ABSTRACT

Diarylethene-containing ligands and their coordination compounds are described. The ligands display photochromism with UV excitation, while the coordination compounds display photochromism with both excitation in the UV region and excitation into lower energy absorption bands characteristic of the coordination compounds, through which the excitation wavelengths for the photocyclization can be extended from λ≤340 nm to wavelengths beyond 470 nm. Switching of the luminescence properties of the compounds has also been achieved through photochromic reactions.
Fig. 2

![Graph 1](image1)

Wavelength / nm

Fig. 3

![Graph 2](image2)

Wavelength / nm
Fig. 4

(a) 

open form  \[ \xrightarrow{313 \text{ nm}} \] close form  \[ \xrightarrow{365, 540 \text{ nm}} \]

(b) 

open form  \[ \xrightarrow{313, 440 \text{ nm}} \] close form  \[ \xrightarrow{365, 540 \text{ nm}} \]

Fig. 5

Normalized Emission Intensity

Wavelength / nm
Fig. 8

\[ \text{Pd(PPh}_3\text{)}_4 \text{, THF / 2M Na}_2\text{CO}_3\text{(aq)} \]

\[ \begin{align*}
  \text{2.5} & \quad \text{S(OH)}_2 \quad \text{Br} \quad \text{N} \quad \text{Br} \quad \text{N} \\
  \text{R} & \quad \text{Ph} \quad \text{Me} \quad \text{Me} \\
\end{align*} \]

\[ \text{R} = \quad \text{Ph} \quad \text{OMe} \quad \text{(L2)} \quad \text{Ph} \quad \text{Me} \quad \text{(L3)} \]

Fig. 9

(a) Absorbance vs. Wavelength / nm (b) Normalized Emission Intensity
Fig. 10

\[ \text{2.5} \quad \text{B(OH)}_2 \quad + \quad \text{Br-N=H} \quad \xrightarrow{\text{Pd(PPh)}_3_4} \quad \text{THF / 2M Na}_2\text{CO}_3(aq) \quad \rightarrow \quad \text{product} \]

\[ \text{MeI, NH}_4\text{PF}_6 \quad \text{MeCN / K}_2\text{CO}_3 \quad \rightarrow \quad \text{product} \quad (1.4) \]

\[ \text{Pd(OAc)}_2, \text{KO'Bu} \quad \text{NaI / THF} \quad \rightarrow \quad \text{product} \quad (7) \]

Fig. 11

[Graph showing absorbance vs. wavelength]
PHOTOCHROMIC DIAREYLETHENE CONTAINING COORDINATION COMPOUNDS AND THE PRODUCTION THEREOF

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application is a continuation-in-part of co-pending U.S. patent application Ser. No. 10/883,677 filed Jul. 6, 2004, which claims the benefit of U.S. provisional patent application No. 60/484,668 filed Jul. 7, 2003, the entirety of both disclosures being incorporated herein by reference.

FIELD OF THE INVENTION

[0002] This invention is related to the design and the photochromatic behavior of novel photochromic ligands and their coordination compounds. The design of these photochromic ligands and their coordination compounds is based on the cis-diarylethene structure, which forms part of a mono- or poly-cyclic ring structure that contains one or more donor atom(s) or donor heteromatom(s) for coordination to an acceptor atom to form photochromic coordination compounds.

BACKGROUND OF THE INVENTION

[0003] Photochromism is defined as “a reversible transformation of a single chemical species being induced in one or both directions by absorption of electromagnetic radiation, with two states having different distinguishable absorption spectra.” Thus, photochromic compounds are compounds that possess at least two isomeric forms which have different physical properties, such as absorption properties, refractivity, and the like, and can be transformed from one form to another by light excitations at prescribed wavelengths.

[0004] Photochromism has been intensively studied due to its potential use for optical recording and other optical functioning devices. To be practically used as optical recording materials, both isomeric forms must be thermally stable and possess excellent durability for reversible photochromic reactivity. Diarylethene is one class of photochromic compounds which possesses these properties, and therefore is a suitable class of compounds for the construction of optical functioning devices. The cis-configuration of both aryl groups in the diarylethenes studied is generally fixed by an upper cycloalkene structure, such as fluorinated alicyclic group, aromatic group, anhydride and maleimide group. Apart from the difference in absorption characteristics and the like between the two forms and their thermal stabilities, the availability of desirable excitation wavelengths that can be tuned and selected for the photochromic reactions also represents an important aspect in the design of materials for optical functioning devices. It has been shown that with the more n-conjugated upper cycloalkene structures, such as maleimide derivatives, in the diarylethene compounds, the photoacyclization proceeded with lower energy excitation in the visible region.


SUMMARY OF THE INVENTION

[0006] The present invention relates to the use of coordination compounds to perturb the properties of the diarylethenes in photochromic compounds. Described below is a report of the design, synthesis and studies of cis-diarylethene-containing ligands, with the upper cycloalkene being part of a mono- or poly-cyclic ring structure that contains one or more donor atom(s) or donor heteromatom(s), such as phenanthrolines, pyridines, imidazoles, pyrazoles, thiazoles, pyroles, diazines, triazines, polypyridines, porphyrins and phthalocyanines and the like, for coordination compound formation.

[0007] The object of the present invention is to provide a new class of diarylethene-containing coordination compounds capable of displaying perturbed and sensitized photochromic properties. The invented photochromic compound is a coordination compound that contains a diarylethene with one or more donor atoms coordinated to an acceptor atom of the coordination compound. Any diarylethene in which the ethene group in a heterocyclic moiety, monocyclic or poly-cyclic, with any donor atom(s) capable of forming a coordination compound can be used in the present invention. There is no restriction on the nature of the aryl groups and they can be heteroaryl groups such as, for instance, thienyl groups. Likewise, any acceptor atoms which can be coordinated with the ethene-containing heterocyclic ligand moiety can be employed.

[0008] In a preferred form, the photochromic coordination compound is expressed by the following general formula (I):

![Diagram](image)

where unit B represents a mono- or poly-cyclic ring structure, such as phenanthroline, pyridine, imidazole, pyrazole, thiazole, pyrrole, drazine, triazine, polypyridine, porphyrin and phthalocyanine and the like, that contains one or more donor atom(s) X, such as carbon, or donor heteromatom(s) X, such as nitrogen, oxygen, sulfur, phosphorus, selenium, etc., n is integer from 0 to 3, [M] represents the coordination unit containing an acceptor atom M, such as rhenium(I), osmium(II), ruthenium(II), osmium(II), rhodium(III), iridium(II), gold(III), gold(I), silver(I), copper(I), copper(II), platinum(II), palladium(II), iron(II), cobalt(III), chromium(III), cadmium(II), boron(III) and the like, R1 and R2 individually represent alkyl groups and alkoxy groups, and R3 to R6 individually represent atoms or groups selected from the group of hydrogen atom,
halogen atom, hydroxyl group, alkyl group, alkoxy group, alkynyl group, cyano group, nitro group, alkyne carbonyl group, alkoxy carbonyl group, perfluoroalkyl group, aryl group, cycloalkyl group, arylene group, aryloxy group, alkylene group, mono- or dialkylaminocarbonyl group, alkylnitro group, arylnitro group, alkylnitro group, arylnitro group, and the like. In general, any alkyl or alkoxy group contains 1 to about 20 carbon atoms, any cycloalkyl group contains 3 to 8 carbon atoms, and any aryl group contains 6 to about 20 carbon atoms.

A non-limiting list of examples of diarylethene compounds containing a heterocyclic ethene-containing ligand moiety includes 5,6-diethynyl-1,10-phenanthroline, 2,3,7,8,12,13,17,18-octathiophenyl-5,10,15,20-tetraphenyl porphyrin, 6,7-dithienyl-dipyrrolo[3,2-\(a\):3',2'-c']phenazine and the like.

A non-limiting list of coordination units includes chlorotricarbonylrhenium(I), dithiolatozinc(II), dihaloplatinum(II), bipyridylplatinum(II), bis[bipyridyl]ruthenium(II), diphosphocopper(I), bipyridylcopper(I) and the like.

One of the advantages of the formation of coordination compounds from their pure organic counterparts (free ligands) in this invention is the extension of the excitation wavelength for the photocyclization of the diarylethene moiety from \(\lambda \leq 340\) nm to lower energy, so that the photorefractive forward reaction can proceed with visible light excitation by utilization of the low-energy absorptions characteristic of coordination compounds. In addition, the photorefractive reactions can be utilized to switch the photoluminescence properties characteristic of the coordination compounds.

**BRIEF DESCRIPTION OF THE DRAWINGS**

FIG. 1 shows a representative synthetic route for a diarylethene-containing ligand and its coordination compounds using 5,6-diethynyl-1,10-phenanthroline and its chlorotricarbonylrhenium(I) and dithiolatozinc(II) compounds as illustrative examples.

FIG. 2 shows the overlaid electronic absorption spectra of the open form (-) and the close form (- -) of a diarylethene-containing nitrogen donor ligand (L1).

FIG. 3 shows the overlaid electronic absorption spectra of the open form (-) and the close form (- -) of a diarylethene-containing coordination compound (1).

FIG. 4 shows the photocyclization reactions of (a) a diarylethene-containing ligand (b) its coordination compound using 5,6-diethynyl-1,10-phenanthroline and its chlorotricarbonylrhenium(I) compound as illustration.

FIG. 5 shows the overlaid corrected emission spectra of the open form (-) and the close form (- -) of (1) in benzene solution at 298 K.

FIG. 6 shows the overlaid corrected emission spectra of the open form (-) and the close form (- -) of (1) in EtOH-MeOH (4:1 v/v) glass at 77 K.

FIG. 7 shows the absorption spectral changes of complex (4) in benzene upon excitation at \(\lambda = 300\) nm.

FIG. 8 shows a representative synthetic route for diarylethene-containing ligands and their coordination compounds using 1-(aryl)-bis-(2,5-dimethyl-3-thienyl)-2-(2-pyridyl)imidazoles and their chlorotricarbonylrhenium(I) compounds as illustrative examples.

FIG. 9 shows (a) the absorption and (b) emission spectral changes of complex (6) in degassed chloroform solution (7.16x10^{-5} M) upon MLCT excitation at \(\lambda = 410\) nm.

**DESCRIPTION OF THE PREFERRED EMBODIMENTS**

**Example 1**

The ligand (L1) is synthesized by the Suzuki cross-coupling reactions of 2,5 equivalents of 2,5-dimethyl-3-thienylboronic acid and 5,6-dibromo-1,10-phenanthroline in the presence of the palladium catalyst, Pd(PPh3)4, and sodium carbonate in a heterogeneous mixture of water and THF according to the synthetic route depicted in FIG. 1. Excitation of (L1) with 313 nm light resulted in the formation of the close form, corresponding to the photocyclization product. The overlaid electronic absorption spectra of the open and close forms (L1) in benzene solution are shown in FIG. 2.

**Example 2**

Upon coordination to a chlorotricarbonylrhenium(I) complex, the open form of the corresponding complex (1) undergoes photocyclization with the excitation of both the intraligand absorptions at \(\lambda \leq 340\) nm and the metal-to-ligand charge transfer (MLCT) absorption characteristic of this coordination compound up to \(\lambda \leq 480\) nm. The electronic absorption data of (L1) and complex (1) are summarized in Table 1. The close forms of these compounds are found to undergo thermal backward reactions. The half-lives of the close forms have been determined and summarized in Table 2. The quantum yields for both photocyclization and photocycloreversion of (L1) and its rhenium complex (1) are summarized in Table 3.

**TABLE 1**

<table>
<thead>
<tr>
<th>Compound</th>
<th>Configuration</th>
<th>Absorption (in benzene) (\lambda_{\text{abs}}/\text{nm} (\text{d}^2\text{m}^{-1}\text{mol}^{-1}\text{cm}^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>(L1)</td>
<td>Open form</td>
<td>304 (8670)</td>
</tr>
<tr>
<td>(L1)</td>
<td>Close form</td>
<td>366 (2480), 510 (4040), 540 (3860)</td>
</tr>
<tr>
<td>(1)</td>
<td>Open form</td>
<td>338 (4930), 396 (4690)</td>
</tr>
<tr>
<td>(1)</td>
<td>Close form</td>
<td>390 (3670), 546 (5390), 586 (5950)</td>
</tr>
</tbody>
</table>

**TABLE 2**

<table>
<thead>
<tr>
<th>Compound</th>
<th>Half-lives (t_{1/2}) at 20°C</th>
<th>Half-lives (t_{1/2}) at 60°C</th>
</tr>
</thead>
<tbody>
<tr>
<td>(L1)</td>
<td>143 hours</td>
<td>222 mins</td>
</tr>
<tr>
<td>(1)</td>
<td>77.7 hours</td>
<td>79.3 mins</td>
</tr>
</tbody>
</table>
TABLE 3

<table>
<thead>
<tr>
<th>Compound</th>
<th>( \Phi_{11} )</th>
<th>( \Phi_{440} )</th>
<th>( \Phi_{650} )</th>
<th>( \Phi_{910} )</th>
</tr>
</thead>
<tbody>
<tr>
<td>(L1)</td>
<td>0.486</td>
<td>0</td>
<td>0.123</td>
<td>0.029</td>
</tr>
<tr>
<td>(L2)</td>
<td>0.532</td>
<td>0.648</td>
<td>0.028</td>
<td>0.009</td>
</tr>
</tbody>
</table>

*Values reported are corrected to the ratio of the photochemical acton conformation, i.e. with respect to the anti-parallel configuration.

[0027] The photoluminescence properties of both the open and closed forms were measured. FIGS. 5 and 6 display the overlay emission spectra of the open form and the close form of complex (1) in benzene at 298 K and in EtOH-MeOH glass (4:1 v/v) at 77 K. The emission of complex (1) was found to change from metal-to-ligand charge transfer (MLCT) phosphorescence to ligand-centered (LC) phosphorescence upon photocyclization of the open form to the close form. These demonstrate the change of emission properties upon photorhomic reactions. Table 4 summarized the emission data of ligand (L1) and complex (L).

TABLE 4

<table>
<thead>
<tr>
<th>Compound</th>
<th>Medium (77K)</th>
<th>Open form</th>
<th>Close form</th>
</tr>
</thead>
<tbody>
<tr>
<td>(L1)</td>
<td>Benzene (298)</td>
<td>583 (&lt;0.1)</td>
<td>644 (&lt;0.1)</td>
</tr>
<tr>
<td></td>
<td>Glass (77)</td>
<td>577 (5.2)</td>
<td></td>
</tr>
<tr>
<td>(L2)</td>
<td>Benzene (298)</td>
<td>955 (2.6)</td>
<td>626 (&lt;0.1)</td>
</tr>
<tr>
<td></td>
<td>Glass (77)</td>
<td>620 (6.4)</td>
<td></td>
</tr>
</tbody>
</table>

*Excitation wavelength at ca. 355 nm. Emission maxima are corrected values.

Example 2

[0028] Upon coordination of (L1) to a dithiolatozinc(II) complex, the open forms of the corresponding complexes (2), (3), and (4) undergo photocyclization with excitation at \( \lambda \leq 340 \) nm. FIG. 7 shows the absorption spectral changes of complex (4) upon excitation at \( \lambda = 300 \) nm. The electronic absorption maxima of both the open and the close forms of complexes (2), (3), and (4) are summarized in Table 5.

TABLE 5

<table>
<thead>
<tr>
<th>Complex</th>
<th>Configuration</th>
<th>Absorption maximum (in chloroform)</th>
</tr>
</thead>
<tbody>
<tr>
<td>(2)</td>
<td>Open form</td>
<td>302, 326, 378</td>
</tr>
<tr>
<td>(3)</td>
<td>Close form</td>
<td>366, 382, 536, 576</td>
</tr>
<tr>
<td>(3)</td>
<td>Open form</td>
<td>302, 326, 382</td>
</tr>
<tr>
<td>(4)</td>
<td>Close form</td>
<td>366, 382, 536, 576</td>
</tr>
<tr>
<td>(4)</td>
<td>Close form</td>
<td>366, 584, 542, 584</td>
</tr>
</tbody>
</table>

[0029] Those skilled in the art will recognize that various changes and modifications can be made in the invention without departing from the spirit and scope thereof. The various embodiments described were for the purpose of further illustrating the invention and were not intended to limit it.

Example 3

[0030] The ligands, 1-(4-methoxyphenyl)-bis-(2,5-dimethyl-3-thienyl)-1,1'-bipyridine (L2) and 1-(4-methylphenyl)-bis-(2,5-dimethyl-3-thienyl)-1,1'-bipyridine (L3), are synthesized by Suzuki cross-coupling reactions of 1.5 equivalents of 2,5-dimethyl-3-thienylboronic acid and 1-aryln-4,5-dibrom-2-(2-pyridyl)imidazol in the presence of the palladium catalyst, Pd(PPh3)4, and sodium carbonate in a heterogeneous mixture of water and THF according to the procedure similar to that of (L1) as depicted in FIG. 8. Excitation of (L2) and (L3) with 313 nm light resulted in the formation of the close form, corresponding to the photocyclization product.

Example 4

[0031] Similar to ligand (L1), upon coordination of (L2) and (L3) to chlorotris(pyridyl)rhodium(I) complex, the open form of the corresponding complexes (5) and (6) undergo photocyclization with the excitation of both the intraligand absorptions at \( \lambda \leq 370 \) nm and the metal-to-ligand charge transfer (MLCT) absorption characteristic of this coordination compound up to \( \lambda \leq 470 \) nm. The electronic absorption data of ligands (L2)–(L3) and complexes (5)–(6) are summarized in Table 6. Apart from the change in the absorption properties, the emissions of all the complexes were also found to drop significantly upon conversion to the close form. These further illustrate the change of emission properties upon photorhomic reactions. The representative emission and electronic absorption spectral changes of the open form in chloroform solution upon photo-irradiation are shown in FIG. 9.

TABLE 6

<table>
<thead>
<tr>
<th>Compound</th>
<th>Configuration</th>
<th>Absorption maximum (in chloroform)</th>
</tr>
</thead>
<tbody>
<tr>
<td>(L2)</td>
<td>Open form</td>
<td>320</td>
</tr>
<tr>
<td>(L2)</td>
<td>Closed form</td>
<td>330, 425, 586</td>
</tr>
<tr>
<td>(L3)</td>
<td>Open form</td>
<td>319</td>
</tr>
<tr>
<td>(L3)</td>
<td>Closed form</td>
<td>334, 410, 576</td>
</tr>
<tr>
<td>(5)</td>
<td>Open form</td>
<td>352, 435, 448</td>
</tr>
<tr>
<td>(5)</td>
<td>Closed form</td>
<td>284, 374, 473, 712</td>
</tr>
<tr>
<td>(6)</td>
<td>Open form</td>
<td>353, 425, 448</td>
</tr>
<tr>
<td>(6)</td>
<td>Closed form</td>
<td>286, 374, 480, 713</td>
</tr>
</tbody>
</table>

[0032] The ligand, 1,3-dimethyl-4,5-bis-(2,5-dimethyl-3-thienyl)-imidazolium hexahloroantimonate (L4), is synthesized by the methylation of 4,5-bis-(2,5-dimethyl-3-thienyl)-1,1'H-imidazol, which is also prepared using Suzuki cross-coupling reactions of 2.5 equivalents of 2,5-dimethyl-3-thienylboronic acid and 4,5-dibromo-1'H-imidazol in the presence of the palladium catalyst, Pd(PPh3)4, and sodium carbonate in a heterogeneous mixture of water and THF according to the procedure similar to that of (L1), as depicted in FIG. 10. Excitation of the ligand with \( \lambda \leq 290 \) nm light resulted in the formation of the close form, corresponding to the photocyclization product.

Example 5

[0033] Upon coordination to palladium(II) iodide to form the bis[1,3-dimethyl-4,5-bis(2,5-dimethyl-3-thienyl)-imidazol-2-ylidene]diodo(II) complex, the open form of the corresponding complex (7) undergoes photocyclization with the excitation of both the intraligand absorptions at \( \lambda \leq 290 \) nm and the absorption characteristic of this coordination compound up to \( \lambda \leq 320 \) nm. The electronic absorption data of ligand (L4) and complex (7) are summarized in Table.
7. The representative electronic absorption spectral changes of complex (7) in dichloromethane solution upon photo-irradiation are shown in FIG. 11.

<table>
<thead>
<tr>
<th>Compound</th>
<th>Configuration</th>
<th>Absorption maximum (in dichloromethane) λ&lt;sub&gt;max&lt;/sub&gt;/nm</th>
</tr>
</thead>
<tbody>
<tr>
<td>(L5)</td>
<td>Open form</td>
<td>235</td>
</tr>
<tr>
<td>(L5)</td>
<td>Close form</td>
<td>580</td>
</tr>
<tr>
<td>(7)</td>
<td>Open form</td>
<td>240, 305</td>
</tr>
<tr>
<td>(7)</td>
<td>Close form</td>
<td>345, 540</td>
</tr>
</tbody>
</table>

Example 5

[0034] 5-Trimethylsilyl-ethyl-2,3-bis-(2-methyl-3-benzyl-3,2-b)thienyl)thiophene is synthesized by the Sonogashira coupling reaction of trimethylsilylacetylene and the bromo-derivative of 2,3-bis-(2-methyl-3-benzyl-3,2-b)thienyl)thiophene, which is also prepared from the Suzuki cross-coupling reaction of 2.5 equivalents of 2-methyl-3-benzyl(3,2-b)thienyl)boronic acid and 2,3-dihalothiophene in the presence of the palladium catalyst, Pd(PPh₃)₄, and sodium carbonate in a heterogeneous mixture of water and THF according to the procedure similar to that of (L1). Subsequent deprotection of the trimethylsilyl group using potassium carbonate in dichloromethane solution gives the target ligand ethyl-2,3-bis-(2-methyl-3-benzyl-3,2-b)thienyl)thiophene (L5). The synthetic route is shown in FIG. 12. Excitation of the ligand with λ≤352 nm light resulted in the formation of the close form, corresponding to the photocyclization product.

[0035] Upon reaction with dichlorobis(triethylphosphine) platinum(II), the open form of the corresponding complex (8) undergoes photocyclization with the excitation up to λ≤430 nm. The electronic absorption data of ligand (L5) and complex (8) are summarized in Table 8. The representative electronic absorption changes of complex (8) in dichloromethane solution upon photo-irradiation are shown in FIG. 13.

<table>
<thead>
<tr>
<th>Compound</th>
<th>Configuration</th>
<th>Absorption maximum (in dichloromethane) λ&lt;sub&gt;max&lt;/sub&gt;/nm</th>
</tr>
</thead>
<tbody>
<tr>
<td>(L5)</td>
<td>Open form</td>
<td>290, 294, 304, 3144</td>
</tr>
<tr>
<td>(L5)</td>
<td>Close form</td>
<td>300, 382, 554</td>
</tr>
<tr>
<td>(8)</td>
<td>Open form</td>
<td>2068, 296, 304, 3186, 384</td>
</tr>
<tr>
<td>(8)</td>
<td>Close form</td>
<td>372, 540</td>
</tr>
</tbody>
</table>

What is claimed is:

1. A photochromic coordination compound comprising a diarylethene, in which the ethene moiety forms part of a mono or poly-cyclic ring structure containing or being connected to one or more donor atoms coordinated to an acceptor atom of the coordination unit, wherein the photochromic coordination compound excludes zinc azaporphyrin, magnesium azaporphyrin, palladium azaporphyrin, magnesium phthalocyanine and zinc phthalocyanine.

2. The photochromic compound which is a diarylethene-containing coordination compound in which the diarylethene contains part of a mono or poly-cyclic ring structure with one or more donor atom(s) or heteroatom(s) coordinated to a coordination unit [M], which contains an acceptor atom, M, expressed by the general formula (I):

   ![Diagram](image)

where unit B represents a mono- or poly-cyclic ring structure that contains one or more donor atom(s) or heteroatom(s) X, n is integer from 0 to 3, [M] represents a coordination unit containing an acceptor atom M and excludes zinc azaporphyrin, magnesium azaporphyrin, palladium azaporphyrin, magnesium phthalocyanine and zinc phthalocyanine, R₁ and R₂ individually represent an alkyl group or an alkoxy group, and R₃ to R₅ individually represent atoms or groups selected from the group of hydrogen atom, halogen atom, hydroxyl group, alkyl group, alkoxy group, alkyln group, cyano group, nitro group, alkylcarbonyl group, alkoxycarbonyl group, perfluoralkyl group, ary group, cyclohexyl group, arylethyl group, arythioether group, aryloxy group, arylenecarbonyl group, and aryloxy carbonyl group.

4. The photochromic coordination compound in accordance with claim 2, wherein the photochromic coordination compound excludes zinc azaporphyrin, magnesium azaporphyrin, palladium azaporphyrin, magnesium phthalocyanine and zinc phthalocyanine.

5. The photochromic compound in accordance with claim 2, wherein B is selected from the group consisting of pyridine, imidazole, pyrazole, thiazole, pyrrole, diazine, triazine, polypyridine, porphyrin and phthalocyanine; and acceptor atom M is selected from the group consisting of phthion(II), zinc(II), ruthenium(II), osmium(II), rhodium (III), iridium(III), gold(III), gold(I), silver(I), copper(I), copper(II), platinum(II), palladium(II), iron(II), cobalt(III), chromium(III), cadmium(II) and boron(III).

6. The photochromic compound in accordance with claim 2, wherein the donor atom or heteroatom X is selected from the group consisting of carbon, nitrogen, oxygen, sulfur, phosphorus and selenium.

7. The photochromic compound in accordance with claim 2, wherein B is a porphyrin.

8. The photochromic compound in accordance with claim 7, wherein R₁ and R₂ represent methyl groups, and R₃ and R₄ represent hydrogen atoms.

9. The photochromic compound in accordance with claim 7, wherein R₁ and R₂ represent hydrogen atoms.

10. The photochromic compound in accordance with claim 7, wherein R₂ and R₃ represent methyl groups.

11. The photochromic compound in accordance with claim 7, wherein R₂ and R₃ represent bromine atoms.

12. The photochromic compound in accordance with claim 7, wherein M represents zinc(II).

13. The photochromic compound in accordance with claim 7, wherein M represents platinum(II).
14. The photochromic compound in accordance with claim 2, wherein unit B is a pyridine.
15. The photochromic compound in accordance with claim 2, wherein unit B is a diazine.
16. The photochromic compound in accordance with claim 2, wherein unit B is a triazine.
17. The photochromic compound in accordance with claim 2, wherein unit B is a polypyridine.
18. The photochromic compound in accordance with claim 2, wherein unit B is a phthalocyanine.
19. The photochromic compound in accordance with claim 2, wherein M represents rhenium(I).

20. A photochromic coordination compound comprising:
a coordination unit having an acceptor atom; and
a diarylethene having one or more donor atoms in an ethene moiety and coordinated to the acceptor atom of the coordination unit;
wherein the coordination compound is photochromic and
excludes zinc azaporphyrin, magnesium azaporphyrin, palladium azaporphyrin, magnesium phthalocyanine and zinc phthalocyanine.

* * * * *