ABSTRACT: The tridentate ligand \( L \) and its complexes with transition-metal ions have been prepared and characterized. The polycondensation reactions of transition-metal complexes with different dialdehydes led to the formation of transition-metal-complex-based polyazomethines, which have been obtained by on-substrate polymerization, and their electrochemical and electrochromic performance have been investigated. The most interesting properties are exhibited by polymers of \( \text{Fe(II)} \) and \( \text{Cu(II)} \) ions obtained by the reaction of the appropriate complexes with a triphenylamine-based dialdehyde. \( \text{Fe(II)} \) polymer \( P_1 \) undergoes a reversible oxidation/reduction process and a color change from orange to gray due to the oxidation of \( \text{Fe(II)} \) to \( \text{Fe(III)} \) ions concomitant with the oxidation of the triphenylamine group. Its electrochromic properties such as long-term stability, switching times, and coloration efficiencies have been investigated, providing evidence of the utility of the on-substrate polycondensation reaction in the formation of thin films of electrochromic metallopolymers.

INTRODUCTION

Polymeric complexes of transition-metal ions have been attracting interest in many scientific and technological fields in recent years due to their multiple applications.\(^{1-3}\) Complexes of transition-metal ions are known to be interesting materials for electrochromic applications, and the color change can be based on the redox reaction of the ligand molecule and/or the metallic center. The incorporation of transition-metal ions into the polymer backbone can also change the electrochromic properties of polymeric materials. This is the result of the formation of additional absorption bands that change the color of the material, such as metal to ligand (MLCT) or ligand to metal charge transfer (LMCT) bands as well as \( d-d \) transitions that are characteristic for transition-metal complexes.\(^{4-6}\)

The synthesis of high-molecular-weight conjugated polymers containing transition metals has been often hampered by synthetic difficulties and/or a solubility problem. Metallosupramolecular polymers can be prepared by the complexation of metal ions with multidentate organic ligands or polymers\(^ {7-11} \) as well as by cross-coupling reactions,\(^ {12} \) radical polymerization,\(^ {13} \) or electropolymerization.\(^ {14-16} \) One of the methods of preparation of polymers is chemical linking of single or multiple kinds of monomers to form long chains, releasing water or a similar simple substance, called a polycondensation reaction. Hasanain et al. described a synthesis via a polycondensation reaction and the electrochromic characterization of a dinuclear ruthenium complex incorporated into the polyimide polymer main chains.\(^ {19} \) The polymers exhibited interesting electrochromic properties, but they were not soluble in common organic solvents, which can hamper their processing. An example of the application of polycondensation to the preparation of metallopolymers was also demonstrated by the synthesis of electrochromic \( \text{Co(III)} \)- and \( \text{Fe(III)} \)-based metallosupramolecular polymers with aromatic azo ligands that conferred good solubility to the metal complex, and due to this the solubility issue has been overcome.\(^ {20} \)

Polyazomethines can also be prepared by an on-substrate method by heating monomers deposited onto the substrate under an acetic atmosphere.\(^ {21-23} \) Polymers prepared in this way are obtained in the form of thin, insoluble layers on the substrate, and they cannot be characterized using conventional methods, such as gel permeation chromatography (GPC), but the insolubility of the polymer layer is desirable in case of application as an active layer in electrochromic devices. The miscibility of the organic layer with the electrolytic gel is problematic, because the electroactive layer can subsequently

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delaminate from the electrode, resulting in a limited device lifetime, poor device performance, localized color defects, and poor color contrast.\textsuperscript{21} An advantage of the on-substrate polymerization method is also the easy introduction of different functional groups, which in turn allows tuning of the polymer properties, and simple preparation and purification with only one byproduct, which is water.

Herein, we present the synthesis and characterization of polyazomethines containing complexes of transition-metal ions. Monomers for the synthesis of electrochromic polyazomethines by the on-substrate polymerization method are usually organic dialdehydes and diamines, but the use of transition-metal complexes has not been presented before. Complexes of transition-metal ions containing four amine groups that are able to undergo a condensation reaction with aldehydes have been obtained by the self-assembly of transition-metal ions with organic ligand 2,6-bis(1-hydrazonooethyl)pyridine (L). Complexes have been characterized by spectroscopic methods and X-ray analysis and have been used in polycondensation reactions with three different dialdehydes, leading to the preparation of several polyazomethines (P1–P6). An investigation of the electrochemical and electrochromic properties of the obtained materials allowed us to identify the key monomers that give polymers with the desired properties.

**EXPERIMENTAL SECTION**

**General Considerations.** Dialdehydes 4 and 5 have been prepared according to a previously reported procedure.\textsuperscript{24} All other reagents are commercially available and were used as received. Acetonitrile was dried by passing over neutral alumina followed by heating at 90 °C under argon atmosphere overnight. The solution was concentrated and cooled with ice, and the obtained white crystals were filtered off, washed with a small amount of cold ethanol, and dried.

**Measurement details of complexes 1–3 are summarized in Table S1.**

**Synthesis.** Ligand L. Ligand L has been obtained according to a previously reported method.\textsuperscript{29} 2,6-Diacetylpyridine (0.5 g, 3.07 mmol) was dissolved in absolute ethanol (10 mL). Afterward, excess hydrazine (5.0 mL) was added to the solution. The mixture was heated at 90 °C under argon atmosphere overnight. The solution was concentrated and cooled with ice, and the obtained white crystals were filtered off, washed with a small amount of cold ethanol, and dried.

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RESULTS AND DISCUSSION

Ligand L has been prepared in a condensation reaction between 2,6-diacylpyridine and hydrazine monohydrate, as outlined in Figure 1.

Figure 1. Synthesis of ligand L

The backbone of ligand L contains a structural unit similar to that of terpyridine, and it has been designed to act as a tridentate donor ligand in reactions with transition-metal ions to form complexes of 1:2 metal to ligand stoichiometry that contain four uncoordinated amine groups able to undergo a condensation reaction with other functional groups. The ligand is known to be potentially pentadentate, but it mainly acts as a tridentate ligand. The reactions of ligand L with iron(II) and copper(II) salts were carried out in a dichloromethylene/acetonitrile mixture (1:1 v/v) at room temperature for 24 h. Complexes have been obtained as colored solids by precipitation using diethyl ether, isolated and characterized. The 1:2 metal:ligand stoichiometry of obtained complexes has been confirmed by high-resolution electrospray ionization mass spectra. For example, the signals at m/z 219.0837 and 525.1710 in the ESI-MS spectra of complex 1 have been assigned to [FeL][BF4] and [FeL][BF4]2, respectively, and clearly indicate the formation of a complex with a 1:2 stoichiometry. In the ESI-MS spectra signals at m/z 437.1605 and 444.1564 are also present, which have been assigned to molecular cations with a 1:2 metal:ligand stoichiometry in which one of the ligand molecules is deprotonated, forming the molecular cations [Fe(LH)][BF4] and [Cu(LH)][BF4], respectively. The infrared spectra of the complexes of transition-metal ions provide useful information about the coordination of the functional groups upon the formation of the complexes in the solid state. The IR spectra of the products have been assigned by comparison with the stretching frequencies of the free ligand L. The peaks in the IR spectra of free ligand L were consistent with those observed for similar hydrazine derivatives. The ligand L and its complexes with transition-metal ions show characteristic IR bands in the range 1647–1630 cm−1, due to the stretching vibration of the imine C=N bond (Figures S1–S4). The weak bands in the spectra of the ligand L at around 3080 and 3026 cm−1 were attributed to an aromatic C–H stretching vibration, whereas the bands at 2933 and 2907 cm−1 were assigned to the C–H stretching vibration of the methyl group. The two bands in the range 3150–3350 cm−1 in the FT-IR spectra of both the ligand and the complexes were assigned to asymmetric and symmetric stretches of the amine group, confirming that –NH2 groups in the transition-metal complexes still exist.
The exact crystal structures of complexes 1−3 have been confirmed by X-ray crystallography. Single crystals appropriate for an X-ray analysis have been obtained by slow diffusion of diethyl ether into acetonitrile solutions of complexes. Figure 2A−C shows the perspective views of the dications 1−3, respectively; Table 1 gives the relevant geometrical parameters. All complexes have an ML₂ structure, and the overall shapes and coordinations of all four dications (in the structure of 1 there are two symmetry-independent complexes) are very similar (Figure 2D). The coordination of the metal cation is best described as distorted octahedral (cf. Table 1), with a quite linear \( \text{N(pyridine)} - \text{M} - \text{N(pyridine)} \) angle.

In the crystal structures the cations, anions, and solvent molecules make three-dimensional networks by means of Coulombic interactions, hydrogen bonds between \( \text{N} - \text{H} \) groups and counterion F or O atoms, and van der Waals contacts.

The polycondensation reactions between various dialdehydes 4−6 and complexes 1 and 2 being tetraamine monomers shown in Figure 3 led to the preparation of polyazomethines containing transition-metal ions.

The dialdehydes have been selected to investigate the influence of the central groups on the electrochemical and electrochromic properties of the obtained polymers. The dialdehyde 6 does not contain an electrochromic group, while dialdehydes 4 and 5 contain electroactive triphenylamine and thiophene groups, respectively, which are known to affect the electrochromic properties of polymers.²⁴

To evaluate the thermal stability of ligand \( \text{L} \) and its complexes and to confirm the stability of the compounds under on-substrate polymerization conditions, a thermogravimetric analysis (TGA) was carried out. The TGA measurements were done in a temperature range of 30−600 °C under a nitrogen atmosphere at a heating rate of 10 °C min⁻¹. It was found that ligand \( \text{L} \) was stable up to 180 °C before it underwent any significant decomposition (Figure S5), whereas complexes with transition-metal ions were found to decompose above 215 °C (1), 190 °C (2), and 310 °C (3) (Figures S6−S8). The small weight loss in the temperature range of 30−100 °C observed in the case of complexes of transition-metal ions has been assigned to the loss of solvent molecules.

Table 1. Relevant Geometrical Parameters (Å, deg) with su Values in Parentheses

|       | 1A (M = Fe)  | 1B (M = Fe)  | 2 (M = Cu)  | 3 (M = Cu)  |
|-------|--------------|--------------|-------------|-------------|
| M1−N1 | 1.874(5)     | 1.878(5)     | 1.960(2)    | 1.9360(16)  |
|       | 1.882(5)     | 1.884(5)     | 1.965(2)    | 1.9749(16)  |
| M1−N23| 1.955(5)     | 1.963(5)     | 2.186(2)    | 2.1344(17)  |
|       | 1.955(6)     | 1.967(5)     | 2.188(2)    | 2.1396(17)  |
| M1−N63| 1.959(5)     | 1.980(5)     | 2.206(2)    | 2.2710(17)  |
|       | 1.978(5)     | 1.988(5)     | 2.214(2)    | 2.2802(17)  |
| angles| 176.6(2)     | 176.9(2)     | 173.31(9)   | 178.80(7)   |
|       | 160.0(2)     | 159.1(2)     | 153.31(8)   | 155.43(6)   |
|       | 159.4(2)     | 159.0(2)     | 151.24(8)   | 151.77(6)   |

Figure 2. Perspective views of one of the symmetry-independent cationic complexes of 1 (A) and cations observed in the structures of 2 (B) and 3 (C). Ellipsoids are drawn at the 50% probability level, and hydrogen atoms are shown as spheres of arbitrary radii. (D) Comparison of four dications, where the metal and coordinated N atoms were fitted onto one another. Color codes: 1A, green; 1B, pale green; 2, red; 3, blue (the asymmetric part of the unit cell of 1 contains two symmetry-independent molecules).
The polycondensation of complexes 1 and 2 with dialdehydes was done by an on-substrate polymerization method. To do this, a mixture of the monomers in a 1:2 transition-metal complex:dialdehyde molar ratio was manually spray coated onto ITO-coated glass slides and the plates were heated at 100 °C for 1 h under a saturated trifluoroacetic acid atmosphere. Afterward, the substrates were rinsed with dichloromethane solution containing triphenylamine and pure dichloromethane to neutralize the acid and remove any unreacted monomers and low-molecular-weight oligomers. To characterize the polymers by TGA analysis and infrared spectroscopy, polymers were also obtained in solution. To do this, a solution of complex 1 or 2 in acetonitrile was added a solution of the appropriate dialdehyde (2 equiv) in chloroform. Then a catalytic amount of TFA (5 mol %) was added to the mixture of monomers and the solution was stirred and heated at 70 °C for 24 h. Afterward triethylamine was added to neutralize the acid and precipitated polymers were filtered off, washed with acetonitrile and chloroform to remove unreacted monomers, and dried on air. Polymers P1−P6 were found to exhibit thermal stability similar to that of complexes of transition-metal ions that were used as monomers (Figures S9−S14). Decomposition of the iron-based polymers P1, P3, and P5 was found to occur above 300 °C, whereas copper-based polymers P2, P4, and P6 decompose at temperatures above 200 °C. The small weight losses at temperatures below 200 °C have been assigned to the loss of solvent molecules adsorbed on the surface of polymer and/or occluded in the polymers. In the FT-IR spectra, due to the existence of two imine bonds in different surroundings in the polymers, two characteristic bands can be attributed to the imino groups (Figures S15−S20). In the spectra there are also very weak signals in the ranges of 3350−3200 and 1690−1670 cm⁻¹ assigned to amine NH₂ and carbonyl C=O groups, respectively, that indicate the existence of unreacted end groups of the polymers.

The composition of polymers obtained by on-substrate polymerization was analyzed using X-ray photoelectron spectroscopy (XPS). As seen in Figure S21, XPS survey spectra of complex 1 and its polymers contain peaks of the core levels C 1s, N 1s, Fe 2p, B 1s, F 2p, and S 2p in case of polymer P3, whereas in case of complex 2 and its polymers core levels of C 1s, N 1s, Cu 2p, S 2p, O 2p, and F 2p have been detected (Figure S22). Additionally, in the case of polymers P1−P6 the XPS survey spectra contain In, Sn, Si, and O core levels of the ITO support. The 2p Fe and 2p Cu peaks for polymers P1 and P2 appeared at 703.96 and 932.94 eV for Fe 2p3/2 and Cu 2p3/2, respectively, which was lower than those of complexes 1 and 2 (708.47 and 935.04 eV for Fe 2p3/2 and Cu 2p3/2, respectively) (Figure S23). These results suggest that metal ions become more cationic in polyazomethines, most likely due to the lower electronegativity of imine nitrogen atoms in comparison to amine nitrogen atoms that increases the positive charge on metal ions.

The obtained compounds have been characterized in terms of their electrochemical and electrochromic properties. First, absorption spectra in the visible region of ligand L and its complexes in a dichloromethane solution in the case of ligand L and in acetonitrile solutions for the complexes have been recorded (Figure 4).

Ligand L exhibits no absorption in the visible range with a strong absorption below 340 nm (Figure S24). The Fe(II) complex 1 exhibits two absorption bands in the visible region with maxima at 516 and 428 nm, which can be assigned to metal to ligand charge transfers (MLCTs). Such an electronic behavior has already been observed for homoletic iron-based compounds with pyridine-based ligands, and it can be explained by the first MLCT band at 428 nm involving the ligand subunit whose energy remained unchanged whatever the substitution and the second MLCT band at lower energy involving the central pyridine ring, which is sensitive to electronic tuning on this ring. In the case of Cu(II) complexes weak d−d bands at 685 and 677 nm for complexes 2 and 3, respectively, are responsible for the green color of the copper(II) complexes.
To examine the influence of the electrochromic properties of transition-metal complexes on polyazomethines, first the electrochemical and spectroelectrochemical measurements of complexes were carried out in solution. The electrochemical properties of the compounds were examined using cyclic voltammetry. The measurements were made in an anhydrous and deaerated 0.1 M solution of tetrabutylammonium hexafluorophosphate in acetonitrile as the supporting electrolyte.

Complex 1, which contains an Fe(II) ion was found to undergo the reversible oxidation/reduction process Fe(II) $\leftrightarrow$ Fe(III) with an anodic peak potential ($E_{pa}$) of +1.27 V and a cathodic peak potential ($E_{pc}$) of +1.17 V. In the case of complex 2, containing a Cu(II) ion, an irreversible reduction peak has been observed at the cathodic peak potential $E_{pc} = -0.41$ V associated with the reduction process Cu(II) $\rightarrow$ Cu(I). In the anodic scan a peak for the irreversible oxidation of Cu(I) ions into Cu(II) ions has been observed at an anodic peak potential of +0.33 V. 2 has been chosen as an example of a Cu(II)-based complex for the examination of electrochemical and electrochromic properties due to the more intense green color in comparison with 3 in solution at the same concentration of the complex.

Due to the irreversible character of oxidation/reduction of the metallic center in complexes of Cu(II) ions, the characterization of the redox process was carried out for an Fe(II) complex. It was done by recording cyclic voltammograms at different scan rates and examining the dependence of the peak current on the scan rate (Figure 5B). In case of complex 1 in solution it was found that both anodic and cathodic peak currents are linearly dependent on the square root of the scan rate with excellent linear correlation coefficients ($R^2$) of 0.9998 and 0.9995 for $i_{pa}$ and $i_{pc}$, respectively (Figure 5C). This indicates a reversible electron transfer process involving freely diffusing redox species.

The oxidation/reduction of metallic centers was accompanied by an intrinsic color change that was tracked by spectroelectrochemistry. It combines both electrochemistry and spectroscopy and allows us to track the color changes in situ. Complex 1 was found to change from red to yellow as a result of the electrooxidation of Fe(II) ions into Fe(III) ions (Figure 6A). This was accompanied by a gradual decrease of the MLCT bands at 516 and 428 nm and increase of the absorbance in the range of 350−400 nm that can be attributed to LMCT of the Fe(III) complex.41,42 The sharp isosbestic point at 402 nm confirms that only oxidized and reduced species are present in the solution and no side products of electrooxidation are formed. In case of Cu(II) complex 2 the observed color change was from green to red-brown as a result of the electroreduction Cu(II) $\rightarrow$ Cu(I). The color change was connected with the disappearance of the d−d band at 685 nm accompanied by an increase in the absorbance in the range below 600 nm (Figure 6B).

The electrochemical and spectroelectrochemical properties of polyazomethines were investigated using polymer-modified ITO-coated glass electrodes. Polymers P1 and P2 obtained by polycondensation of complexes of Fe(II) and Cu(II), respectively, with a triphenylamine-based dialdehyde were found to undergo oxidation/reduction processes of both metallic centers and the triphenylamine group (Figure 7).

The oxidation potentials of polymer P1 were found to be highly dependent on the scan rate. At a scan rate of 20 mV/s in the anodic part of the cyclic voltammogram an oxidation peak

![Figure 4. Absorption spectra of ligand L (■ black) measured in dichloromethane and complexes 1 (● red), 2 (▲ blue) and 3 (▼ magenta) in acetonitrile.](https://doi.org/10.1021/acs.inorgchem.1c01249)

![Figure 5. (A) Cyclic voltammetry of complexes 1 (red) and 2 (black) measured in anhydrous and deaerated 0.1 M solutions of TRAPF$_6$ in acetonitrile at a scan speed of 100 mV/s. (B) CV profiles of complex 1 at different scan rates. (C) Linear dependence of the peak currents on the square root of the scan rate for complex 1.](https://doi.org/10.1021/acs.inorgchem.1c01249)
TBAPF$_6$ in acetonitrile at a scan rate of 20 mV/s. Measured in an anhydrous and deaerated 0.1 M solution of TBAPF$_6$ in acetonitrile as the supporting electrolyte with applied potentials of 0 (black), 1.0 (red), 0.9 (black), 1.1 (magenta), 1.2 (green), and 1.3 V (navy) versus Ag/Ag$^+$. The linear behavior ($R^2 = 0.9944$) of the peak current as a function of the scan rates (Figure S27) indicates that the redox events occur at the electrode surface, demonstrating that the polymer was successfully attached to the ITO electrode. In the cyclic voltammogram of polymer P2 an irreversible oxidation wave at $E_{pa} = +0.49$ V connected with the electrooxidation of Cu(I) ions to Cu(II) ions and a quasi-reversible redox event with a half-wave potential of $E_{1/2} = +0.97$ V ($E_{pa} = +1.15$ V and $E_{pc} = +0.79$ V) associated with the redox reaction of triphenylamine group have been observed (Figure 7B). Additionally in the cathodic part of the voltammogram irreversible reduction of Cu(II) ions to Cu(I) ions at a potential of $-0.58$ V has been detected. The oxidation and reduction potentials of polymer P2 are shifted to more positive and more negative potentials, respectively, in comparison to complex 2, indicating that the metal centers in polymers are becoming hard to oxidize or reduce. This is consistent with an increase in the resistance of the film.

Polymers P3 and P4 exhibited redox peaks from electrochemical reactions of both the metallic centers and the thiophene group (Figure S28). An irreversible oxidation process of Fe(II) ions in polymer P3 was observed at $E_{pa} = +1.61$ V, whereas irreversible oxidation and reduction processes based on the redox reaction of Cu(II) ions in the case of polymer P4 were observed at $+0.85$ and $-0.62$ V, respectively. Additionally, irreversible oxidation waves at $E_{pa} = +1.99$ and $+2.12$ V for polymers P3 and P4, respectively, were observed that are consistent with oxidation of the thiophene moiety. In the case of polymers P5 and P6 obtained by the polycrystallization of transition-metal ions complexes with dialdehyde 6, which does not contain any electroactive group, we expected to obtain thin films that exhibit oxidation/reduction processes based on the redox reaction of transition-metal ions, but no oxidation/reduction peaks were observed in the investigated potential window. This was probably due to the high insulating character of the obtained polyazomethines.

Due to the best electrochemical properties of polymers P1 and P2 containing a triphenylamine group, they were chosen for further investigation of their electrochromic properties. The polymer P1 was found to undergo a color change from orange to gray as a result of electrooxidation. This was accompanied by a decrease in the MLCT band at 475 nm and the formation of a band below 400 nm, which was assigned to a ligand to metal charge transfer (LMCT) band of the Fe(III) complex, as well as the formation of new, broad band in the NIR region with a maximum at 930 nm (Figure 8), which was connected with the formation of a radical cation on the triphenylamine group in azomethines.

The isosbestic point at 580 nm confirms that only two forms, oxidized and neutral forms, are present, which indicates that oxidation of Fe(II) to Fe(III) is concomitant with the oxidation of Fe(II) ions. With an increase in the scan speed the shift of the first oxidation potential to more positive values has been observed and it overlaps with the second oxidation peak, and at a scan speed of 1000 mV/s it appears at a potential of $+1.7$ V (Figure S26). Similarly the reverse peak in the cathodic part of the cyclic voltammograms shifts toward more negative potential upon an increase in the scan rate from $+1.0$ V at a scan rate of 20 mV/s to $+0.5$ V at a scan rate of 1000 mV/s. Indeed, the peak to peak CV separation ($\Delta E_p$) increases with the scan rate, which indicates the presence of electrochemical irreversibility and a quasi-reversible redox process; this is because the peak potentials of the forward and backward scans are always shifted by a non-negligible voltage value, while a true reversible process should feature the same (reduction and oxidation) potential values. The linear behavior ($R^2 = 0.9944$) of the peak current as a function of the scan rates (Figure S27) indicates that the redox events occur at the electrode surface, demonstrating that the polymer was successfully attached to the ITO electrode.
reversible, and after application of a slightly negative potential (−0.1 V), neutralization of the triphenylamine group and reduction of Fe(III) to Fe(II) occur, which resulted in the restoration of the initial UV−vis−NIR spectra.

To investigate the time that is required for the polymer to change in color from orange to gray, switching between +1.5 and −0.1 V was carried out 10 times in 30 s intervals to ensure the completion of the electrochromic reaction and the changes in transmittance were monitored at the absorption maximum of the oxidized state at 930 nm (Figure S29). The switching times, coloration ($T_{c,90}$) and bleaching ($T_{b,90}$) times, were calculated as the times required to reach 90% of the final change in transmittance between neutral and oxidized states (Figure 9A), and they were found to be 9.6 and 6.4 s for the coloring and bleaching steps, respectively.

The maximum transmittance difference between the oxidized and reduced states was calculated to be 31%, and it did not change after 10 switching cycles. Coloration efficiency is an important characteristic for electrochromic materials, and it is a proportionality factor between the optical absorbance change of an electrochrome at a designated wavelength (optical density ($\Delta OD$)) and the density of injected/ejected electrochemical charge ($Q_d$) necessary to induce a full color change. A quantitative calculation of the coloration efficiency of materials can be done from a plot of the in situ optical density at an appropriate wavelength versus the inserted/extracted charge density. A plot of the optical density at 930 nm versus charge density for polymer P1 is shown in Figure 9B. The coloration efficiency value was extracted as the slope of the line fitted to the linear region of the curve and it was...
found to be 136.5 cm$^2$/C, which is similar to values for other triphenylamine-based polymers obtained via a polycondensation method.$^{51}$

To investigate the stability of the film as a function of time, multiple switching between orange and gray colors was investigated in 10 s intervals (Figure 9C). The 10 s interval was chosen to obtain a high enough color contrast and, due to the fact that the measurement was done using an acetonitrile-based electrolyte, to shorten the duration of the measurement to avoid evaporation of the solvent. It was found that the polymer P1 was quite stable during ∼275 switching cycles and the transmittance difference dropped by only ∼3.5% from 30.9% to 27.4% but then decreased to 23% when ∼310 switching cycles were carried out.

In the case of polymer P2 the color change from yellow to gray-blue was observed as a result of oxidation of the triphenylamine group followed by a color change to red as a result of neutralization of the radical cation on the triphenylamine group and reduction of Cu(II) ions to Cu(I) ions (Figure S30). The color changes, although visible by the naked eye, were found to be of very low intensity in terms of differences in both absorbance and transmittance (Figures S30 and S31). Additionally, the polymer was found to exhibit low stability during multiple oxidation/reduction cycles and it delaminated from the surface during 10 oxidation/reduction cycles (Figure S31). This was probably due to the necessity of changes in the conformation of the coordination subunits caused by a change in the coordination preferences of the metallic centers. In the case of polymer P3 a color change from red to orange has been observed (Figure S32), whereas the polymer P4 exhibited a color change from yellow to brown after electroreduction (Figure S33). The color changes of polymers P3 and P4 were both found to be irreversible.

To investigate the influence of transition-metal ions on the electrochromic properties of the materials, we attempted to prepare thin films of the investigated ligands using the on-substrate polymerization method. It was found that the material obtained by polycondensation of ligand L and dialdehyde 4 exhibited reversible oxidation/reduction properties with the anodic and cathodic potentials $E_{pa} = +1.13$ V and $E_{pc} = +0.72$ V, respectively (Figure S34A). In case of the material obtained by polycondensation of ligand L and dialdehyde 5, an irreversible oxidation at $E_{pa} = +1.99$ V was observed (Figure S34B), while the polycondensation of L with dialdehyde 6 gave an electrochemically inactive material. The electrochemically induced color change of the thin film was observed only in the case of a material containing a triphenylamine group (Figure S35). The material in its neutral state was yellow with an absorption maxima at 378 nm, and after oxidation of the triphenylamine group, material turned blue, which was concomitant with the appearance of a new, broad absorption band at 700 nm. This indicates that the coordination with metal ions changes the optical properties of the materials.

The surface morphology of the layers of polymers P1 and P2 was investigated using scanning electron microscopy (SEM) as well as atomic force electron microscopy (AFM).

As seen in Figure 10, the polymers do not form a continuous and uniform layer on the surface; instead, the ITO surface is covered by small droplets of the polymers. This is the result of the coating method that was applied. Due to spray-coating of the mixture of monomers onto ITO substrates, small droplets of the mixture of monomers formed and then polymerization occurred within the droplets. During the SEM observation, in order to determine the composition of a certain part of the polymer, an EDX analysis was performed to obtain the elemental composition. EDX spectra of polymers P1 and P2 are shown in Figure S36 and S37, and they confirm that polymers P1 and P2 contain the transition-metal ions iron and copper, respectively, in the structure, which has also been confirmed by an XPS analysis. Due to the nonuniform nature of the layers, it was difficult to measure their average thickness. The thickness of the islands of polymers was investigated using AFM imaging (Figures S38 and S39). It was found that the thickness of selected islands is as much as 1 µm for polymer P1 and as much as 2 µm for P2.

## CONCLUSIONS

In summary, we have prepared the series of polyazomethines P1–P6 containing complexes of transition-metal ions by polycondensation reactions of tetraamines bearing complexes of transition-metal ions with the 2,6-bis(1-hydrazonoethyl)-pyridine ligand L and different dialdehydes and investigated them with respect to their performance as electrochromic materials. The polymers were obtained by on-substrate polymerization methods, which allowed for the production of polymers as insoluble thin layers on a transparent ITO electrode. Electrochemical and spectrophotometric studies reveal that the most interesting properties are exhibited by polymers P1 and P2, obtained by the polycondensation of complexes of both Fe(II) and Cu(II) ions, respectively, with
the triphenylamine-based dialdehyde. Polyazomethine P1 containing Fe(II) ions exhibited a color change from orange to gray, while the Cu(II)-based polyazomethine P2 changed a color from yellow to blue as a result of electrooxidation and from blue to red after electroreduction of the metallic center. Additionally, polymer P1 exhibited good electrochemical stability during ~300 oxidation/reduction cycles, switching times of 9.6 and 6.4 s for the coloring and bleaching steps, respectively, and a coloration efficiency of 136.5 cm²/C, which makes it a promising material for electrochromic applications. The research conducted also showed the usefulness of an oun-substrate polycrystallization reaction for the preparation of thin films of electrochromic metallopolymers.

■ ASSOCIATED CONTENT

Supporting Information
The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.inorgchem.1c01249.

Additional figures and a table as described in the text (PDF)

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CCDC 2075985–2075987 contain the supplementary crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/data_request/cif, or by emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223 336033.

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