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Fabrication of an Active Electronic Device Using a Hetero-bimetallic Coordination Polymer

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Supporting Information

ABSTRACT: A nickel(II)/lead(II) coordination polymer [(NCS)Pb(H2O)-LNi(NCS)]n {H2L = N,N'-bis(3-methoxysalicylidene)propane-1,3-diamine} has been synthesized and characterized. The band gap (3.18 eV) calculated from Tauc’s plot suggests the semiconducting nature of the complex. The material has a photosensitivity of 5.76, indicating its applicability in the fabrication of photosensitive devices. The complex has been successfully applied in a technologically challenging thin-film photosensitive Schottky device.

INTRODUCTION

Coordination polymers (CPs) have attracted the interest of coordination chemists for the last few years for their application in gas absorption, magnetic material, catalysis, luminescence, drug delivery, sensor technology, etc.1,2 Although application of some organic materials in light-emitting diodes3 and field effect transistors4 is known, fabrication of an active electronic device using CPs is not well explored till date. Some d10 metals have the ability to increase the intensity and shift the wavelength of emission of organic ligands by metal coordination, and therefore d10 metal-based CPs were used in optoelectronic devices.5 There are reports in the literature regarding the fascinating application of some semiconducting CPs in photovoltaics6 and other optoelectronic devices.7 N2O2O2− donor compartmental Schiff base ligands have been widely used for the preparation of hetero-dinuclear complexes.8 These could then be linked using suitable pseudohalides [NCS−, N3−, NCO−, and N(CN)2−] to generate various CPs.9 Although any member of the first transition series is a good choice to be placed in the inner N2O2− compartment of such ligands, nickel(II) is preferred for its ability and flexibility to bind both hard and soft donor centers with equal ease and to adopt various geometries with different coordination numbers, oxidation states, and its applicability extending from organometallic to biological chemistry.10 On the other hand, the coordination chemistry of lead(II) is also interesting for its potential to form hemi- as well as holo-directed complexes with coordination numbers ranging from 2 to 10.11 However, it must be remembered that lead is hazardous to pollution and health12 and must be handled with great care. Any semiconducting material showing rectifying nature in a metal–semiconductor (MS) junction can be applied in the Schottky device.13 The nonlinear rectifying behavior of I−V characteristics of that material-based device measured under dark and illumination conditions proves its photosensitive Schottky diode character.14 However, to the best of our knowledge, there is no report in the literature to use any X-ray characterized hetero-bimetallic CP for the fabrication of any optoelectronic devices till date. In this work, a compartmental Schiff base ligand has been used to synthesize a nickel(II) complex, which could then be used as a metalloligand to prepare a hetero-bimetallic nickel(II)/lead(II) building block, which in turn, is linked via thiocyanate to form a coordination polymer (Scheme 1). The band gap (3.18 eV) has been calculated from Tauc’s plot, and it suggests the semiconducting nature of the complex. The rectifying nature in I−V characteristics hints toward its Schottky diode characteristics. The on−off ratio (70) of the semiconductor device fabricated with CP is high. This is eventually the first report of any hetero-bimetallic coordination polymer with photosensitivity, which could be used in the fabrication of a Schottky diode.

EXPERIMENTAL SECTION

Nickel(II) thiocyanate tetrahydrate has been prepared by us following the literature method.15 Other reagents and solvents...
geometrically idealized positions and constrained to ride on maps. All other hydrogen atoms were placed in their attached with oxygen atoms were located by dihydrogen atoms were re−}

...spectrum in KBr (4500−12789 cm −1 ). FT−IR (KBr, cm −1 ): C, 34.06; H, 2.99; N, 7.57; found: C, 33.9; H, 2.8; N, 7.7%. UV−vis [ε max (L mol −1 cm −1 )] (DMF): 271 (1.7 × 10 3 ); 352 (8.6 × 10 3 ); 584 (9.5); 814 (74.1).

Physical Measurements. C, H, and N analysis was performed using a PerkinElmer 240C elemental analyzer. IR spectrum in KBr (4500−500 cm −1 ) was recorded with a PerkinElmer Spectrum Two spectrophotometer. Electronic spectrum of the complex in N,N-dimethylformamide (DMF) was recorded on a PerkinElmer Lambda 35 UV−visible spectrophotometer. Steady-state photoluminescence spectrum of the complex in DMF was obtained in Shimadzu RF−5301PC spectrofluorometer at room temperature. Time−dependent photoluminescence spectrum was recorded using a Hamamatsu MCP photomultiplier (R3809) and was analyzed by using IBHDAS6 software. X−ray diffraction of the powdered sample was performed on a Bruker D8 instrument with Cu Kα radiation (λ = 1.5406 Å).

X−ray Crystallography. A suitable single crystal of the complex was picked and mounted on a glass fiber, and diffraction intensities were measured with an Oxford Diffraction X−Calibur diffractometer equipped with Mo Kα radiation (λ = 0.71073 Å, 50 kV, 40 mA) at 150 K. Data collection and reduction were performed with the Crysalis software. The structure of the complex was solved by direct methods using SHELXS−97 and refined by full−matrix least squares on F 2 , using the SHELXL−2016 program. Non−hydrogen atoms were refined anisotropically. Hydrogen atoms attached with oxygen atoms were located by difference Fourier maps. All other hydrogen atoms were placed in their geometrically idealized positions and constrained to ride on their parent atoms. Programs used included PLATON, DIAMOND, ORTEP, and MERCURY.

Device Fabrication. Electrical characterization was performed in the complex-based metal−semiconductor (MS) junction devices. These devices were fabricated by depositing a thin film of a well−dispersed solution of the synthesized complex. To develop the thin films, indium tin oxide (ITO)−coated glass substrates were cleaned by acetone, ethanol, and distilled water with the help of an ultrasonicator. Thereafter, the complex was separated mixed with N,N-dimethylformamide (DMF) in the right proportion (20 mg mL −1 ) and sonicated for several minutes until it produces a well−stabilized dispersion. Then, on the top of the cleaned ITO-coated substrates, stable dispersion of the just prepared complex was spun first at 600 rpm for 4 min and thereafter at 1000 rpm for 6 min with the help of SCU 2700 spin coating unit. To evaporate the solvent, as−deposited thin films were annealed under vacuum at 80 °C for 30 min. For the characterization of the developed thin film, thickness was measured by a surface profiler as ∼1 μm. Pure Al (99.99%) was chosen as the metal electrode and was deposited on to the active thin film to prepare the metal−semiconductor (MS) interface through a shadow mask by thermal evaporation technique with the help of a vacuum−coating unit under a base pressure of 10 −6 Torr. The effective areas of the films were maintained as 7.065 × 10 −2 cm 2 . For electrical characterization of the devices, the current−voltage (I−V) characteristics were recorded under dark and AM 1.5G radiation with the help of a Keithley 2635B source meter by a two−probe technique. All preparation and measurements were performed at room temperature and under ambient conditions.

RESULTS AND DISCUSSION

Synthesis. H 2 L is a N 2 O 2 donor Schiff base ligand, and it has been prepared by refluxing 1,3-diaminopropane and 3-methoxysalicylaldehyde in methanol following the literature method. The ligand (H 2 L) on reaction with lead(II) nitrate followed by the addition of nickel(II) thiocyanate tetrahydrate formed the complex. Formation of the complex has been shown in Scheme 1.

Structure Description. [(NCS)Pb(H 2 O)LNi(NCS)] n. The complex crystallizes in monoclinic Space group P2 1/n. Selected crystallographic data have been summarized in Table 1. Significant bond lengths and bond angles have been gathered in Tables 2 and S1 (Supporting Information), respectively.

The complex consists of a chain system with the repeating hetero-dinuclear neutral units, [(NCS)Pb(H 2 O)LNi(NCS)] n, joined through two μ 1-thiocyanate bridge. In each bimetallic unit, the nickel(II) and lead(II) centers, respectively, occupy

Scheme 1. Schematic Representation for the Synthetic Route to the Complex
Table 1. Crystal Data and Refinement Details

| parameter                  | value                  |
|----------------------------|------------------------|
| formula                    | C₃₂H₂₂NiO₅PbS₂         |
| formula weight             | 740.46                 |
| temperature (K)            | 150                    |
| crystal system             | monoclinic             |
| space group                | P2₁/n                  |
| a (Å)                      | 12.3716(6)             |
| b (Å)                      | 9.6865(3)              |
| c (Å)                      | 20.7826(13)            |
| β (°)                      | 105.941(6)             |
| Z                          | 4                      |
| d_{ab} (g cm⁻¹)            | 2.054                  |
| μ (mm⁻¹)                   | 8.022                  |
| F (3000)                   | 1432                   |
| total reflections          | 15388                  |
| unique reflections         | 6823                   |
| observed data [I > 2σ (I)]| 5172                   |
| no. of parameters          | 326                    |
| R (int)                    | 0.051                  |
| R_f, wR_f (all data)       | 0.0798, 0.0987         |
| R_f, wR_f [I > 2σ (I)]    | 0.0538, 0.0916         |

Table 2. Selected Bond Lengths (Å) of the Complex

| bond                     | length (Å) |
|--------------------------|------------|
| Ni(1)–O(31)              | 2.034(4)   |
| Ni(1)–N(23)              | 2.030(5)   |
| Ni(1)–N(19)              | 2.029(6)   |
| Ni(1)–O(11)              | 2.039(4)   |
| Ni(1)–N(2)               | 2.154(6)   |
| Ni(1)–N(1)               | 2.112(5)   |
| Pb(1)–O(11)              | 2.328(4)   |
| Pb(1)–O(131)             | 2.655(6)   |
| Pb(1)–O(31)              | 2.310(6)   |
| Pb(1)–O(31)              | 2.645(6)   |
| Pb(1)–S(1)               | 2.867(3)   |
| Pb(1)–O(1)               | 2.703(7)   |

Note: Symmetry transformations = 1/2 - x, -1/2 + y, 1/2 - z.

The inner N₂O₃ and outer O₂O' sites of the potential N,O,O', donor deprotonated compartmental Schiff base, L⁺⁻. A perspective view of the asymmetric unit of the complex with selective atom numbering scheme is depicted in Figure 1. Both nickel(II) and lead(II) have pseudo octahedral geometries. Ni(1) is coordinated by two imine nitrogen atoms, N(19) and N(23), and two phenoxo oxygen atoms, O(11) and O(31), of the deprotonated potential N₂O₃ donor compartmental Schiff base ligand (L)⁺⁻, which constitute the equatorial plane. The deviation of the coordinating atoms N(19), N(23), O(11), and O(31) from the least-square mean plane through them is -0.037(5), 0.038(5), 0.044(4), and -0.043(4) Å, respectively, and that of Ni(1) from the same plane is 0.0025(7) Å. The fifth coordination site is occupied by one nitrogen atom, N(2), from a thiocyanate anion. A nitrogen atom, N(1)⁺, from a symmetry-related (1/2 - x, -1/2 + y, 1/2 - z) thiocyanate anion coordinates nickel(II) in its sixth coordination site to form a distorted octahedron. On the other hand, lead(II) is coordinated by two phenoxo oxygen atoms, O(11) and O(31), and two ethoxy oxygen atoms, O(131) and O(291), which constitute the equatorial plane. The fifth coordination site is occupied by an oxygen atom, O(1), from a water molecule. A sulfur atom, S(1), from an end-to-end bridged thiocyanate coordinates with lead(II) in its sixth coordination site to complete its distorted octahedral geometry. Nickel(II) and lead(II) centers are bridged by two phenoxo oxygen atoms, O(11) and O(31), with Ni(1)−Pb(1) distance being 3.458(7) Å. The bridging angles, Ni(1)−O(11)−Pb(1) and Ni(1)−O(31)−Pb(1), are 104.5(2) and 105.3(2)°, respectively. The Ni(1)−O(11)−O(31)−Pb(1) core is almost planar (the dihedral angle is close to ~5.32°).

Supramolecular Interaction. The complex forms an interesting supramolecular structure via π···π interactions. π···π interactions have been observed between the phenyl ring C(12)−C(13)−C(14)−C(15)−C(16)−C(17) and symmetry-related (−x, −y, −z) phenyl ring C(12)−C(13)−C(14)−C(15)−C(16)−C(17). These interactions form a two-dimensional (2D) structure (Figure 2). Details of the geometric features of π···π interactions have been given in Table 3.

Morphological Analysis. Microstructure of the complex has been studied by field emission scanning electron microscope (FESEM) and presented in Figure 3. The micrograph states featherlike morphology of the complex.

Optical Analysis. The Tauc plot (the details have been given in the Supporting Information) of the material has been presented in Figure 4. From Tauc’s plot, the direct energy band gap (E₀) of the material has been estimated as 3.18 eV by measuring the x-intercept of the extrapolated linear part of the plot (hν)² vs (hν). The analysis for taking direct band gap has been provided in the Supporting Information.

As the band gap suggests that the complex is semiconducting in nature, there is a possibility to apply the complex in a

Figure 1. (a) Perspective view of the complex (asymmetric unit) with a selective atom numbering scheme; (b) the polymeric structure of the complex. Only the relevant atoms have been shown for clarity in both cases.
semiconducting device. But there has always been technological difficulty in application of these materials in active devices. Thus, to check the possibility of device application, fabrication of a metal–semiconductor-based thin-film Schottky diode has been tried. As the electronic charge transport properties determine the performance of the devices, it is very important to analyze charge transport properties in detail.

**Electrical Characterization.** The current–voltage ($I$–$V$) measurements of the device [schematic diagram (Figure S2) has been given in the Supporting Information] fabricated with CP have been done in dark and under AM 1.5G light illumination in the voltage range of $-2$ to $+2$ V. Experimentally obtained $I$–$V$ curves of the device under dark and illumination are shown in Figure 5. The inset in Figure 5 represents the curve in log scale.

The nonlinear $I$–$V$ curve of the device under both the conditions indicates that the fabricated metal–semiconductor junction is a Schottky junction. From the $I$–$V$ curve, rectification ratios have been calculated, which are 70 and 89 under dark and illumination condition, respectively. This proves the successful application of the complex material in an active electronic device. The estimated conductivity of the thin film has been increased from $2.02 \times 10^{-4}$ Sc m$^{-1}$ under dark to $8.55 \times 10^{-4}$ Sc m$^{-1}$ under light. The conductivity data for the crystal has also been provided in the Supporting Information.

To assess the potentiality of the fabricated device for application in a photodetector, we have estimated important parameters like photosensitivity ($S = I_{ph}/I_D$ where $I_{ph}$ is the photocurrent and $I_D$ is the dark current $I_{dark}$), specific detectivity ($D^*$), and responsivity ($R$). Photosensitivity of our device is 5.76, and this has been compared with the other reported devices and presented in Table 4.23

Responsivity ($R$) of the device can be estimated from the following equation

$$R = \frac{I_{ph}}{P_{in}A}$$

Here, $A$ is the effective area of the diode and $P_{in}$ is the incident power of the light. For our device, the responsivity is 1.69 A W$^{-1}$, which is significant for this kind of device. The specific detectivity ($D^*$) has been estimated from the equation

$$D^* = \frac{R}{(2qI_D)^{1/2}}$$

where $q$ is the charge of electron. The specific detectivity of the device has been estimated as $6.55 \times 10^{10}$ Jones, which is again
significant. All device parameters indicate that the fabricated device has a great potential in the field of active electronic device.

Further analysis of the different device parameters has been done by thermionic emission theory given by the following equation

\[ I = I_0 \exp\left( \frac{qV}{nKT} \right) - 1 \]

Here, the forward current is represented by \( I \), reverse saturation current by \( I_0 \), \( V \) denotes the applied bias voltage, \( q \) is the electronic charge, \( k \) represents the Boltzmann constant, \( T \) gives the absolute temperature, and \( n \) denotes the ideality factor (IF). \( I_0 \) is expressed by

\[ I_0 = A \exp\left( -\frac{q\phi_B}{kT} \right) \]

The effective area of the diode (A) is taken to be \( 7.065 \times 10^{-2} \) cm\(^2\). \( A^* \) denotes the Richardson constant whose value is 32 A K\(^{-2}\) cm\(^{-1}\), and \( \phi_B \) is the Schottky barrier height.

By using following Cheung’s equations\(^{23}\) given below, other different Schottky parameters have also been calculated

\[ \frac{dV}{d \ln I} = \left( \frac{\eta kT}{q} \right) + IR_S \]

\[ H(I) = V - \frac{\eta kT}{q} \ln \left( \frac{I}{AA^*T^\gamma} \right) \]

\[ H(I) = IR_S + \eta\phi_B \]

Here, the series resistance is given by \( R_S \) and all other notations are kept same.

The intercept of the \( dV/d \ln I \) vs \( I \) plot shown in Figure 6 is used to estimate the ideality factor. Determination of the barrier potential height and the series resistance is done using the intercept and slope of \( H(I) \) vs \( I \) plot given in Figure 7. The various parameters calculated are enlisted in Table 5.

The ideality factor should be unity, but here it is less than 1, which can be due to the presence of inhomogeneities at the Schottky junction.\(^{25}\)

For the assessment of charge transport behavior of the device, mobility and diffusion length of charge carriers are the two important parameters. So the charge transport mechanism has been analyzed through the metal–semiconductor contact. Here, we have estimated mobility (\( \mu_{al} \)), lifetime (\( \tau \)), carrier concentration (\( N \)), and diffusion length (\( L_D \)) by applying space charge limited current (SCLC) theory. Figure 8 shows the \( I-V \) curve in a double logarithmic scale. The graph presented in Figure 8 can be divided into three regions with different slopes. Region 1 (Reg 1), which represents the lower bias, follows ohmic nature with current being directly proportional to voltage. The origin of the current is from the bulk generated electrons of the film, rather than the injected free carriers. Here, tunneling is the main contributing factor that can be observed from the \( I-V \) curve.\(^{26}\)

In Figure 8, the slope of Reg 2 is increased compared with Reg 1. Within the material, a part of the injected carriers is trapped within the traps present and a part remains free. With the increase of applied voltage, traps within the material are filled with a higher number of injected carriers.\(^{27}\) As the current increases rapidly, the Fermi level (\( E_F \)) moves through a trap level.\(^{28}\) The slope of this region increases sharply as the traps are being filled up rapidly by the carriers. The second region is known as the trap filling region.\(^{28}\) When almost all traps are filled, the slope of the third region decreases as result of slower carrier movement. Here, current follows the relationship \( I \propto V^2 \), which is the feature of trap-free space charge limited current (SCLC) regime.\(^{27,28}\)

So, effective carrier mobility (\( \mu_{al} \)) has been estimated from \( I \) vs \( V^2 \) graph (Figure 9) of the trap-free SCLC region using the Mott–Gurney equation:\(^{29}\) given by

\[ \frac{dV}{d \ln I} = \left( \frac{\eta kT}{q} \right) + IR_S \]

\[ H(I) = V - \frac{\eta kT}{q} \ln \left( \frac{I}{AA^*T^\gamma} \right) \]

\[ H(I) = IR_S + \eta\phi_B \]

Table 4. Previously Reported X-ray Characterized Photosensitive Complexes\(^{44}\)

| complex | conductivity in dark | conductivity under light | photosensitivity | ref |
|---------|----------------------|--------------------------|------------------|----|
| [CdL\(^{(1)}\)(μ\(_{1,1,3\text{-NCS}}\))\(_{3}\)]\(_{2}\) | 1.01 × 10\(^{-8}\) | 2.16 × 10\(^{-8}\) | 14.36 | 23a |
| [Cu\(_{6,4}\)(4-pic)\(_{2}\)(H\(_{2}\)O)\(_{2}\)]\(_{2}\)[ClO\(_{4}\)] | 8.21 × 10\(^{-8}\) | 11.84 × 10\(^{-8}\) | 1.83 | 23b |
| [Zn(acd)(4-spy)\(_{2}\)(H\(_{2}\)O)\(_{2}\)] | 5.12 × 10\(^{-8}\) | 16.48 × 10\(^{-8}\) | 1.96–2 | 23c |
| [Cd(acd)(4-spy)\(_{2}\)(H\(_{2}\)O)\(_{2}\)] | 6.54 × 10\(^{-8}\) | 28.77 × 10\(^{-8}\) | 3.5 | 23c |
| [Cd(2,2′-dsb)(4-nvp)(DMF)(H\(_{2}\)O)] | 6.60 × 10\(^{-8}\) | 10.71 × 10\(^{-8}\) | 0.54 | 23d |
| [Cu\(_{6,4}\)(μ\(_{1,1,3\text{-NCS}}\))\(_{2}\)] | 3.63 × 10\(^{-8}\) | 4.13 × 10\(^{-8}\) | 2.83 | 23e |
| [Cu\(_{6,4}\)(μ\(_{1,1,3\text{-NCS}}\))[HgCl(μ\(_{1,1,3\text{-NCS}}\))] | 1.48 × 10\(^{-8}\) | 8.40 × 10\(^{-8}\) | 57 | 23f |
| [(NCS)Pb(H\(_{2}\)O)\(_{4}\)](NCS)\(_{2}\) | 6.02 × 10\(^{-8}\) | 3.47 × 10\(^{-8}\) | 5.76 | this work |

\(^{44}\)Where HL\(^{1}\) = 2-(2-(ethylamino)ethyliminomethyl)-6-ethoxyphenol, H\(_{2}\)acd = acetylenedicarboxylic acid, 4-pic = 4-picoline, 4-spy = 4-styrylpyridine, H\(_{2}\)2,2′-dsb = 2,2′-disulfanediyldibenzoic acid (2,2′-dsb), and 4-nvp = 4-(1-naphthylvinyl)pyridine.
\[ I = \frac{9q_{\text{eff}}\varepsilon_0\varepsilon_r AV^2}{8d^3} \]

where \( I \) is the current, \( \varepsilon_0 \) is the permittivity of free space, \( \varepsilon_r \) is the dielectric constant, and \( d \) is the thickness of the film (about 1 \( \mu m \)).

There are few more important charge transport parameters like carrier concentration and transit time, which determine the device performance. The carrier concentration \( (N) \), transit time \( (\tau) \), and diffusion length \( (L_D) \) is estimated with the following set of equations\(^3^0\)

\[ N = \frac{\sigma}{q\mu_{\text{eff}}} \]

\[ \tau = \frac{9\varepsilon_0\varepsilon_r A}{8d} \left( \frac{V}{I} \right) \]

where \( D \) is the diffusion coefficient. The diffusion coefficient has been calculated by employing the Einstein–Smoluchowski equation\(^3^1\)

\[ \mu_{\text{eff}} = \frac{qD}{kT} \]

By analyzing the charge transport parameters, it is seen that the diffusion length and carrier mobility increased and transit time decreased after light illumination, which indicates that the device could be very useful for photosensitive device applications.

\section*{CONCLUSIONS}

Although the nonlinear rectifying behavior of \( I-V \) characteristics of few semiconducting material-based devices measured under dark and illumination conditions proves their photo-

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**Table 5. Different Charge Transport and Device-Related Parameters**

| Conditions  | On/off Ratio | \( \sigma \) (S cm\(^{-1}\)) | \( \frac{dV}{d\ln I} \) | \( H(I) \) | \( \Phi_B \) (eV) | \( \mu_{\text{eff}} \) (cm\(^2\) V\(^{-1}\) s\(^{-1}\)) | \( N \) (m\(^{-3}\)) \times 10\(^{23}\) | \( \tau \) (\( \mu s \)) | \( D \) (m\(^2\) s\(^{-1}\)) \times 10\(^{-8}\) | \( L_D \) (nm) |
|------------|-------------|-----------------|-----------------|---------|------------|-----------------|----------------|--------|----------------|--------|
| Dark       | 70          | \( 2.02 \times 10^{-4} \) | 0.52            | 0.94    | 1.07       | 0.71            | 9.46 \times 10^{-3} | 1.33   | 0.35           | 8.85   | 248       |
| Light      | 89          | \( 8.55 \times 10^{-4} \) | 0.79            | 0.80    | 0.86       | 0.55            | 3.42 \times 10^{-2} | 1.56   | 0.11           | 2.45   | 273       |

**Figure 6.** \( \frac{dV}{d\ln I} \) vs current plot in dark and light conditions.

**Figure 7.** \( H \) vs current graph in dark and light conditions.
sensitive Schottky diode character, there was no report in the literature to use any X-ray-characterized hetero-bimetallic coordination polymer for the fabrication of any optoelectronic device before the present work. In this article, synthesis, X-ray structure, and electronic properties of a hetero-bimetallic nickel(II)/lead(II) CP has been described. The band gap (3.18 eV) of the material indicates that it belongs to the semiconductor family. Intermolecular interactions play a key role for molecular conductors and semiconductors. The present complex forms a one-dimensional (supramolecular) structure via coordinate bond. This coordinate bond formation helps the complex to show very high conductivity. The complex has been successfully applied in technologically challenging thin-film Schottky diodes with appraised device related parameters. The results indicate that the material can be applied to photosensitive active devices. The preparation of the complex thus illustrates a potentially versatile approach to the construction of hetero-bimetallic coordination polymers and their potential application in the field of optoelectronics.

We are still working in the laboratory to get better yield of the product and to develop and characterize more systems based on this strategy to generalize the concept.
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