Novel phthalonitrile derivatives as potential compounds for extraction and complexation of metal cations

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ABSTRACT

The synthesis and the binding properties of novel phthalonitrile derivatives 1-3 towards metal cations have been described in this paper. The complexation and extraction of some transition and heavy metal cations have been followed by UV-visible spectrophotometry absorption in methanol. The conductivity studies have been used in order to confirm complex's stoichiometries. The treatment of UV spectra by digital program showed the formation of ML (with ML2 in some cases) (M=metal, L=ligand) species. Beyond the discussion of the stability profiles of complexes particular attention is paid to the selectivity towards Cu2+ in the 1st sequence of transition metal cations and towards Hg2+ in the sequence of heavy metal cations.

Indexing terms/Keywords
Phthalonitriles, cation binding properties, selectivity, complexation, extraction.

Academic Discipline And Sub-Disciplines
Organic and organometallic chemistry

SUBJECT CLASSIFICATION
Organic chemistry
INTRODUCTION

Since a long time, pharmaceutical industry required green chemistry to find new compounds able to copy the metalloenzymes and to find a better way to mimic their reactions. Continuous challenge of researches is devoted in this way to create new and efficient complexes. Among nitrile compounds, phthalonitriles are known as good candidates for the complexation of metal cations. They are also thermally stable with high performance composite structures. Besides, they constitute a class of high-temperature polymers that has a number of exceptional properties such as outstanding thermal and thermo-oxidative stability, good moisture resistance and superior fire resistance. [1-12]

As a continuation of our interests in phthalonitriles [13-15], we report here the synthesis, complexation and extraction properties of novel phthalonitriles 1-3 towards transition and heavy metal cations. The three compounds were prepared by Hamdi et coll.[16] The studies were performed by UV-visible spectrophotometry absorption and by conductometry. The last technique was used in order to test some stoichiometries of complexes formed. The liquid - liquid extraction of metallic picrate salts by ligands 1 - 3 were followed from water into dichloromethane.

![Scheme 1: Structure of studied phthalonitriles derivatives 1-3.](image)

Results and discussion

Synthesis

**Synthesis of 1**

4- nitrophthalonitrile (0.38 g, 1.92 mmol), 2,4,6 Trimethylphenylamine (0.55 g, 1.92 mmol) and anhydrous DMF (15 mL) were added to a round bottom flask under a nitrogen atmosphere. A fine powder of anhydrous potassium carbonate (0.8 g, 5.76 mmol) was added to this mixture. The resulting mixture was then heated to 80 °C in an oil bath under a nitrogen atmosphere for 6 hours. The crude product was collected by filtration and washed with distilled water. The crude product was recrystallized from THF-petroleum ether to afford a white powder. Yield: 0.77 g (98 %). M.p. = 400 °C. IR (KBr pellet): 1305 (C-N), 1568 (C=C), 2235 (C≡N), 3049 (C-H, aromatic). ^1^H NMR (DMSO-d$_6$): 8.38-8.98 (m, 5H, Harom), 6.57 (s, 2H, H$_3$), 4.21 (s, 1H, NH), 2.09 (s, 3H, CH$_3$), 2.02 (s, 3H, CH$_3$), 2.08 (s, 3H, CH$_3$). Anal. Calc. for C$_{28}$H$_{14}$N$_2$O$_2$: C, 70.1; H, 5.2; N, 9.0%.

**Synthesis of 2**

The synthesis of 2 was similar to 1, except 4-Hydroxybenzaldehyde (1.27 g, 5.77 mmol) was employed instead of 2,4,6 Trimethylphenylamine. The amounts of the other reagents were: 4-nitrophthalonitrile (1 g, 5.8 mmol) and anhydrous potassium carbonate (2 g, 13.88 mmol). Yield: 1.12 g (76%) M.p. = 350 °C. IR spectrum (cm$^{-1}$): 1263 (C-O-C), 1568 (C-C), 2236 (C≡N), 3049 (C-H, aromatic). ^1^H NMR (DMSO-d$_6$): 9.1 (s, 1H, CHO), 7.1-8.98 (m, 8H, H$_a$). Anal. Calc. for C$_{28}$H$_{14}$N$_2$O$_2$: C, 78.1; H, 5.6; N, 16.0%.

**Synthesis of 3**

The synthesis of 3 was similar to 1, 2-(3,4-dimethoxyphenyl)ethanol was employed instead of 2,4,6 Trimethylphenylamine. The amounts of the other reagents were 4-nitrophthalonitrile (1 g, 5.55 mmol) and anhydrous potassium carbonate, 2 g (13.88 mmol). Yield: 0.77 g (85%) M.p. = 370 °C. IR spectrum (cm$^{-1}$): 3077 (Ar-CH), 2227 (C-N), 1601 (C-C), 836, 797 (C-F): 1263 (C-O-C), 1568 (C=C), 2227 (C≡N), 3049 (C-H, aromatic). ^1^H NMR (DMSO-d$_6$): 6.50-7.77 (m, 8H, H$_a$), 3.74 (s, 6H, OCH$_3$). 2.86 (t, 2H, CH$_2$), 4.1 (t, 2H, CH$_2$). Anal. Calc. for C$_{28}$H$_{14}$N$_2$O$_2$: C, 70.1; H, 5.2; N, 9.0%.

Instrumentations

The spectrophotometric and conductometric studies were followed in methanol (Riedel-de Haën for HPLC) and the dichloromethane (Fluka, Purum) used in extraction, were commercial and used without further purification. The metal salts chosen were chlorides (Fluka, Purum) and were dried under vacuum for at least 24 h before use. The concentrations of the stock solutions of cations were standardized by complexometry using the appropriate colored indicators [17]. The UV absorption spectra were recorded on a Perkin Elmer Lambda 11 spectrophotometer. The supporting electrolyte used in the stability constant determinations was NEt$_4$Cl (Acros Organics) according to the procedure already described [18].
picrate salts employed in extraction were prepared as described in literature [19]. A conductivity measurement was made by using Cyber Scan PCS10 conductivity meter. The conductivity cell constant is K = 0.9 cm.

Melting points were determined using an Electrothermal apparatus and are uncorrected. 1H NMR spectra were carried on a Varian Gemini 400 (400 MHz) spectrometer using TMS as internal standard (δ = 0 ppm). IR spectra were recorded on a Perkin Elmer 398 Spectrophotometer. Elemental analyses were performed on Perkin Elmer 2400 elemental analyzer, and the values found were within ±0.3% of the theoretical values.

**Complexation by UV - visible spectrophotometer**

The stability constants β* are obtained by the least squares method of a differences square sum of a differences (U = Σ (A*exp - A*calc)^2), where A*exp and A*calc are, respectively, the experimental and calculated absorbances (U = Σ (A*exp - A*calc)^2). The β* values correspond to the average of at least three independent experiments [20, 21].

The UV absorption spectra have been recorded between 220 and 360 nm. The free phthalonitriles 1-3 exhibit one maximum of absorption in all cases of Metal/Ligand systems (λmax~300nm). Another selectively is detected with ligand 1, which leads to a selectivity of this ligand with respect to the Irving-Williams rules which is translated by the increase of the complex stability from Mn** to Cu** and then a decrease to Zn**. Furthermore, the complexes ML3 in the case of Cu** and Zn** with ligand 3 are less stable than their corresponding ML. Their stability constants are equal to 3.45 and 3.25 logarithmic units, respectively. In the case of heavy metal cations, Hg** is more interacted by the three derivatives 1-3 and the best value of stability constants is 4.78 logarithmic units found with 1. Comparing between the three phthalonitriles, 1 seems to have more affinity towards transition and heavy metal cations. In both case of series of metal cations, complexes ML formed by ligand 1 are more stable than with 2 and 3 and even more stable than ML2. The presence of nitrile functions and nitrogen

**Table 1**: Complexation of transition metal cations by 1-3 in methanol, at 25°C, I = 10^-2 mol.L^-1.

<table>
<thead>
<tr>
<th>M:L</th>
<th>Mn**</th>
<th>Co**</th>
<th>Ni**</th>
<th>Cu**</th>
<th>Zn**</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1:1</td>
<td>a</td>
<td>3.45</td>
<td>4.12</td>
<td>4.76</td>
</tr>
<tr>
<td>2</td>
<td>1:1</td>
<td>a</td>
<td>2.80</td>
<td>3.60</td>
<td>4.05</td>
</tr>
<tr>
<td>3</td>
<td>1:1</td>
<td>a</td>
<td>3.34</td>
<td>3.44</td>
<td>3.87</td>
</tr>
<tr>
<td></td>
<td>1:2</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>a:</td>
<td>Absorbance changes too small to enable satisfactory fitting.</td>
<td>0.01 &lt; σ_N^** &lt; 0.2</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Table 2**: Complexation of heavy metal cations by 1-3 in methanol, at 25°C, I = 10^-2 mol.L^-1.

<table>
<thead>
<tr>
<th>M:L</th>
<th>Hg**</th>
<th>Pb**</th>
<th>Ag**</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1:1</td>
<td>4.78</td>
<td>4.09</td>
</tr>
<tr>
<td>2</td>
<td>1:1</td>
<td>4.00</td>
<td>a</td>
</tr>
<tr>
<td>3</td>
<td>1:1</td>
<td>3.89</td>
<td>a</td>
</tr>
</tbody>
</table>

| a: Absorbance changes too small to enable satisfactory fitting. | 0.01 < σ_N^** < 0.06 |

As regards to the complexes ML in the 1st sequence of transition metal cations, the stability constants shift between 2.80 and 4.76 logarithmic units. The highest value is in favor of Cu** with ligand 1 which leads to a selectivity of this ligand towards Cu** (S_Cu**/Co** = 26). Another selectively is detected with ligand 2 towards Cu** with an approximate value S Cu**/Zn** equals to 18. The profiles of stability of ML (figure 1) illustrate the resemblance of the stability sequence of high spin octahedral metal complexes with the Irving-Williams rules which is translated by the increase of the complex stability from Mn** to Cu** and then a decrease to Zn**. Furthermore, the complexes ML3 in the case of Cu** and Zn** with ligand 3 are less stable than their corresponding ML. Their stability constants are equal to 3.45 and 3.25 logarithmic units, respectively. In the case of heavy metal cations, Hg** is more interacted by the three derivatives 1-3 and the best value of stability constants is 4.78 logarithmic units found with 1. Comparing between the three phthalonitriles, 1 seems to have more affinity towards transition and heavy metal cations. In both case of series of metal cations, complexes ML formed by ligand 1 are more stable than with 2 and 3 and even more stable than ML2. The presence of nitrile functions and nitrogen
on the bridge between the two aromatic units seems to increase the affinity of ligand 1. The triple bond between N and C atoms extends aromaticity and then leads to a higher interaction between the ligand 1 and the cation. Moreover, the soft character of both nitrogen atom and aromatic units could be the reason of the high interaction with soft acid as transition and heavy metal cations according to Lewis.

Complexation by conductometry

The present work proposes a simple, accurate and low-cost method employing conductometric detection during the complexation titration using metal cation chloride as titrant. The process was based on the chemical interaction between cation coming from chloride ions and phthalonitriles 1-3. Under optimized experimental conditions the method was applied for some cases of systems Metal/Ligand which created previously some difficulties in their treatments in UV-visible spectrophotometer absorption studies.

This method is carried on by the projection of the meeting point of the slopes corresponding to the variation of conductance values via the ratio \( R = C_M/C_L \) according to the literature. \([29, 30]\) The results obtained are in close agreement with those obtained using the first technique. Only system Cu\(^{2+}/3\) (figure 2) is presented here. In fact, the titration of the solution of phthalonitriles 3 by a solution of CuCl\(_2\) shows a variation of conductance values, proportionally to the ratio \( R = C_M/C_L \). After plotting the conductance values, variation of the curve was observed and the projection of the meeting point of the slopes corresponds to the stoichiometry of the complex formed during the titration. In the case of Cu\(^{2+}/3\), complexes ML and ML\(_2\), as shown on figure 2, are formed simultaneously.

Extraction studies

The extraction experiments from water into dichloromethane were performed according to the following procedure: 5 mL of 2.5x10\(^{-4}\) M aqueous picrate solution and 5 mL of 2.5x10\(^{-4}\) M solution of ligand in CH\(_2\)Cl\(_2\) were mechanically shaken in stoppered glass tube for 3 min, then magnetically stirred in a thermostatted water bath at 20 ± 0.1 °C for 30 min and finally left standing for a further 30 min in order to obtain a complete separation of the two phases. The absorbance \( A \) of the metal picrates remaining in the aqueous phase was then determined spectrophotometrically at 355 nm. The percentage extraction, %E, are derived from the expression 100(A\(_0\) - A)/A\(_0\), where A\(_0\) is the absorbance of the aqueous solution of a blank experiment without phthalonitrile. \([22]\]

The percentages of extraction %E calculated are generally minor in all cases of cations, slightly improved for ligand 1, where %E shifts between 4 and 36. This enhancement is probably due to the presence of N in addition comparing with 2 and 3. The highest value of %E is in favor of Cu\(^{2+}\) (36%) and Ag\(^{+}\) (27%). The first result confirms once again the high affinity of this ligand towards Cu\(^{2+}\). While the second, towards Ag\(^{+}\), could be interpreted by the soft character of both base N and acid Ag\(^{+}\) though this preference have not been seen in the complexation study.
Conclusion

This review is dealing about the synthesis and the binding properties of new phthalonitriles derivatives 1-3 towards some transition and heavy metal cations. The UV spectrophotometric and conductometric studies illustrates the formation of complexes ML and ML₂ in addition of ML in case of Cu²⁺ and Zn²⁺ with ligand 3. Moreover, the profiles of stability of complexes ML is following the Irving-Williams rules and making evidence of the high stability of complexes of Cu²⁺. A selectivity S(Cu²+/Co²⁺) of 26 with 1 and 18 with 2 are detected. As regards to the stability, the presence of nitrogen atom in the bridge between the two units in ligand 1 seems to improve its affinity comparing with 2 and 3 carrying oxygen, a hard atom, in their structures.

REFERENCES

[16] Naceur HAMDI et al., unpublished results