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Charged Dioxomolybdenum(VI) Complexes with Pyridoxal Thiosemicarbazone Ligands as Molybdenum(V) Precursors in Oxygen Atom Transfer Process and Epoxidation (pre)Catalysts

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Abstract

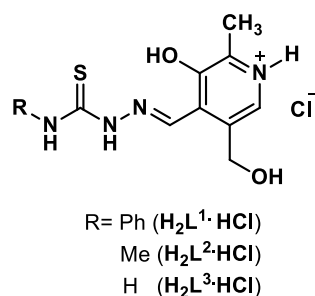
The reaction of $[\text{MoO}_2(\text{acac})_2]$ with hydrochloric salts of pyridoxal thiosemicarbazone ligands, $\text{H}_2\text{L}^{1-3}\cdot\text{HCl}$, in dry methanol or acetonitrile resulted in the formation of the mononuclear complex $[\text{MoO}_2(\text{HL}^1)(\text{CH}_3\text{OH})]\text{Cl}\cdot 1.5\text{CH}_3\text{OH}$ (**1a**) or polynuclear complexes $[\{\text{MoO}_2(\text{HL}^{1-3})\}\text{Cl}]_n$ (**1-3**) containing a different substituent on the nitrogen atom of the N-thioureide fragment: Ph (**1** and **1a**), Me (**2**) or H (**3**). New dinuclear molybdenum(V) complexes $[\text{Mo}_2\text{O}_3(\text{HL}^{1-3})_2]\text{Cl}_2$ (**4-6**) were obtained by oxygen atom abstraction from **1-3** with triphenylphosphine. All compounds were characterized by chemical analysis, IR spectroscopy, one- and two-dimensional NMR methods, TGA and in selected cases by DSC measurements. In addition, the crystal and molecular structure of **1a** was determined by single crystal X-ray diffraction. All molybdenum(VI) species were used as (pre)catalysts for olefin epoxidation under solvent-free conditions with a 0.05% Mo loading vs. olefin.

Keywords: Molybdenum/ Thiosemicarbazones / Pyridoxal / Epoxidation / Solvent-free catalysis / OAT

1. Introduction

Oxygen atom transfer (OAT) processes, as well as oxidation processes, are important in Nature and in the chemical industry [1]. Molybdenum enzymes catalyze a number of such reactions. These oxotransferases contain oxo- (or dioxo)-molybdenum units additionally coordinated by sulfur atoms [2]. We are currently interested in dioxomolybdenum(VI) complexes with ligands that mimic the coordination environment of the metal ion in enzymes. Among them, tridentate ligands containing ONS donor atoms are attracting our interest.

Transition metal complexes with thiosemicarbazone ligands are of particular importance because of their interesting biological properties, e.g. antitumoral, antifungal or antiviral properties [3]. Ligands derived from pyridoxal have attracted considerable attention due to different coordination modes to the central metal atom [4-8]. In these complexes, the pyridoxal thiosemicarbazone ligand can be in neutral (H_2L), singly-deprotonated (HL)⁻, or doubly-deprotonated (L)²⁻ form [4]. The protonation state of these ligands in metal complexes plays an important role since it offers fine-tuning of properties such as electrochemical, photophysical or catalytic activity [9]. Although numerous transition metal complexes (Cu, Fe, Ni, Mn, Co) [8a,10] with pyridoxal thiosemicarbazone derivatives are known, related molybdenum complexes are scarce [11]. Mo^{VI} complexes are also known to be very active as epoxidation catalysts, generally in solution of an organic solvent, especially for neutral complexes [12]. We have recently demonstrated that these epoxidation reactions show very good results under solvent-free conditions [13].



Scheme 1. Pyridoxal based ligands.

In this contribution, we present the syntheses and characterization of dioxomolybdenum(VI) complexes with different pyridoxal thiosemicarbazone charged ligands (Scheme 1), as well as their activity as OAT agents and as epoxidation catalysts under solvent-free conditions. The influence of protonation on the spectroscopic and structural properties of the metal complexes was studied by UV-Vis and NMR spectroscopies, as well as by a single crystal X-ray diffraction.

2. Experimental Section

2.1. Measurements.

Thermogravimetric (TG) analyses were performed using a Mettler TG 50 thermobalance with aluminum crucibles. All experiments were recorded with a heating rate of 5 °C min⁻¹ in a dynamic atmosphere with a flow rate of 200 cm³ min⁻¹. Differential scanning calorimetry (DSC) measurements were carried out with a Mettler–Toledo DSC823e calorimeter and analyzed by the Mettler STARE 9.01. software. Infrared spectra were recorded in KBr pellets with a Perkin–Elmer 502 spectrophotometer in the 4500–450 cm⁻¹ region. Electronic absorption spectra were recorded at 25 °C on a Cary 100 UV/Vis spectrophotometer. Magnetic measurements were made by the standard Gouy method. NMR spectra were obtained using a Bruker Avance DRX500 spectrometer. The spectra were recorded in DMSO-d⁶ with TMS as internal standard. ¹H and ¹³C resonances were assigned by the combination of 1D (¹H and APT sequences) and 2D homo- and heteronuclear experiments (COSY, HSQC and HMBC sequences). Elemental analyses (C, H, N, S) were provided by the Analytical Services Laboratory of the Ruđer Bošković Institute, Zagreb. Chlorine was determined by the gravimetric method as silver chloride [14].

2.2. Starting materials

The starting complex [MoO₂(acac)₂] was prepared as described in the literature [15]. Acetonitrile was dried over P₂O₅ and then distilled. Methanol was dried using magnesium turnings and iodine, and then distilled. Pyridoxal hydrochloride, thiosemicarbazides (H₂NNHC(S)NHR, R = C₆H₅, CH₃ or H) and triphenylphosphine (PPh₃) were commercially available. Ligands H₂L¹⁻³·HCl were prepared according to procedures described in the literature [16].

2.3. Synthesis of [MoO₂(HL¹)(CH₃OH)]Cl·1.5CH₃OH (**1a**)

A mixture of [MoO₂(acac)₂] (0.16 g, 0.5 mmol) and pyridoxal 4-phenylthiosemicarbazone hydrochloride H₂L¹·HCl (0.14 g, 0.4 mmol) in dry acetonitrile (10 mL) and dry methanol (10 mL) was refluxed for 5 hours. The solution was left at room temperature for one day and the obtained orange crystalline precipitate was filtered, rinsed with acetonitrile and dried in a desiccator down to constant weight. Yield: 0.14 g (55%). Anal. Calc. for C_{17.5}H₂₅ClMoN₄O_{6.5}S (M_r = 558.868): C, 37.61; H, 4.50; Cl, 6.34; N, 10.03; S, 5.74%. Found: C, 37.73; H, 3.95; Cl, 6.55; N, 10.56; S, 6.05%. TG: CH₃OH, 13.9% (Calcd. 14.3%); MoO₃, 26.5% (Calcd. 26.5%). Selected IR data (cm⁻¹): 1602, 1553 (C=N), 1367 (C–O), 939 (MoO₂)_{asym}, 904 (MoO₂)_{sym}, 639 (C–S).

2.4. Synthesis of $\{[MoO_2(HL)]Cl\}_n$ complexes

The appropriate pyridoxal thiosemicarbazone ligand $H_2L^{1-3}\cdot HCl$ (0.4 mmol) was dissolved in dry acetonitrile (20 mL), $[MoO_2(acac)_2]$ (0.16 g, 0.5 mmol) was added and the resulting solution was refluxed for the specified time. The precipitate that formed was filtered, washed with acetonitrile and dried in a desiccator down to constant weight.

R = Ph: Complex $\{[MoO_2(HL^1)]Cl\}_n$ (**1**): Dark red product. Reaction time: 5 hours. Yield: 0.22 g (94%). Anal. Calc. for $C_{15}H_{15}ClMoN_4O_4S$ (Mr = 478.76): C, 37.63; H, 3.16; Cl, 7.41; N, 11.70; S, 6.70%. Found: C, 37.41; H, 3.11; Cl, 7.14; N, 11.80; S, 6.32%. TG: MoO_3 , 29.8% (Calcd. 30.1%). Selected IR data (cm^{-1}): 1604, 1557 (C=N), 1362 (C-O), 954 (MoO_2)_{asym}, 935 (MoO_2)_{sym}, 641 (C-S). UV-VIS(methanol): λ/nm ($\epsilon/dm^3 mol^{-1} cm^{-1}$): 354 (14888) and 434 (8368).

R = Me: Complex $\{[MoO_2(HL^2)]Cl\}_n$ (**2**): Brown product. Reaction time: 12 hours. Yield: 0.087 g (43%). Anal. Calc. for $C_{10}H_{13}ClMoN_4O_4S$ (Mr = 416.69): C, 28.82; H, 3.14; Cl, 8.51; N, 13.45; S, 7.70%. Found: C, 28.66; H, 3.25; Cl, 8.21; N, 13.50; S, 7.65%. TG: MoO_3 , 34.3 (Calcd. 34.5%). Selected IR data (cm^{-1}): 1591, 1540 (C=N), 1358 (C-O), 930 (Mo=O), 801 (Mo=O...Mo), 621 (C-S). UV-VIS(methanol): λ/nm ($\epsilon/dm^3 mol^{-1} cm^{-1}$): 325 (17992) and 405 (2336).

R = H: Complex $\{[MoO_2(HL^3)]Cl\}_n$ (**3**): Dark brown product. Reaction time: 12 hours. Yield: 0.08 g (46 %). Anal. Calc. for $C_9H_{11}ClMoN_4O_4S$ (Mr = 402.66): C, 26.85; H, 2.75; Cl, 8.80; N, 13.91; S, 7.96%. Found: C, 26.53; H, 2.92; 8.54; N, 13.65; S, 7.18%. TG: MoO_3 , 35.5% (Calcd. 35.8%). Selected IR data (cm^{-1}): 1617, 1559 (C=N), 1357 (C-O), 930 (Mo=O), 815 (Mo=O...Mo), 621 (C-S). UV-VIS(methanol): λ/nm ($\epsilon/dm^3 mol^{-1} cm^{-1}$): 342 (177500) and 427 (1927).

2.5. X-Ray Crystallography.

The single-crystal X-ray diffraction data of **1a** was collected by ω -scans on an Oxford Diffraction Xcalibur 3 CCD diffractometer with graphite-monochromated Mo-K α radiation ($\lambda = 0.71073 \text{ \AA}$). The data reduction was performed using the CrysAlis software package [17]. Solution, refinement and analysis of the structure were done using the programs integrated in the WinGX system [18]. The structure was solved using SHELXS [19] by the Patterson method. The refinement procedure was performed by the full-matrix least-squares method based on F^2 against all reflections using SHELXL-97 [20]. One methanol molecule was found to be disordered over two sites and over a center of symmetry. The geometry of the two disordered methanol molecules were restrained to be as that of the methanol that is not disordered. The occupancy of the two sites refined to 0.263(8) and 0.237(8),

while they had a common temperature factor. The non-hydrogen atoms were refined anisotropically, except for those of the disordered methanol molecule which were refined isotropically. All hydrogen atoms, except those on the disordered methanol molecule, were located in the difference Fourier maps. Because of poor geometry for some of them they were placed in calculated positions and refined using the riding model. Only the hydrogen atoms of one free and the bound methanol hydroxyl group were fixed. Geometrical calculations were done using PLATON [21]. The structure drawings were prepared using PLATON and MERCURY [22] programs. The crystallographic data are summarized in Table 1, whereas the selected bond distances and angles are listed in Table 2.

Table 1. Crystallographic data for compound **1a**.

chemical formula	C _{17.5} H ₂₅ ClMoN ₄ O _{6.5} S
Mr	558.87
crystal colour, habit	orange-brown, plate
crystal size (mm ³)	0.593, 0.208, 0.042
crystal system	Triclinic
space group	P ⁻¹
unit cell parameters	
a (Å)	7.2147(5)
b (Å)	12.3246(10)
c (Å)	13.8100(11)
α, β, γ (°)	101.115(7), 100.204(6), 103.460(6)
V (Å ³)	1139.67(16)
Z	2
D _{calc} (g cm ⁻³)	1.623
temperature (K)	295
μ (mm ⁻¹)	0.827
F(000)	566
number of unique data	4807
number of data [F _o ≥ 4σ(F _o)]	3755
number of parameters	290
R ₁ ^a , [F _o ≥ 4σ(F _o)]	0.0486
wR ₂ ^b	0.1278
Min. and max.	-0.86, 0.49
electron density (e Å ⁻³)	

$$^a R = \frac{\sum ||F_o| - |F_c||}{\sum |F_o|}, \quad ^b wR = [\frac{\sum (F_o^2 - F_c^2)^2}{\sum w(F_o^2)^2}]^{1/2}$$

2.6. Reaction of complexes **1-3** with PPh_3

All reactions were performed under a dry argon atmosphere. PPh_3 (0.02 g, 0.08 mmol) and γ -picoline (8 μ L) were dissolved in dry acetonitrile (15 mL) and then **1**, **2** or **3** (0.08 mmol) was added. The suspension was heated for 10 h. The dark red-brown microcrystalline product was collected by filtration, washed with acetonitrile and dried.

R = Ph: Complex $[Mo_2O_3(HL^1)_2]Cl_2$ (**4**): Yield: 0.020 g (52%) (obtained from **1**). Anal. Calcd. for $C_{30}H_{30}Cl_2Mo_2N_8O_9S_2$ (Mr = 941.52): C, 38.27; H, 3.21; Cl, 7.53; N, 11.90; S, 6.81. Found: C, 38.11; H, 3.42; Cl, 7.21; N, 11.85; S, 6.55. TG: 30.8% MoO_3 (Calcd. 30.6%). Selected IR data (cm^{-1}): 1601, 1552 (C=N), 1363 (C-O), 948 (Mo=O), 788 (Mo-O-Mo), 625 (C-S).

R = Me: Complex $[Mo_2O_3(HL^2)_2]Cl_2$ (**5**): Yield: 0.015 g (46%) (obtained from **2**). Anal. Calcd. for $C_{20}H_{26}Cl_2Mo_2N_8O_7S_2$ (Mr = 817.38): C, 29.39; H, 3.21; Cl, 8.67; N, 13.71; S, 7.85. Found: C, 29.24; H, 3.68; Cl, 8.23; N, 13.64; S, 7.66. TG: 35.0% MoO_3 (Calcd. 35.2%). Selected IR data (cm^{-1}): 1606, 1553 (C=N), 1366 (C-O), 953 (Mo=O), 783 (Mo-O-Mo), 611 (C-S).

R = H: Complex $[Mo_2O_3(HL^3)_2]Cl_2$ (**6**): Yield: 0.013 g (41 %) (obtained from **3**). Anal. Calcd. for $C_{18}H_{22}Cl_2Mo_2N_8O_7S_2$ (Mr = 789.33): C, 27.39; H, 2.81; Cl, 8.98; N, 14.20; S, 8.12. Found: C, 27.25; H, 3.10; Cl, 8.65; N, 14.12; S, 7.98. TG: 36.5% MoO_3 (Calcd. 36.5%). Selected IR data (cm^{-1}): 1617, 1560 (C=N), 1343 (C-O), 948 (Mo=O), 779 (Mo-O-Mo), 621 (C-S).

2.7. General procedure for the epoxidation of cyclooctene by aqueous TBHP

A mixture of cyclooctene (2.76 mL, 20 mmol), acetophenone (0.1 mL, internal reference) and Mo (pre)catalyst (0.01 mmol Mo) was stirred and heated up to 80 °C before adding aqueous TBHP (70 wt %, 5.48 mL, 40 mmol). The reaction was followed for 6 h. Aliquots (0.1 mL) of the organic phase were taken at required times from the reaction media, mixed with 2 mL of Et_2O and a small quantity of MnO_2 was added. The mixture was then filtered through silica and analyzed by 1H NMR in $CDCl_3$.

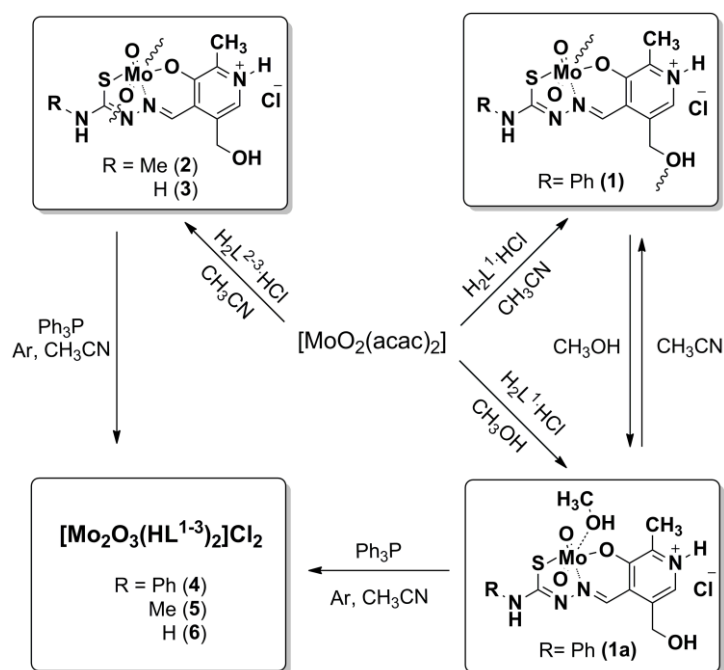
2.8. Computational Details

All geometries were optimized without any symmetry constraint with the Gaussian 03 program suite [23]. The input geometries were adapted from the X-ray structures of $[MoO_2(L^1)(MeOH)] \cdot CH_3OH$ (**1a***) [11] and $[MoO_2(HL^1)(MeOH)]Cl \cdot 1.5CH_3OH$ (**1a**). The calculations used the standard B3LYP three-parameter functional [24] with the 6-31G** basis set [25] for C, H, N and O atoms and the CEP-31G* basis set [26] for molybdenum. All optimized geometries were confirmed to be local minima by frequency analyses (Table S1 in the Supporting Information). The calculated IR spectra were

generated from the DFT-generated frequencies and intensities by applying Lorentzian functions and adjusting the linewidth to the best fit with experimental spectra.

3. Results and Discussion

New mononuclear $[\text{MoO}_2(\text{HL}^1)(\text{CH}_3\text{OH})]\text{Cl}\cdot 1.5\text{CH}_3\text{OH}$ (**1a**), and polynuclear molybdenum(VI) complexes $[\{\text{MoO}_2(\text{HL}^{1-3})\}\text{Cl}]_n$ (**1-3**) were prepared by the reaction of $[\text{MoO}_2(\text{acac})_2]$ with different pyridoxal thiosemicarbazone hydrochlorides (Scheme 2). Dry solvents (methanol and acetonitrile) were used to prevent ligand hydrolysis.



Scheme 2. Synthesis of Mo^{VI} complexes (**1a** and **1-3**) and Mo^{V} complexes (**4-6**).

Complexes **1a** and **1** were identified by the presence of $\nu(\text{MoO}_2)$ bands at 939 and 904 cm^{-1} for **1a**, and 954 and 935 cm^{-1} for **1** [27]. A band around 1071 cm^{-1} for both complexes was assigned to the $\nu(\text{C-O})$ of the coordinated methanol molecule in **1a** or the hydroxymethyl group in **1** (Scheme 2) [28]. On the other hand, the sixth coordination site in $[\{\text{MoO}_2(\text{HL}^{2,3})\}\text{Cl}]_n$ (**2** and **3**) is occupied by the oxygen atom from the $\text{Mo}=\text{O}$ unit from the neighbouring complex molecule, thus forming a coordination polymer. This is suggested by the intense IR bands observed at 801 cm^{-1} (**2**) and 815 cm^{-1} (**3**), as well as by a single strong absorption band at 930 cm^{-1} assigned to the $\text{Mo}=\text{O}\cdots\text{Mo}$ bridging and terminal $\text{Mo}=\text{O}$ groups, respectively [29]. The broad absorption bands in the $3000\text{-}2600\text{ cm}^{-1}$ range, appearing in the spectra of all complexes, confirm the presence of the pyridinium moiety. Two vibration bands in the $1610\text{-}1540\text{ cm}^{-1}$ range were assigned to the stretching vibrations of the $\text{C}=\text{N}$ groups of the

thiosemicarbazones [30], whereas those at about 625 cm^{-1} were attributed to the coordinated C–S_{thiol} group [31].

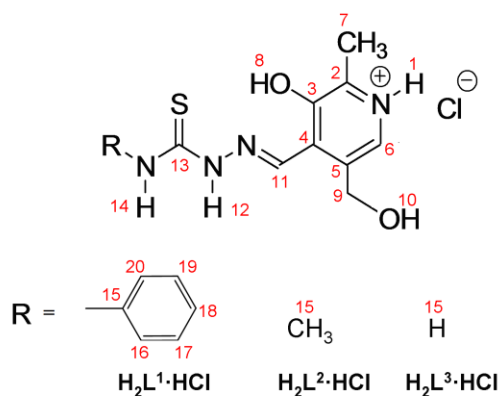
Polynuclear complexes **1-3** are not solvated. Upon heating the $\{[\text{MoO}_2(\text{HL}^{1-3})]\text{Cl}\}_n$ complexes under a pure oxygen atmosphere, only weight losses corresponding to complex decomposition occurred in the $227\text{--}430\text{ }^\circ\text{C}$ (**1**), $207\text{--}457\text{ }^\circ\text{C}$ (**2**) and $209\text{--}441\text{ }^\circ\text{C}$ (**3**) ranges. Crystals of **1a** were used for a TG analysis without grinding, to limit the loss of the solvent molecule. The first step in the TGA curve (weight loss within the $25\text{--}105\text{ }^\circ\text{C}$ range) was related to the solvent molecule release. Afterward, the intermediate product is stable up to $225\text{ }^\circ\text{C}$ when it starts to decompose and afford MoO_3 as the final residue. The mononuclear complex **1a** was heated from ambient temperature up to $175\text{ }^\circ\text{C}$. The substance obtained by this treatment was identified by chemical and TG analyses as the pentacoordinated $\{[\text{MoO}_2(\text{HL}^1)]\text{Cl}\}$ compound.

Even after prolonged heating of **1a** at $175\text{ }^\circ\text{C}$ in the solid state, the resulting material does not convert into **1**. Such thermally induced solid-state reaction is expected to occur only between molecules that are suitably arranged, with reacting centres close to each other within the crystal [32]. In the structure of **1a** the hydroxymethyl groups are not sufficiently close to the nearest Mo^{VI} centres of the $[\text{MoO}_2(\text{HL}^1)]^+$ cations (the $\text{Mo}\cdots\text{O}_2$ distance is 5.483 \AA , Figure S1) [32]. This was also confirmed by comparison of the IR spectrum of the non solvated product with those observed for **1a** and **1** (Figure S2 in the Supporting Information). Also, transformation of **1a** into a μ -oxo bridged dimer is not possible in the solid-state, since the $\text{Mo}=\text{O}_t$ group is not directed toward the vacant coordination site of the neighbouring $[\text{MoO}_2(\text{HL})]^+$ cation. This is additionally supported by absence of a strong band at $\sim 800\text{ cm}^{-1}$, which is characteristic for a molybdenum \cdots oxygen interaction.

On the other hand, transformation of the mononuclear complex **1a** into the coordination polymer **1** can be achieved by dissolution of **1a** in acetonitrile. Similarly, the reverse transformation can be obtained after exposure of the samples of **1** to methanol. Depending on the solvent nature, the methanol molecule in $[\text{MoO}_2(\text{HL}^1)(\text{CH}_3\text{OH})]^+$ cation can be easily replaced by the hydroxymethyl group of a neighbouring molecule and vice versa as already observed for the related neutral complexes [33].

3.1. NMR Spectroscopy

The NMR spectra of compounds **1-3** were recorded in DMSO-d^6 because of their very low solubility in non-coordinating solvents (the numbering scheme is given in Scheme 3 and all data are collected in Table S2, in Supporting Information). The solubilisation of the coordination polymers **1-3** is probably related to the coordination of DMSO to the Mo^{VI} center, breaking the infinite chain structure and yielding mononuclear $[\text{MoO}_2(\text{HL}^{1-3})(\text{DMSO})]^+$ complexes. The spectra of **1a** and **1** are very similar, the only difference being the presence of the MeOH molecule signal for the solution of **1a**.



Scheme 3. NMR numbering scheme of $\text{H}_2\text{L}^{1-3}\cdot\text{HCl}$.

^1H and ^{13}C chemical shifts are compared with those of the free ligand $\text{H}_2\text{L}^{1-3}\cdot\text{HCl}$. In the ^1H spectra of compounds **1-3**, the azomethine $\text{CH}=\text{N}$ proton signal H_{11} is shifted downfield up to 0.23 ppm as a result of the complexation. The absence of signals due to OH and $=\text{N}-\text{NH}$ (H_8 and H_{12} , respectively) suggests coordination through the oxygen atom and thioenolization of the $-\text{N}-(\text{C}=\text{S})-$ group into $-\text{N}=(\text{C}-\text{S})-$. Also, the H_{14} protons exhibit an upfield shift (up to 0.89 ppm) as a consequence of the coordination through the sulphur atom from the deprotonated thiol group. The ^{13}C chemical shifts exhibit larger changes, up to 7.35 ppm as observed for C_{11} and 11.49 ppm for C_{13} . These effects are downfield for the former and upfield for the latter site and confirm the coordination of imine-nitrogen and thiol-sulphur atoms to the $\{\text{MoO}_2\}^{2+}$ core. The effects observed for C_3 are all downfield and appreciably smaller (up to 1.57 ppm).

The ^1H and ^{13}C chemical shift values of $[\{\text{MoO}_2(\text{HL}^{1-3})\}\text{Cl}]_n$ (**1-3**) may be compared to those previously reported for the related complexes containing the doubly deprotonated ligands $[\text{MoO}_2(\text{L}^{1-3})]_n$ (Table S2 in SI, Figure S3). The spectral variations can be correlated with the protonation of the pyridoxal ligand. The signal at ca. 12 ppm (assigned to H_1) found in the spectra of compounds **1-3** is absent in the spectra of $[\text{MoO}_2(\text{L}^{1-3})]_n$. This confirms protonation of the pyridine ring N atom and the presence of the ligand as $(\text{HL})^-$. Such protonation induces a shift of all pyridine ring signals, particularly those of H_6 (up to +0.13 ppm) and C_6 atoms (up to -6.15 ppm). Additionally, the broad signal assigned to the hydroxymethyl H_{10} proton is also fairly shifted downfield by approximately 0.25 ppm. There is also a significant downfield shift of the H_{14} signal (up to 0.41 ppm) in complexes **1-3** with respect to the corresponding deprotonated complexes, which is probably the result of hydrogen bonding between the thiosemicarbazone NH group and chloride [34]. This suggests that the chloride counterion maintains this interaction in solution, as also found in the solid state (see next section).

3.2. X-Ray Crystallography

In the molecular structure of **1a** the coordination sphere around the Mo atom is a distorted octahedron with a tridentate ligand (HL^1)⁻ bound to the *cis*-{MoO₂}²⁺ core. The ligand is coordinated to molybdenum as *ONS*-donor through the phenolic-oxygen, the imine-nitrogen and the thiol-sulphur atoms. The sixth coordination site is occupied by the methanol molecule. The C1-S bond of 1.737(4) Å indicates the thiol form. Two chelate rings are formed by complexation of the ligand to Mo, a five- (Mo, S, C1, N2, N3) and a six-membered (Mo, N3, C2, C3, C7, O1) chelate ring that form a dihedral angle of 8.70(13)°. Therefore, the ligand is relatively planar with π -electron delocalization throughout the ligand. The substituted pyridine and phenyl rings form an angle of only 3.9(2)°. An ORTEP drawing of the complex cation and the chloride anion is shown in Figure 1. Relevant bond lengths and angles are given in Table 2.

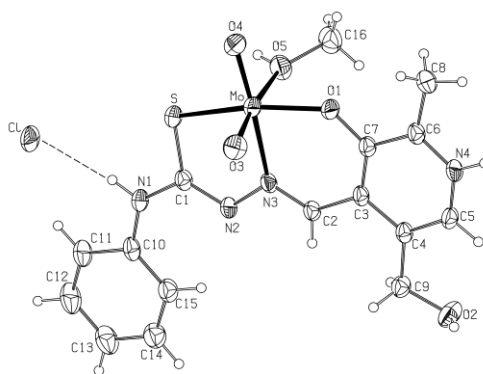


Figure 1. ORTEP plot of the structure of **1a** with displacement ellipsoids of non-hydrogen atoms drawn at the 30% probability level. Molecules of crystal methanol are omitted. Hydrogen bond is shown by a dashed line.

Table 2. Selected bond lengths (Å) and angles (°) for compound **1a**.

Bond lengths (Å)		Bond angles (°)	
Mo1-S	2.3936(10)	S-Mo-O1	149.40(10)
Mo1-O1	1.943(3)	S-Mo-O3	98.59(10)
Mo1-O3	1.684(4)	S-Mo-O4	92.97(10)
Mo1-O4	1.696(4)	S-Mo-O5	79.33(8)
Mo1-O5	2.332(4)	S-Mo-N3	76.11(7)
Mo1-N3	2.312(4)	O1-Mo-O3	100.22(14)
S-C1	1.737(4)	O1-Mo-O4	105.04(13)
N1-C1	1.350(5)	O1-Mo-O5	77.22(12)
N1-C10	1.407(6)	O1-Mo-N3	80.84(11)

N2-C1	1.318(5)	O3-Mo-O4	105.15(17)
N2-N3	1.377(5)	O3-Mo-O5	167.93(15)
N3-C2	1.290(4)	O3-Mo-N3	87.45(14)
N4-C5	1.319(5)	O4-Mo-O5	86.87(15)
N4-C6	1.330(5)	O4-Mo-N3	164.58(13)
C2-C3	1.457(4)	O5-Mo-N3	80.50(12)

The crystal structure of **1a** consists of $[\text{MoO}_2(\text{HL}^1)(\text{CH}_3\text{OH})]^+$ cations, chloride anions and molecules of methanol of which one is in a general position while the other one is disordered over a center of symmetry. Both protonated nitrogen atoms (N1 and N4) and the hydroxyl group from the pyridoxal moiety (O2) are involved in hydrogen bonding with the chloride ion, thus connecting three neighbouring complex cations and forming double-chains along the *c*-axis. The chains are linked through hydrogen bonding with one of the free methanol molecules along the *a*-axis. The solvent methanol molecule that is not disordered is a hydrogen bond acceptor from the bound methanol and a donor to the pyridoxal hydroxyl group. Hydrogen bonds are given in Table 3. Interactions between chains along the *b*-axis involve the disordered methanol molecule. The packing diagram is shown in Figure 2 (and Figure S4 in the Supporting Information).

Table 3. Geometry of the hydrogen bonds (Å, °) for compound **1a**.

D-H...A	D-H (Å)	H...A (Å)	D...A (Å)	D-H...A(°)
N1-H1...Cl _a	0.86	2.39	3.221(3)	162
N4-H4...Cl _b	0.86	2.18	3.033(3)	174
O2-H2 _a ...Cl	0.82	2.30	3.105(4)	169
O5-H5 _a ...O6	0.81	1.83	2.639(5)	175
O6-H6 _a ...O2 _c	0.99	1.91	2.878(6)	164

^a 2-x,1-y,1-z; ^b 2-x,1-y,2-z, ^c 1-x,1-y,1-z

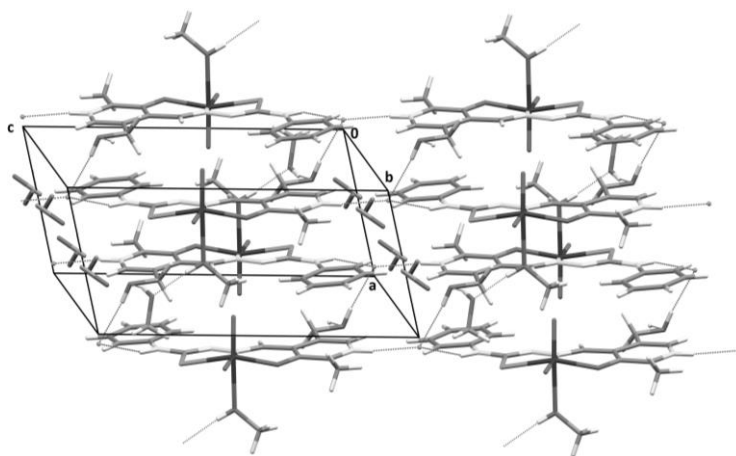


Figure 2. Packing of **1a** in the unit cell. Hydrogen bonds are shown by dotted lines. All four possible molecule sites of the disordered methanol molecule are shown (without their hydrogen atoms).

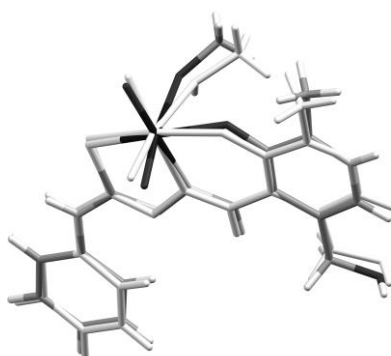


Figure 3. Superimposed molecules of **1a** (light grey) and **1a*** (dark grey).

In contrast to **1a** where N4 is a hydrogen bond donor, the same atom in the non protonated complex, $[\text{MoO}_2(\text{L}_1)(\text{CH}_3\text{OH})]\cdot\text{CH}_3\text{OH}$ (**1a***), is a hydrogen bond acceptor [11]. Significant changes in the geometry of **1a** in comparison to **1a*** include angles of the Mo coordination polyhedron and the C5-N4-C6 angle in the pyridoxal ring, which increases upon protonation from $119.6(2)^\circ$ to $124.6(3)^\circ$, Figure 3. The largest difference in the coordination polyhedron concerns the S-Mo-O5 angle, most probably caused by involvement in different hydrogen bonding pattern and also different packing in the crystal.

3.3. DFT calculations

DFT geometry optimizations were carried out to assign the IR absorption bands in **1a** and **1a***, in order to assess the effect of ligand protonation. Before examining these results, we present a brief description of the optimized geometries. Relevant results are collected and compared to the experimental crystallographic data in Table 4. The agreement between experimental and calculated geometries for the core is generally very good and small differences may be at least in part attributed to crystal packing, which is not accounted for in the single molecule calculation. All calculated distances are overestimated,

very little for the terminal oxido ligands (maximum deviation 0.01 Å), to an intermediate degree for Mo–O(1), Mo–S and Mo–N (0.03 Å to 0.06 Å), and to a greater extent for the coordinated methanol O atom (by 0.14 Å on average).

Table 4. Comparison of relevant bond lengths (Å) between experimental data for **1a** and **1a*** and related calculated structures [MoO₂(HL¹)(MeOH)]⁺ (**A**) and [MoO₂(L¹)(MeOH)] (**A***). For **A**, only the cation had been taken into account for the calculation. Atom labels refer to the crystallographic structure.

Bonds	1a	A	1a*	A*
Mo-O(1)	1.943	1.987	1.918	1.975
Mo-O(3)	1.684	1.691	1.689	1.693
Mo-O(4)	1.696	1.697	1.706	1.706
Mo-O(5)	2.332	2.455	2.336	2.484
Mo-N(3)	2.312	2.373	2.280	2.339
Mo-S(1)	2.394	2.448	2.415	2.449
N(3)-N(2)	1.378	1.345	1.364	1.370

The elongation/shortening of bond lengths experimentally observed between neutral and charged complexes (**1a*** and **1a**) is reproduced by the DFT calculations, with similar differences. Only the most relevant data is listed in Table 4.

The experimental IR spectra of **1a** and **1a*** are compared with the calculated ones for **A** and **A*** in Figure 4. Although shifted by around 60 cm⁻¹ on average, the patterns are similar. Three vibrations at 981 (asym), 1006 (sym), 1020 (sym) cm⁻¹ for **A** can be attributed to MoO₂ (the symmetric vibration is coupled with other vibrations and contributes significantly to two normal modes) and two vibrations at 968 (asym) and 1001 (sym) cm⁻¹ for **A***. The calculations are also useful for the assignment of other vibrational modes, such as the C–S vibrations (685 cm⁻¹ for **A** and 679 cm⁻¹ for **A***) the N–N vibrations (1190 cm⁻¹ for **A** and 1152 cm⁻¹ for **A***), the CH=N vibrations (coupled with other vibrations in bands at 1617, 1656 and 1658 cm⁻¹ for **A** and at 1560, 1624 and 1653 cm⁻¹ for **A***) and finally the pyridoxal N–H vibration in case of **A** (3569 cm⁻¹). The shift relatively to the experimental values is observed in all cases, as typically observed for calculations carried out at this level of theory.

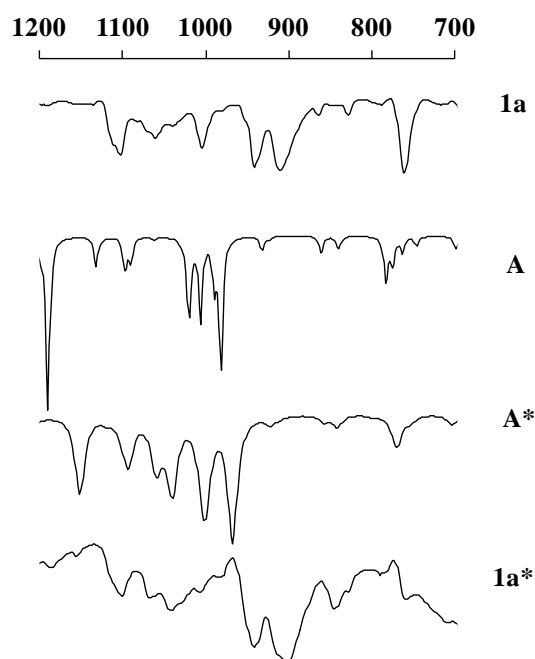


Figure 4. Comparison of experimental (**1a** and **1a***) and theoretical (**A** and **A***) IR vibration pattern in the range 700-1200 cm^{-1} .

3.4. UV-Vis Spectroscopy

The electronic spectra of the thiosemicarbazone complexes **1–3** recorded in methanol exhibited $S(p\pi)\text{--}Mo(d\pi)$ LMCT in the regions 405 – 434 nm as well as $N(p\pi)\text{--}Mo(d\pi)$ and $O(p\pi)\text{--}Mo(d\pi)$ LMCT bands at 325-354 nm [35]. These spectral characteristics are influenced by the nature of the ligand and by its degree of protonation. Titrations of **1–3** with HCl or Et_3N were performed by adding aliquots of HCl or Et_3N solution (2 μL) into cuvettes in methanol with UV-Vis monitoring and compared with those of the corresponding $\text{H}_2\text{L}^{1-3}\cdot\text{HCl}$ ligands. A titration experiment was also carried out by adding HCl to **1a***. Spectral changes associated to system **1** and the ligand $\text{H}_2\text{L}^1\cdot\text{HCl}$ are presented in Figure 5.

The addition of HCl to the solution obtained from **1** resulted in a shift of the 434 and 354 nm bands to higher wavelengths (461 and 363 nm, respectively) with a change of molar absorptivity. These changes are presumably due to the protonation of the ligand and formation of the $[\text{MoO}_2(\text{H}_2\text{L}^1)(\text{CH}_3\text{OH})]^{2+}$ cation. Therefore, the corresponding bands of **1a***, containing the ligand in the most deprotonated form, are at lower wavelengths than for **1**, exhibiting two bands at 416 and 349 nm.

The addition of Et_3N to the methanolic solution of **1** resulted in the bands at 390 and 312 nm. The spectrum of the obtained solution is different from the spectrum of the $[\text{MoO}_2(\text{L}^1)(\text{CH}_3\text{OH})]$ complex. However, it is similar to that of the deprotonated ligand $(\text{L}^1)^{2-}$ obtained upon addition of Et_3N to the solution of $\text{H}_2\text{L}^1\cdot\text{HCl}$. Apparently, addition of Et_3N to the solution destabilizes the association between

the ligand and the molybdenum atom, probably yielding the free doubly deprotonated ligand and the molybdate ion, MoO_4^{2-} . The same result was obtained also using KOH in methanol instead of Et_3N . The corresponding behaviour of compounds **2** and **3** is identical. In the case of **2**, addition of HCl caused a shift of the 405 and 325 nm bands to 442 and 360 nm, respectively. Similarly, the spectrum of **3** has absorption maxima at 427 and 342 nm, whereas acidification with HCl shifted the bands to 438 and 361 nm, respectively.

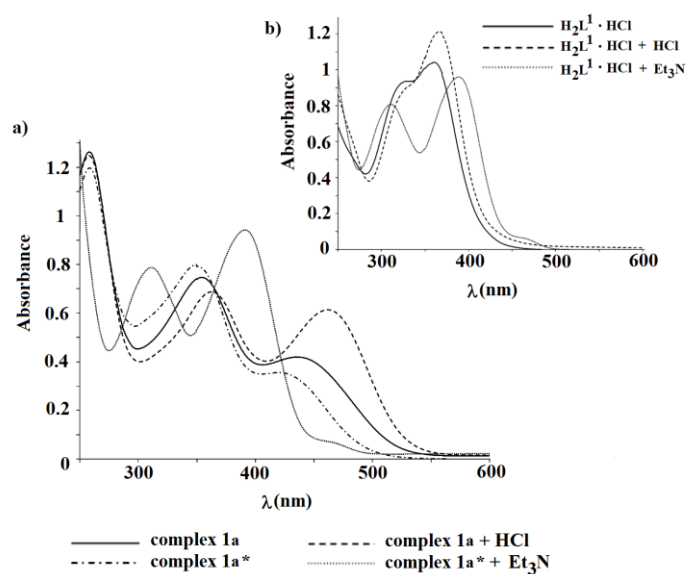


Figure 5. (a) UV-Vis spectra of thiosemicarbazone complexes $[\text{MoO}_2(\text{HL}^1)(\text{CH}_3\text{OH})]\text{Cl} \cdot 1.5\text{CH}_3\text{OH}$ (**1a**) and $[\text{MoO}_2(\text{L}^1)(\text{CH}_3\text{OH})] \cdot \text{CH}_3\text{OH}$ (**1a***) in methanol at 25 °C. (b) UV-Vis spectrum of $\text{H}_2\text{L}^1 \cdot \text{HCl}$ in methanol at 25 °C.

3.5. Reactivity of Mo(VI) complexes towards oxygen acceptor: extension of the OAT reaction

Reactions of dioxomolybdenum(VI) complexes **1-3** with triphenylphosphine (PPh_3) were performed in dry acetonitrile under an inert atmosphere and in the presence of γ -picoline to improve the solubility. OAT reactions resulted in the formation of dinuclear oxo-bridged molybdenum(V) complexes of general formula $[\text{Mo}^{\text{V}}_2\text{O}_3(\text{HL}^{1-3})_2]\text{Cl}_2$ (**4-6**) (Scheme 1), the nature of which was proven by chemical analyses, IR spectroscopy and TGA measurements.

The IR spectra of **4-6** exhibit bands around 950 cm^{-1} and 780 cm^{-1} , assigned to terminal $\text{Mo}=\text{O}$ and $\text{Mo}-\text{O}-\text{Mo}$ bridging groups, respectively (Figure S5 Supporting Information) [36]. All compounds were obtained pure with a 40% yield when the reaction proceeded for 10 hours. The diamagnetism, which is characteristic for oxo bridged Mo(V) dimers [37], was confirmed by magnetochemical measurements. Our results are consistent with findings reported by Young et al. [38]. As it has been previously shown, formation of $\{\text{Mo}^{\text{V}}_2\text{O}_3\}^{4+}$ species from $\{\text{Mo}^{\text{VI}}\text{O}_2\}^{2+}$ seems to be the result of OAT from $[\text{Mo}^{\text{VI}}\text{O}_2(\text{HL}^{1-}$

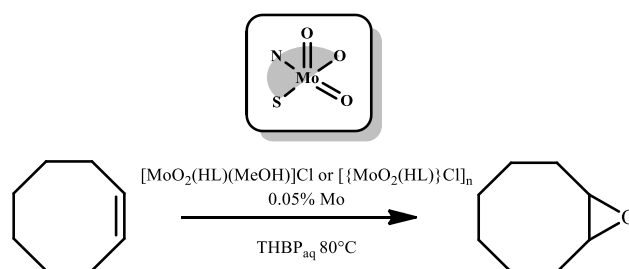
³)Cl (**1-3**) to PPh₃ [38,39]. Weight losses corresponding to the complex decomposition were observed by TGA (under a pure oxygen atmosphere) in the 246-446 °C (**4**), 239-431 °C (**5**) and 223-448 °C (**6**) ranges, leading to MoO₃.

The reactivity of complexes **4-6** towards oxygen donors is revealed by dissolution in DMSO, which leads to an immediate colour change from dark red to pale orange, as a consequence of the reoxidation to **1-3** [40]. For this reason, NMR spectra of these compounds could not be recorded. Unfortunately, all other attempts to characterize **4-6** were hampered by their poor solubility even in very polar solvents.

3.6. Catalytic Epoxidation Reactions

We have recently reported the catalytic activity of the neutral [MoO₂(L¹)(MeOH)] (**1a***) and [MoO₂(L¹⁻³)]_n (**1*-3***) complexes for the epoxidation of cyclooctene by aqueous TBHP (Scheme 4 and Scheme S1 in the Supporting Information) under solvent-free conditions and with a low Mo:olefin ratio (1:2000) [13]. We have now carried out a comparative study of the same catalytic reactions under the same conditions for the cationic complexes in **1-3** and **1a** in order to evaluate and clarify the effect of ligand protonation.

Like the corresponding neutral complexes, the protonated analogues were sparingly soluble in cyclooctene and insoluble in water at room temperature, but completely dissolved in the organic phase after addition of aqueous TBHP at 80 °C. The reactions were followed by ¹H NMR during 6 hours. The relevant results are summarized in Table 5 and the kinetic profiles are shown in Figure 6. The ligands alone were not active and the reference [MoO₂(acac)₂] (pre)catalyst gives a conversion of 54% with a 82% epoxide selectivity under these conditions [13].



Scheme 4. General equation of Mo-catalyzed epoxidation of cyclooctene.

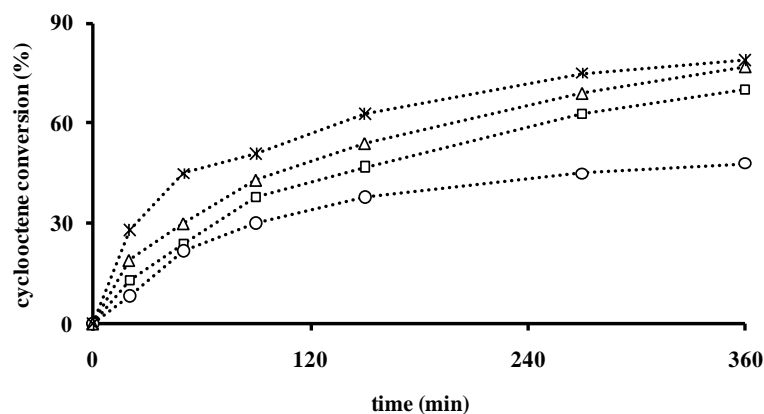


Figure 6. Converted cyclooctene vs. time with dioxomolybdenum(VI) (pre)catalysts: □ **1**, ○ **1a**, △ **2**, * **3**. Conditions: Mo/substrate/TBHP = 1/2000/4000, T = 80 °C.

Table 5. Comparison of epoxidation data for dioxomolybdenum(VI) (pre)catalysts.

Complex	1a	1a* ^[13]	1	1* ^[13]	2	2* ^[13]	3	3* ^[13]	[MoO ₂ (acac) ₂] ^[13]
Conversion (%) ^a	48	97	70	54	78	71	79	69	54
Selectivity (%) ^b	74	97	82	84	86	92	89	99	82
TOF (h ⁻¹) ^c	480	3360	787	645	1080	960	1680	1080	530
TON ^d	960	1940	1960	1040	1253	1420	1580	1380	1300

^a Calculated after 6 h, ^b formed epoxide per converted olefin after 6 h, ^c cyclooctene transformed / catalyst / time at 20 minutes, ^d cyclooctene transformed / catalyst at 6 h

All compounds show relatively high selectivities (74-89%) with moderate to very good conversions after 6 h (48-79%) in the order **1a** < **1** < **2** ~ **3**. Surprisingly, the selectivity and the conversion with the mononuclear charged compound **1a** are the lowest (even lower than for [MoO₂(acac)₂]), at variance with the results obtained with the corresponding neutral Mo^{VI} compound **1a***, where the mononuclear MeOH adduct performed better than the coordination polymers. The TON with **1**, **2** and **3** (1960, 1253 and 1580) may be compared with those of the corresponding neutral systems (1040 for **1***, 1420 for **2*** and 1380 for **3***, respectively [13]). Hence, the ligand protonation has a strongly positive influence for L¹, a moderately negative one in the case of L² and a weakly positive one for L³. However, ligand protonation has a lower effect on the activity than methanol coordination. A visual comparison of the conversion and selectivity within the four complexes (mononuclear or polynuclear, neutral or cationic) containing ligand L¹ is shown in Figure 7. All these results might be linked to the nature of the ligands, to the protonation state and to the energy required to remove methanol in case of mononuclear compounds. To our knowledge, under solvent-free conditions, neutral compounds with similar ligands and based on vanadium instead of molybdenum as the catalytic metal performed less efficiently as greater metal

loadings were required [41]. Several literature references concern the epoxidation of olefins using molybdenum containing complexes (1% Mo *vs.* substrate being the usual amount) with similar tridentate ligands as catalyst. Some of these catalysts exhibit very high TOF and are highly selective, but most of them were only used in solution of organic solvents (often dichloroethane) and/or with TBHP in decane as the oxidant [42]. The results presented herein are relevant because the process does not use solvent and employs TBHP in water as oxidant with a low catalysts loading.

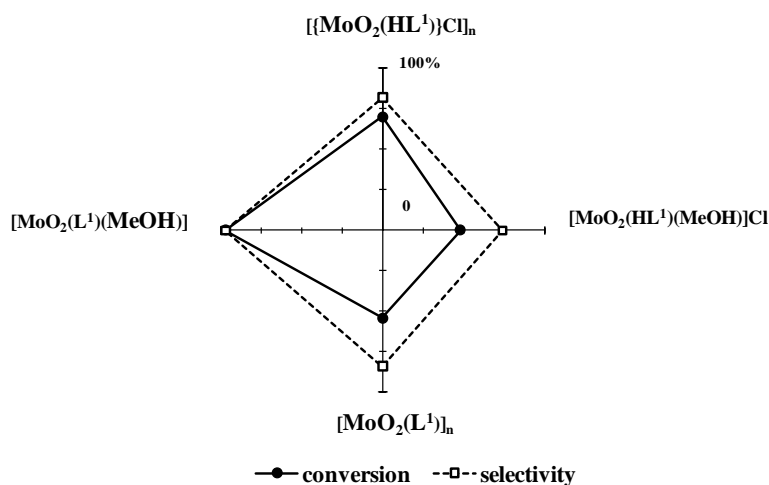


Figure 7. Conversion and selectivity comparison in epoxidation of cyclooctene by aqueous TBHP catalyzed by charged/neutral and/or monomeric/polymeric Mo^{VI} complexes synthesized from the H_2L^1 ligand. The results for the neutral compounds are taken from ref [13].

4. Conclusion

Mononuclear and polynuclear charged molybdenum(VI) complexes with singly-deprotonated pyridoxal thiosemicarbazonato ligands have been synthesised and characterized. They have been used as precursors for the synthesis of corresponding dinuclear oxo-bridged molybdenum(V) complexes by reaction with PPh_3 . The catalytic activity has been tested for cyclooctene epoxidation by aqueous TBHP under solvent-free conditions. The emphasis was placed on very low Mo loading (0.05% *vs.* substrate). These compounds are quite active although generally not as much as the corresponding neutral analogues. Nevertheless, the results obtained with the charged polymeric materials provide a good basis for further optimization aiming at efficient catalyst recovery and recycling.

Supporting Information

Appendix A. Supplementary data - CCDC 829512 contains the supplementary crystallographic data for **1a**. These data can be obtained free of charge via <http://www.ccdc.cam.ac.uk/conts/retrieving.html>, or

from the Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: (+44) 1223-336-033; or e-mail: deposit@ccdc.cam.ac.uk.

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