy resin. Thereafter different grade emery papers ranging from 600 to 1200 were used to polish the exposed surface to eliminate surface rusts and scales. The steel surface was thereafter washed thoroughly with distilled water, degreased with acetone and dried at room temperature. The aggressive electrolytic solution of 1 M HCl was prepared by dilution of 32% HCl. The complexes were first dissolved in a small volume of DMSO (10% of total solvent volume) to ensure uniform dissolution without turbidity. Stock solutions of the complexes were subsequently prepared in 1 M HCl. Preliminary checks on the performance of the complexes as corrosion inhibitors suggested that a concentration of 100 ppm is optimum and the inhibition efficiencies were thus tested at 100 ppm.

Electrochemical studies were conducted on the AUTOLAB (model 302N) potentiostat/galvanostat coupled with a threeelectrode assemblage driven by Nova 1.10.1.9 software. A typical electrochemical system setup has the polished mild steel as the working electrode, Ag/AgCl, 3 M KCl as the reference electrode and a platinum rod as the counter electrode which are all submerged in the electrolyte. A waiting time of 1800 s was allowed for the mild steel in the electrolyte to corrode freely without applying external potential or current in order to obtain a stable open circuit potential (OCP). After the steady-state potential was achieved, EIS tests were conducted at the OCP by evaluating the frequency response of the system in the range of 100 kHz to 0.01 Hz with an amplitude of 10 mV. These EIS measurement settings are considered to be wide enough to sufficiently reveal the resistive and capacitive behaviour of the electrochemical system, and also comply with the recommended ASTM practice G106-89.38 All electrochemical tests conducted were done under aerated unstirred conditions at 303 \pm 1 K. Values of charge transfer resistance were generated from the obtained impedance spectra. Percentage inhibition efficiencies (η_{EIS}) of the studied compounds were calculated from the measured charge transfer resistance (R_{ct}) values using the following equation:39

$$\eta_{\mathrm{EIS}} = rac{R_{\mathrm{ct}} - R_{\mathrm{ct}}^0}{R_{\mathrm{ct}}} imes 100$$

where R_{ct} and R_{ct}^{0} represent the values of the charge transfer parameters with and without the inhibitors, respectively.

Polarization measurements were performed by sweeping the potential from -250 to +250 mV with respect to the OCP at a constant scan rate of 1 mV s⁻¹. The measurement parameters are chosen within the recommended limits in the ASTM F 2129 standard practice.⁴⁰ Corrosion current densities (i_{corr}) were obtained by extrapolating the linear segments of the anodic and cathodic Tafel slopes to corrosion potential. The percentage inhibition efficiency (η_{PDP}) was calculated from the obtained values of i_{corr} using the equation:⁴¹

$$\eta_{ ext{PDP}} = rac{i_{ ext{corr}}^0 - i_{ ext{corr}}}{i_{ ext{corr}}^0} imes 100$$

where i_{corr} and i_{corr}^{0} represent the corrosion current densities with and without the studied inhibitors, respectively.

In order to measure the linear polarization resistance, a potential range of ± 10 mV relative to OCP was applied to the system and current response was monitored at a scan rate of 0.125 mV s⁻¹. Values of polarization resistance (R_p) for the blank and inhibited system were obtained from this method and the percentage inhibition efficiency was calculated thus:⁴²

$$\eta_{ ext{LPR}} = rac{R_{ ext{p}} \ - R_{ ext{p}}^0}{R_{ ext{p}}} imes 100$$

where R_p and R_p^0 represent the polarization resistances in the presence and absence of the studied inhibitors, respectively.

2.6 Computational studies

The quantum chemical calculations for the inhibitors were performed in the gas phase using density functional theory



Scheme 1 Summary of the synthesis methodology for the preparation of complexes 1–4.

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Table 1 Crystallographic data and structure refinement for 1a, 1b, 2, 3 and 4

Compound	2	1h	6	3	
Compound Emnirical formula				STON H J	S d O NPJ H J
	C321136CUIV2C4F 254	U321136UU1V2U4F204	U3211361V21VIU4F204	C3211361V2O4F204L11	C321136CULV2C4F204
Formula weight	/61./4	/61./4	/01.52	/68.18	815.21
Temperature/K	100(2)	150(2)	100(2)	100(2)	100
Crystal system	Monoclinic	Triclinic	Triclinic	Triclinic	Monoclinic
Space group	C2/c	$P\overline{1}$	$P\bar{1}$	$P\overline{1}$	C2/c
a/Å	22.9770(4)	10.8831(2)	10.8316(6)	10.02980(10)	49.6546(10)
$b/ m \AA$	10.7463(2)	12.4185(2)	13.2433(8)	12.6887(2)	11.0919(2)
$c/{ m \AA}$	15.6178(3)	14.1718(3)	14.1413(9)	14.1918(2)	34.7070(7)
$\alpha / ^{\circ}$	90	91.5350(10)	65.148(3)	87.0380(10)	06
$\beta/^{\circ}$	113.0040(10)	97.5390(10)	89.961(3)	78.8780(10)	112.9760(10)
γ°	90	109.8070(10)	71.017(3)	73.5350(10)	06
Volume/Å ³	3549.65(12)	1781.23(6)	1718.55(18)	1699.52(4)	17598.9(6)
Z	4	2	2	2	20
$ ho_{ m calc}~{ m g}~{ m cm}^{-3}$	1.425	1.420	1.472	1.501	1.538
μ/mm^{-1}	0.847	0.844	0.940	1.102	0.987
F(000)	1580.0	790.0	792.0	796.0	8320.0
Crystal size/mm ³	$0.41 \times 0.32 \times 0.23$	$0.33 \times 0.22 \times 0.12$	$0.24 \times 0.21 \times 0.16$	$0.42\times0.22\times0.191$	0.3 imes 0.24 imes 0.21
Radiation	$\mathrm{MoK}lpha\left(\lambda=0.71073 ight)$	$\mathrm{MoK}\alpha~(\lambda=0.71073)$	$\mathrm{MoK}\alpha~(\lambda=0.71073)$	$\mathrm{MoK}\alpha~(\lambda=0.71073)$	$\mathrm{MoK}\alpha~(\lambda=0.71073)$
20 range for data collection/ $^{\circ}$	3.852 to 57.042	3.496 to 54.392	3.214 to 52.448	3.348 to 54.81	1.782 to 56.772
Index ranges	$-30 \le h \le 27$,	$-13 \leq h \leq 13,$	$-13 \leq h \leq 13,$		$-66 \leq h \leq 65,$
	$-14 \leq k \leq 14, -20 \leq l \leq 20$	$-15 \leq k \leq 15, -18 \leq l \leq 18$	$-16 \leq k \leq 16, -16 \leq l \leq 17$		$-14 \le k \le 14, -46 \le l \le 46$
Reflections collected	23 063	20 669	23 029	20 706	139572
Independent reflections	$4403 \; [R_{ m int} = 0.0168,$	$7650 \; [R_{ m int} = 0.0182,$	$6684 \left[R_{ m int} = 0.0234, ight.$	$7546 [R_{ m int} = 0.0117,$	$21 \ 937 \ [R_{ m int} = 0.0337,$
	$R_{ m sigma}=0.0131]$	$R_{ m sigma}=0.0227]$	$R_{ m sigma}=0.0245]$	$R_{ m sigma}=0.0137]$	$R_{ m sigma}=0.0247]$
Data/restraints/parameters	4403/0/207	7650/1/412	6684/0/410	7546/0/431	21 937/0/1024
Goodness-of-fit on F^2	1.050	1.040	1.112	1.048	1.188
Final <i>R</i> indexes $[I \ge 2\sigma (I)]$	$R_1=0.0226, \mathrm{w} R_2=0.0614$	$R_1=0.0266, \mathrm{w} R_2=0.0621$	$R_1=0.0562, \mathrm{w} R_2=0.1379$	$R_1 = 0.0297, \mathrm{w} R_2 = 0.0800$	$R_1=0.0417, \mathrm{w}R_2=0.0891$
Final R indexes [all data]	$R_1 = 0.0247, wR_2 = 0.0629$	$R_1 = 0.0344, wR_2 = 0.0664$	$R_1 = 0.0611, WR_2 = 0.1413$	$R_1=0.0323,\mathrm{w}R_2=0.0820$	$R_1 = 0.0525, wR_2 = 0.0930$
Largest diff. peak/hole/e A	0.51/-0.23	$0.34/{-}0.27$	$1.55/{-0.85}$		$1.80/{-1.44}$

(DFT). The density functional theory (DFT) calculations were executed applying the hybrid BP86 exchange-correlation functional with the Def2TZVP43 basis set for geometry optimization of the structures using Gaussian'16 program package.44 The synergy between the BP86 functional and the Def2TZVP basis set have been reported.43,45 However, for the nickel complex, B3LYP⁴⁶ functional with LANL2DZ was utilized and have been previously reported for Ni complexes.47 Optimizations to a local minimum for all structures using the Berny algorithm⁴⁸ was confirmed with no negative imaginary frequencies. This indicates that the structure geometry was obtained with a stationary point on the potential surface. The highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO) energies were obtained for all the metal complexes. The difference between the energy of the LUMO and HOMO results in the energy gap (ΔE). Other reactivity descriptors were derived such as the ionization potential (I) and electron affinity (A), absolute hardness (η) , softness (S), electrophilicity index (ω) and absolute electronegativity (χ) of the complexes based on Koopman's theorem.⁴⁹ These descriptors are derived as follows:



3. Results and discussion

3.1 Synthesis of complexes

The methodology for the synthesis of complexes **1–4** are summarized in Scheme 1. The molecular structure for any homoleptic cobalt(π) dithiophosphonato complex has to date eluded isolation and the formation of such complexes are thus performed by the introduction of N-donor ligands such as 1,10phenanthroline and 2,2-bipyridine. The cobalt(π) complex **1** was isolated from a mixture of CoCl₂·6H₂O precursor, ammonium dithiophosphonate ligand (**L1**) and 1,10-phenanthroline in 1 : 2 : 1 molar ratio. Complex **1** is soluble in chlorinated solvents, acetone, and THF but insoluble in *n*-hexane and water.

The nickel(II) complex 2 was synthesized from the reaction between NiCl₂·6H₂O, 1,10-phenanthroline and L2, in a stoichiometric ratio of 1:1:2. The complex was isolated in high yield as an air and moisture stable powder. Complex 2 was soluble in DCM, chloroform, acetone and THF but insoluble in alcohols, water, ether and *n*-hexane. The zinc(II) complex 3 was synthesized in high yield as air and moisture stable white solid by the reaction between $Zn(NO_3)_2 \cdot 6H_2O_1$, 1,10-phenanthroline and L2 in a stoichiometric ratio of 1 : 1 : 2 at room temperature. The complex is soluble in DCM, acetone, chloroform and THF but insoluble in alcohols, ether, n-hexane and water. The low molar conductance values of 10^{-3} M solutions of the complexes in DMSO at room temperature (8.94–22.60 Ω^{-1} cm² mol⁻¹) indicate that the complexes are non-electrolytes.⁵⁰ Complex 3 is diamagnetic and the ¹H, ¹³C and ³¹P NMR data were generated from CDCl₃ solution, whilst FT-IR data was obtained in the solid-state. A zinc(II) complex of dithiophosphate with 4,4bipyridine was reported by Glinskaya et al., adopting a distorted



Fig. 1 Molecular structure for one of the polymorphs of 1 with thermal ellipsoids drawn at 50% probability. Hydrogen atoms omitted for clarity.

Table 2Selected bond lengths (Å) and angles (°) for 1a and 1b, 2 withthe estimated standard deviation (e.s.d.) in parentheses

Complex	1a	1b	Complex	2
Co1-S1	2.5063(3)	2.5496(5)	Ni1-S1	2.3217(5)
Co1-S2	2.5061(3)	2.4574(5)	Ni1-S2	2.3034(5)
Co1-N1	2.1526(10)	2.1486(13)	Ni1-N1	2.1247(16)
Co1-N2	2.1526(9)	2.1473(14)	Ni1-N2	2.0588(15)
S1-P1	1.9992(4)	1.9931(6)	S1-P1	2.0309(8)
S2-P2	1.9902(4)	1.9898(6)	S2-P2	2.0287(6)
P1-O1	1.6027(9)	1.6006(12)	P1-O1	1.6013(16)
P2-C13	1.7928(12)	1.8038(17)	P2-O2	1.6125(14)
O1-C1	1.4580(14)	1.451(2)	P2-C13	1.800(2)
N3-C11	1.3602(14)	1.362(2)	O1-C3	1.443(3)
S1-Co1-S2	96 - 542(14)	99.263(16)	S1-Ni1-S2	81.540(15)
N1-Co-S1	94.63(3)	93.15(4)	N1-Ni-S1	93.33(4)
N1-Co-N2	76.73(5)	76.81(5)	N1-Ni-N2	80.18(5)

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tetrahedron geometry,⁵¹ but the present study reports a mononuclear zinc(II) complex for a dithiophosphonate ligand with a 4-coordinate geometry for the first time. The cadmium(II) complex, 4 was isolated as colourless solid from a mixture of $Cd(NO_3)_2 \cdot 4H_2O$ salt, ligand L2 and 1,10-phenanthroline in alcohol in a 1:2:1 stoichiometric ratio. The complex is soluble in DCM, chloroform and acetone but insoluble in water, *n*hexane and alcohols.

3.2 NMR spectroscopy

The NMR spectra for complexes **1** and **2** were not recorded due to their paramagnetic nature. Magnetic susceptibility measurements gave magnetic moments of 4.82 and 2.83 B.M for **1** and **2** respectively, indicative of a high spin complexes with four unpaired electrons for **1** and two unpaired electrons for **2**, both of which are paramagnetic with octahedral geometries. The NMR data for complex 3 are presented in the experimental section. The methoxy protons for 3 were observed at 3.75 ppm as singlet peaks and the terminal methyl peak in the (OR) fragment was observed at 0.79 ppm. The other propoxyl protons in 3 were observed at 3.64 ppm for OCH₂– and 1.46 ppm for $-OCH_2CH_2CH_3$, with the aromatic protons, observed 9.30–6.56 ppm. The ¹H NMR spectrum for 4 showed peaks associated with 1,10-phenanthroline in the range of 7.83–9.48 ppm; while the singlet peak at 3.72 ppm is diagnostic of OCH₃ of the anisyl fragment of Lawesson's reagent.

3.3 X-ray crystallography

The X-ray crystal structure of **1** consisted of discrete mononuclear molecular units. The crystallographic and structure



Fig. 2 Molecular representation of 2 (Thermal ellipsoids are drawn at 50% and hydrogen atoms omitted for clarity) (above) and crystal packing diagram of 2 viewed along *c*-axis (below).

refinement data are presented in Table 1. The complex is neutral and the Co center adopts a distorted octahedral geometry. Two structures of 1 were obtained (1a and 1b) as polymorphs, 1a crystallized in a monoclinic C2/c space group and 1b crystallized in a triclinic $P\bar{1}$ space group. It is a neutral 6-coordinate complex whose coordination sites include the four sulphur donor atoms of the two bidentate dithiophosphonate ligands, and two nitrogen donor atoms of 1,10-phenanthroline. The bond angles for S-Co-S average at 86° and N-Co-N average at 76°. The bond angles are smaller than 90°, typically expected for ideal octahedral complexes due to restricted intraligand bite. The bond distances for Co-S, Co-N and P-S range from 2.457(5)-2.5496(5) Å, 2.1473(14)-2.1526(9) Å and 1.9898(6)-1.9992(4) Å, respectively. The intraligand S…S bite average at 3.326 Å. The crystal structure for 1 is shown in Fig. 1 and selected bond lengths (Å) and angles (°) for 1a and 1b are shown in Table 2.

The Ni(II) complex, 2 crystallized in a triclinic space group $P\bar{1}$ and its crystallographic data are listed in Table 1. The complex is neutral, mononuclear and 6-coordinate with a distorted octahedral geometry around the Ni(II) center stabilized by four S donor atoms from two bidentate dithiophosphonate ligands and two N donor atoms from the 1,10-phenanthroline, shown in Fig. 2. The Ni-S distances are in the range 2.3034(4) - 2.3451(4) Å, which are in good agreement with previous reports.⁵² The Ni-N bond distances range from 2.1336(13) - 2.0588(15) Å and the P-S bonds average at 2.028 Å. The S-Ni-S bond angles average at 87° and the N-Ni-N bond angles average at 79°, which are smaller than 90° expected for a regular octahedron due to restricted ligand bite.52 Selected bond distances and angles are listed in Table 2.

The zinc(II) complex 3 crystallized in the triclinic space group $P\bar{1}$ and the molecular structure is shown in Fig. 3 and the crystallographic data are presented in Table 1. The crystal structure analysis revealed that 3 was a discrete mononuclear neutral species. The geometry around the zinc(II) center adopts

Table 3 Selected bond lengths (Å) and angles (°) for 3 and 4 with the estimated standard deviation (e.s.d.) in parentheses

Complex	3	Complex	4
Zn1-S1	2.3217(5)	Cd1-S1	2.6775(7)
Zn1-S2	2.3034(5)	Cd1-S2	2.6915(7)
Zn1-N1	2.1247(16)	Cd1-N1	2.374(2)
Zn1-N2	2.0588(15)	Cd1-N2	2.386(2)
S1-P1	2.0309(8)	S1-P1	1.9999(10)
S2-P2	2.0287(6)	S2-P2	1.9945(10)
S3-P1	1.9564(9)	S3-P1	2.0035(10)
S4-P2	1.9414(7)	S4-P2	1.9894(10)
P1-O1	1.6013(16)	P1-O1	1.606(2)
P2-O2	1.6125(14)	P2-O2	1.603(2)
P2-C13	1.800(2)	P2-C13	1.795(3)
O1-C3	1.443(3)	O1-C3	1.422(4)
S1-Zn1-S3	118.655(18)	S1-Cd1-S2	89.41(2)
N1-Zn-S1	114.73(4)	N1-Cd-S1	95.22(6)
N1-Zn-N2	80.26(6)	N1-Cd1-N2	70.13(8)

a distorted tetrahedral configuration, coordinating through one S-donor atom from each of the two dithiophosphonate ligands in a monodentate fashion and two N-donor atoms of one 1,10phenanthroline (bidentate) ligand in a bidentate chelating fashion. The Zn-S and Zn-N bond distances range from 2.2837(9) -2.3451(4) Å and 2.052(3) to 2.1336(3), respectively, which is within the range previously reported for a $zinc(\pi)$ dithiophosphate and N-donor ligand (Zn-S: 2.285(8)- 2.3065(7) Å and Zn–N: 2.047(2) Å).⁵¹ The P–S bond distance for the coordinated sulphur is in the range of 2.0274(6)-2.0396(13) Å and expectedly longer than the uncoordinated sulphur, which is in the range 1.9613(7)-1.9457(11) Å. The bond angles N1–Zn1–S1 and S1-Zn-S2 of the distorted tetrahedron range from 79.80(5)-120.162(15)° and is in good agreement with related reported values.53 Selected bond distances and angles are listed in Table 3.



Fig. 3 Molecular structure of 3 (thermal ellipsoids are drawn at 50% and hydrogen atoms omitted for clarity).



Fig. 4 Molecular structure representation of 4. Thermal ellipsoids drawn at 50% with hydrogen atoms omitted for clarity.

Slow evaporation from acetone yielded single crystals of 4 suitable for crystallographic studies. Complex 4 crystallizes in the monoclinic space group C2/c; the unit cell consists of 2.5 symmetrically non-equivalent molecules, shown in Fig. 4. The structure of 4 revealed the first mononuclear cadmium(II) complex with this set of N and S donor ligands. The Cd(II) center adopts a distorted octahedral geometry formed by two dithiophosphonate ligands coordinating through four sulphur atoms and two N atoms of the 1,10-phenanthroline. The Cd-S distances average at 2.626 Å, which is in agreement with reported values for similar bonds reported.54 The Cd-N bond distances average at 2.380 Å and the P-S bonds average at 1.9998 Å. The N-ligand and S-ligands are bidentate and form five and four-membered CdN₂C₂ and CdS₂P chelate rings, respectively. The S-Cd-S bond angles average at 80° and the N-Cd-N bond angles average at 70°. These angles are smaller than 90° expected for a regular octahedron due to restricted ligand bite.⁵⁴ Selected bond distances and angles are listed in Table 3.

3.4 Electrochemical studies

The molecular structures of the complexes **1–4** are summarized in Scheme 1 and for the purpose of electrochemical corrosion inhibition studies, complexes **1**, **2**, **3** and **4** are referred to as Co, Ni, Zn, and Cd, respectively.

3.4.1 Open-circuit potential (OCP) scanning. The OCP-time profile for mild steel corrosion in 1 M HCl solution in the absence and presence of 100 ppm of the studied complexes is shown in Fig. 5. The OCP *versus* time plot provides information on the stability of the electrochemical system before applying external potential or current. The obtained profile revealed that



Fig. 5 OCP-time curves of mild steel in 1 M HCl without inhibitor (blank) and with 100 ppm of studied complexes at 303 K.

the system achieved a steady-state potential before the first 600 s after submerging the working electrode in the corrosive medium. The observed slight initial fluctuation of OCP can be attributed to the delayed dissolution of iron oxide layers (Fe₂O₃ and Fe₃O₄) that pose as the native protective oxide layers for the steel, and formation of defensive films by the studied complexes in the electrolytic medium.⁵⁵ At complete dissolution of the oxide layers and adsorption of studied molecules on the steel surface, the steady-state potential becomes relatively stable. The OCPs of the systems are generally stable with differentials not more than 3 mV over the entire 30 min waiting period, suggesting that the systems had assumed a sufficiently stable OCP before electrochemical perturbation.

3.4.2 Potentiodynamic polarization and linear polarization resistance (PDP and LPR). The Tafel plots obtained from the potentiodynamic polarization measurements on the working electrode in 1 M HCl in the presence and absence of 100 ppm of the studied complexes at 303 K is presented in Fig. 6. The values of electrochemical parameters such as anodic and cathodic



Fig. 6 Potentiodynamic polarization curves of mild steel in 1 M HCl without inhibitor (blank) and with 100 ppm of studied complexes at 303 K.

Table 4	Polarization parameters for mild stee	in 1 M HCl without inhibitor	[.] (blank) and with 100	opm of studied complexes at 303 K
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	PDP					LPR	
Inhibitors	$-E_{\rm corr}$ (mV)	$i_{ m corr}$ (µA cm ⁻²)	$\beta_{\rm a} \left({\rm mV \ dec^{-1}} \right)$	$\beta_{\rm c} ({\rm mV} {\rm dec}^{-1})$	$\eta_{ m PDP}$	$R_{\rm PR} \left(\Omega \ {\rm cm}^2\right)$	$\eta_{ m LPR}$
Blank	417.74	560.49	138.33	90.09	_	42.28	
Со	422.26	153.99	112.89	83.60	72.53	135.45	68.79
Zn	438.17	49.64	102.89	107.97	91.14	460.70	90.82
Cd	420.31	43.27	91.04	96.66	92.28	470.53	91.01
Ni	418.00	32.85	99.58	90.92	94.14	628.24	93.27

Tafel slopes (β_a and β_c), corrosion current density (i_{corr}), corrosion potential (E_{corr}) and inhibition performance (η_{PDP}) are reported in Table 4. It is clear from the anodic and cathodic branches of the plots that the introduction of Co resulted in a shift to a region of lower current density as compared to the blank solution. Similar shifts were observed for the other three complexes, suggesting that all the inhibitors show similar corrosion inhibition mechanism with Ni showing the most prominent shift from the blank. This observation suggests that the four studied complexes impede steel corrosion by forming an inhibitive film at the active sites on the metallic surface.^{41,56}

The mode of corrosion inhibition exhibited by Co, Zn, Cd and Ni can be classified as anodic, cathodic or mixed-type based on the magnitude of displacement of their E_{corr} compared to that of the blank. It is generally reported that a shift in E_{corr} value less than 85 mV describes a mixed-type inhibitor while

a shift in $E_{\rm corr}$ greater than 85 mV is categorized as cathodic or anodic.57,58 In the present study, the maximum displacement in $E_{\rm corr}$ ranged from 0.26 to 20.43 mV, which proposes that the studied compounds are mixed-type corrosion inhibitors. In other words, the studied inhibitors exhibited the ability to simultaneously restrain anodic dissolution of mild steel and cathodic evolution of hydrogen gas. The obtained values of the Tafel slopes showed a small deviation on the addition of the inhibitors as compared to the blank solution, which suggests that the adsorption mechanism is unaffected by the addition of the inhibitors. The studied inhibitors simply adsorb on the metal surface and block the active sites.50 This was further substantiated by the changes in the values of the anodic and cathodic slopes for the inhibited systems. Factors under consideration when discussing the efficiency of corrosion inhibitors include the presence of heteroatoms, molecular size,



Fig. 7 (a-c). Nyquist plot (A), Bode plot (B) and Randle equivalent circuit (C) for mild steel in 1 M HCl without inhibitor (blank) and with 100 ppm of complexes studied at 303 K.

Inhibitor	$R_{\rm s} \left(\Omega \ {\rm cm}^2 \right)$	$R_{\rm ct} \left(\Omega \ {\rm cm}^2\right)$	n	$Y_0 \left(\mu\Omega \ \mathrm{s}^n \ \mathrm{cm}^{-2}\right)$	$-\alpha$	-S	$C_{\rm dl} \left(\mu {\rm F} \ {\rm cm}^{-2} \right)$	χ^2	η_{EIS}
Blank	2.69	12.5	0.835	613.0	34.86	0.41	234.10	0.18807	_
Со	2.37	50.4	0.835	361.0	53.16	0.56	163.56	0.24371	75.20
Zn	1.98	118.0	0.853	235.0	60.91	0.69	126.69	0.33546	89.41
Cd	2.73	133.0	0.845	151.0	62.95	0.64	73.73	0.31865	90.60
Ni	2.42	152.0	0.856	141.0	63.48	0.71	73.87	0.35772	91.78

Table 5 Electrochemical impedance spectroscopy parameters obtained for studied compounds at 303 K

molecular geometry, electron charge density, mode of interaction, number of functional groups, and so on.56-58 The large molecular size, molecular geometry and the presence of donor atoms such as N, O, S and P in the studied complexes might contribute to their excellent inhibition properties. The order of inhibition efficiencies is Ni (94.14%) > Cd (92.28%) > Zn (91.14%) > Co (72.53%). The observed higher inhibition efficiency for Ni follows the trend previously reported.59,60 Furthermore, electrochemical parameters obtained from linear polarization measurements revealed that the studied complexes behaved as excellent corrosion inhibitors for mild steel in the studied acidic medium. The values of polarization resistance $(R_{\rm p})$ which are obtained from the slope of current densitypotential graph for each studied complex ranged from 135.45 to 628.24 $\Omega~\text{cm}^2$ and were greater than the 42.28 $\Omega~\text{cm}^2$ of the uninhibited solution. This suggests that the adsorbed portions of the complex on the steel surface have successfully blocked the active sites and inhibited dissolution of the metal.

3.4.3 Electrochemical impedance spectroscopy (EIS). EIS was employed as a reliable method for gaining better insight into the behaviour of mild steel-inhibitor interfaces. The Nyquist and Bode plots for steel corrosion in the corrosive medium in the absence and presence of 100 ppm Co, Zn, Cd and Ni at 303 K are shown in Fig. 7a and b. The Nyquist plots of the uninhibited and additive-containing media (with the studied complexes) revealed depressed semicircles indicating the mechanism of steel corrosion was the same in all cases. Careful interpretation of the impedance spectra showed that the introduction of 100 ppm of different complexes resulted in a significant increase in the size of the semicircles and the charge transfer resistance. This indicated that the studied complexes adsorbed on the surface of the metal specimen, thereby forming a protective layer on the mild surface.^{39,61} A simple Randle circuit (equivalent circuit) comprising a constant phase element (CPE), solution resistance (R_s) and charge transfer resistance $(R_{\rm ct})$ shown in Fig. 7c employed to fit the impedance spectra showed excellent fitting with the goodness of fit values ranging from 0.18807 to 0.35772. The EIS parameters derived from the spectra are presented in Table 5.

The tabulated results (Table 5) showed an increase in the values of $R_{\rm ct}$ from 12.5 Ω cm² (for the blank solution) to values ranging from 50.4 to 152.0 Ω cm² with the introduction of 100 ppm of Co, Ni, Cd and Zn complexes, which correspond to the ability of the complexes to impede charge transfer across steel/electrolyte interface. An inhibition efficiency of 91.78% was recorded at 100 ppm of Ni complex at 303 K. Additionally, a drop was observed in the values of $C_{\rm dl}$ with the introduction of

the complexes. The significant increase in the $R_{\rm ct}$ values is due to the formation of a shielding layer on the mild steel surface by the inhibitor molecules.⁵⁸ The observed decrease in $C_{\rm dl}$ can be correlated to the thickness of the protective layer on the steel/acid interface, which is attributed to gradual replacement of water molecules by the inhibitor molecules.⁶² The double-layer capacitance ($C_{\rm dl}$) for mild steel corrosion in the acid and in the presence of the inhibitors was calculated using the equation:⁵⁰

$$C_{\rm dl} = (Y_{\rm o} R_{\rm ct}^{1-n})^{1/n}$$

where Y_0 is the CPE constant, R_{ct} represents the charge-transfer resistance and n (a CPE exponent) is the phase shift which is often used to categorize the nature of the CPE, and also as a measure of the degree of roughness of the electrode surface.⁵⁰ A unit value of n characterizes the CPE to be purely capacitive, while near unity values have been associated with pseudocapacitive electrode surface. The values of n obtained from the EIS as listed in Table 5 are near unity, suggesting the pseudocapacitive behaviour of the electrode surface. Moreover, the values of n in the presence of inhibitors are slightly higher than that for the inhibitor-free medium (except for Co), suggesting that the steel surface assumes less inhomogeneity in the inhibitorcontaining solutions, due to uniform coverage of the electrode surface by the adsorbed inhibitor molecules.⁵⁰

Furthermore, the Bode diagram (Fig. 7b) for the steel corrosion in the presence and absence of the studied complexes features a one-phase peak which confirms that the corrosion process exhibited a single time-constant and is mainly controlled by a single charge transfer mechanism.⁶¹

The values of the phase angles (α) and slopes (S) obtained from the linear portion of the Bode plot at the middle frequency range are recorded in Table 5. The observed trend of increase in the value of the phase angle on introducing the studied complexes to the blank solution indicate the formation of a protective film on the mild steel surface. In addition, the obtained $-\alpha$ and -S which ranged from 53.16–63.48 and 0.56– 0.71 respectively for the studied complexes suggest that the electrode/electrolyte interface acts a pseudocapacitor where an ideal capacitor has a α of -90 and S of $-1.^{63.64}$ The computed inhibition potentials from the different electrochemical methods employed in the study showed excellent agreement.

3.5 Computational study

3.5.1 Quantum chemical descriptor calculations. Quantum chemical descriptors of the four metal complexes (Co, Ni, Zn, and Cd) were investigated to correlate their electronic structures

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Fig. 8 The optimized structures and HOMO and LUMO electron density isosurfaces of Co, Zn, Cd and Ni complexes at BP86/Def2TZVP and B3LYP/LANL2DZ levels.

to their reactivities and viz-a-viz performances as corrosion inhibitors.⁶⁵ The structures of the metal complexes were optimized using BP86 method with Def2TZVP for Zn, Co and Cd complexes while B3LYP with LANL2DZ basis set for Ni. The optimized structures, highest occupied molecular orbitals (HOMO) and lowest unoccupied molecular orbitals (LUMO) electron density isosurfaces for the studied metal complexes are shown in Fig. 8.

The HOMO and LUMO electron density isosurfaces of all the metal complexes showed similar delocalization patterns. The

 Table 6
 The calculated quantum chemical parameters for the metal complexes using BP86/Def2TZVP and B3LYP/LANL2DZ levels

Parameters	Со	Zn	Cd	Ni
Еномо	-4.099	-4.751	-4.691	-5.096
ELUMO	-2.99	-3.293	-3.156	-2.744
ΔE	1.109	1.458	1.535	2.352
η (eV)	0.5545	0.729	0.7675	0.4252
$S(eV^{-1})$	0.9017	0.6859	0.6515	0.4255
I (eV)	4.099	4.751	4.691	5.096
A (eV)	2.99	3.293	3.156	2.774
χ (eV)	-3.6275	-4.022	-3.9235	-3.92
ω (eV)	11.8654	11.095	10.0285	6.533
μ (debye)	4.5615	8.2503	5.4076	5.9329

HOMOs were localized on the central metal atom coordinated with highly electronegative atoms such as S, O and P in all cases while the LUMOs are delocalized on the 1,10-phenanthroline aromatic rings. Similar observations have been previously reported.⁶⁶

The obtained quantum chemical parameters such as the $E_{\text{HOMO}}, E_{\text{LUMO}}, \Delta E, \eta, S, A, I, \chi, \mu$ and ω for the studied metal complexes are shown in Table 6. The energy of the HOMOs (E_{HOMO}) was higher indicating its ability to release electrons from the metal to be accepted by the LUMOs having lower energy (E_{LUMO}) . Co has the lowest energy gap (1.109 eV), that is, it is more reactive than Zn, Cd and Ni. The energy gap of Co corresponds to its higher value of softness showing the greater tendency to donate an electron to the metal surface. As reported in Table 6, the electrophilicity value of the Co metal complex was higher than Zn, Cd and Ni, this supported the observation that Co complex is most reactive. The theoretical and experimental results are in good agreement except for Co²⁺ which showed the least inhibition property thereby contradicting the experimental. The tendency of Co^{2+} to oxidize to Co^{3+} in the presence of HCl (a strong oxidizing agent) accounts for why the theoretical results deviates from the experimental results presented. It, therefore, implies that Co²⁺ was partially oxidized in HCl solution.

4. Conclusions

Heteroleptic complexes of four divalent metals, Co, Ni, Zn, and Cd with dithiophosphonates and N-donor ligands were successfully synthesized and fully characterized by FT-IR, magnetic moment measurements, elemental analysis and mass spectrometry. Structural elucidation of all the four complexes was achieved by single-crystal X-ray studies. The crystal structures revealed the first known Co²⁺ complexes of dithiophosphonates, as well as first mononuclear 4-coordinate zinc complexes. The metal complexes showed excellent corrosion inhibition potentials for mild steel in 1 M HCl. The complexes gave inhibition efficiencies of 75.20% (Co), 89.41% (Zn), 90.60% (Cd) and 91.78% (Ni) when applied at 100 ppm to mild steel in 1 M HCl.

Conflicts of interest

The authors declare no conflict of interest.

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