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Cytotoxic, antibacterial activity and physico-chemical properties of some acid catalyzed Schiff bases

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Some novel Schiff bases have been derived using aromatic aldehydes and aromatic amine in ethanol under acidic conditions. The geometry of these Schiff bases was established using various spectroscopic methods like infrared (IR), nuclear magnetic resonance (NMR) and mass spectrometry. The conductivity studies reveal the non electrolytic behavior of Schiff bases. The synthesized Schiff bases were tested in vitro with the aim of identifying novel lead compounds active against various types of bacteria and shrimps.

Key words: Acid catalyzed, Schiff bases, NMR, antibacterial activity, brine shrimp.

INTRODUCTION

In chemistry, Schiff bases finds versatile use (Sheehan and Grenda, 1962); some of them are the basic units in certain dyes, whereas, some are used as liquid crystals. In organic synthesis, Schiff base reactions are useful in making carbon-nitrogen bonds. Schiff bases appear to be important intermediates in a number of enzymatic reactions involving interaction of an enzyme with an amino or a carbonyl group of the substrate (Liimatainen et al., 2000). Many biological important Schiff bases ligands have been reported which posses antibacterial (Shi et al., 2010), antifungal (Zhong et al., 2009), antimicrobial (Raman et al., 2003), anticonvulsant (Verma et al., 2004), antiHIV (Pandeya et al., 1999), anti-inflammatory (Janos et al, 2009), and antitumor activity (Billman and Schmidgall, 2006). The antitumor activity of the bases towards ascetic tumors increases considerably with a slight increase in water solubility.

Previously a number of biologically important complexes have been reported by our group (Rehman et. al., 2008, 2009; Jamil et al., 2010). In order to broaden the scale of investigations on the Schiff bases, we have now synthesized, structurally characterized and determined cytotoxic and antibacterial activity of a number of Schiff bases, derived from various aromatic aldehydes and aromatic amines and the results are reported herein.

MATERIALS AND METHODS

All the glassware used were cleaned with chromic mixture, detergent and finally with distilled water and then dried in oven at 110°C. All the solvents used in the experiment were purified and dried according to standard procedures (Perrin and Armengo, 1988).

All the chemicals used were of analytical grade and used as such without further purification. Thin layer chromatography (TLC) was performed on precoated silica gel (E-Merck) 60F-254 analytical plates (0.25 mm). The solvent system used was n-hexane/ethyl acetate (1:9).

Melting points were determined on Gallen Kamp digital (England) melting point apparatus. Fourier transform infrared (FTIR) spectra were recorded on Bruker FTIR Tensor 27 spectrophotometer; nuclear magnetic resonance (NMR) spectra were recorded on Bruker 300MHZ FTNMR spectrophotometer, mass spectra were recorded on gas chromatography-mass spectrometry (GC: MS) model MAT 312.

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Abbreviations: FTIR, Fourier transform infrared; IR, infrared; NMR, nuclear magnetic resonance; TLC, thin layer chromatography; DMSO, dimethyl sulfoxide; GC-MS, gas chromatography-mass spectrometry; MIC, minimum inhibitory concentration.
Synthesis of Schiff bases

General procedure

10 m moles of aromatic amine were dissolved in a 15 ml of ethanol in a 100 ml 2-neck round bottom flask equipped with a reflux condenser protected by a calcium chloride drying tube and a quick fit thermometer, then 10 m mole of aromatic aldehyde were dissolved with constant stirring; to this reaction mixture 0.2 to 0.4ml HCl were added with cooling on an ice bath. Yellow color solid separates just after the addition of HCl. The mixture was stirred for 1 - 2 h at room temperature. The reaction mixture was refluxed on a water bath at 90 to 95°C for 3 to 4 h, the reaction was examined by TLC with time to time till completion. The excess of solvent was removed under reduced pressure; the product was was hed with column chromatography using hexane-ethyl acetate (3:7) system.

3-methoxy-benzylidene-p-tolyl-amine (Schiff base 1)

This compound was obtained as yellow solid, 75% yield, m.p. 112°C. IR (KBr) Cm⁻¹ 1588 (C=N), 3044 (C=H Aro), 1590 (C=C Aro) 1H-NMR 2.39 (s,3H,CH₃), 3.95 (s, 3H, O-CH₃), 8.39 (s, 1H, N=CH), 7.1 to 7.7 (m, 7H, Aro), 13C-NMR 18.8 (CH₃), 55.5 (OCH₃), 115 to 135 (Aro), 155.5 (NCH), 160.9, (HC=N), 167.1 (OC) M/z 225, 118, 91.

3,4-Dimethoxy-benzylidene-p-tolyl-amine (Schiff base 2)

This compound was obtained as yellow solid, 75% yield, m.p. 132°C. IR (KBr) Cm⁻¹ 1576 (C=N), 3090 (C=H Aro), 1601 (C=C Aro) 1H-NMR 2.39 (s,3H,CH₃), 3.95 (s, 3H, O-CH₃), 8.41 (s, 1H, N=CH), 7.0 to 7.7 (m, 7H, Aro), 13C-NMR 18.2 (CH₃), 55.5 (OCH₃), 119 to 145 (Aro), 153.9 (NCH), 160.0, (HC=N), 165.9 (OC) M/z 255, 118, 91.

3,4,5-Trimethoxy-benzylidene-p-tolyl-amine (Schiff base 3)

This compound was obtained as yellow solid, 75% yield, m.p. 170°C. IR (KBr) Cm⁻¹ 1580 (C=N), 3078 (C=H Aro), 1610(C=C Aro) 1H-NMR 2.35 (s,3H,CH₃), 3.92 (s, 3H, 3-OCH₃), 8.34 (s, 1H, N=CH), 7.1-7.24 (m, 7H, Aro), 13C-NMR 18.5 (CH₃), 55.5 (p-OCH₃), 50.0 (p-OCH₃), 44.8 (m-OCH₃), 114-141 (Aro), 152.9 (NCH), 160.4, (HC=N), 165.2 (OC) M/z 285, 270, 118, 91.

2-Methoxy-4-(p-tolylimino-methyl)-phenol (Schiff base 4)

This compound was obtained as yellow solid, 75% yield, m.p. 159°C. IR (KBr) Cm⁻¹ 1577 (C=N), 3065 (C=H Aro), 1605 (C=C Aro) 1H-NMR 2.40 (s,3H,CH₃), 3.71 (s, 3H, O-CH₃), 8.37 (s, 1H, N=CH), 7.1-7.7 (m, 7H, Aro), 4.31 (s, 1H, OH) 13C-NMR 18.7 (CH₃), 55.0 (OCH₃), 114-135 (Aro), 148 (COH Aro), 153.5 (NCH), 160.4, (HC=N), 165.5 (OC) M/z 241,118,91.

4-Flouro-benzylidene-p-tolyl-amine (Schiff base 5)

This compound was obtained as yellow solid, 75% yield, m.p. 110°C. IR (KBr) Cm⁻¹ 1587 (C=N), 3077 (C=H Aro), 1612(C=C Aro) 1H-NMR 2.36 (s,3H,CH₃), 8.41 (s, 1H, N=CH),7.1-7.8 (m, 8H, Aro), 13C-NMR 18.6 (CH₃), 119-140 (Aro), 155.0 (NCH), 160.8, (HC=N), 163.4 (CF), 167.4 (OC) M/z 213, 118, 91.

4-Chloro-benzylidene-p-tolyl-amine (Schiff base 6)

This compound was obtained as yellow solid, 75% yield, m.p. 144°C. IR (KBr) Cm⁻¹ 1570 (C=N), 3055 (C=H Aro), 1600 (C=C Aro) 1H-NMR 2.36 (s,3H,CH₃), 8.41 (s, 1H, N=CH),7.1-7.8 (m, 8H, Aro) 13C-NMR 18.5 (CH₃), 117-130 (Aro), 155.1 (NCH), 160.3, 159.1 (OC), (HC=N), 167.0 (OC) M/z 229,118,91.

4-Bromo-benzylidene-p-tolyl-amine (Schiff base 7)

This compound was obtained as yellow solid, 75% yield, m.p. 129°C. IR (KBr) Cm⁻¹ 1579 (C=N), 3076 (C=H Aro), 1597(C=C Aro) 1H-NMR 2.36 (s,3H,CH₃), 8.40 (s, 1H, N=CH),7.0-7.7 (m, 8H, Aro), 13C-NMR 18.9 (CH₃), 121-140 (Aro), 155.9 (NCH), 160.8, 151.1 (CBr), (HC=N), 167.0 (OC) M/z 273,118,91.

4-Ido-benzylidene-p-tolyl-amine (Schiff base 8)

This compound was obtained as yellow solid, 75% yield, m.p. 121°C. IR (KBr) Cm⁻¹ 1585 (C=N), 3066(C=H Aro), 1595(C=C Aro) 1H-NMR 2.36 (s,3H,CH₃), 8.39 (s, 1H, N=CH),7.1-7.7 (m, 7H, Aro) 13C-NMR 18.5 (CH₃), 115-130 (Aro), 155.0 (NCH), 160.8, 149.0 (Cl), (HC=N), 167.5 (OC) M/z 321, 118, 91.

Biological studies

All the bases were tested against various organisms for the following activities:

Brine shrimp lethality bioassay

Artificial sea water was prepared by dissolving 3.8 g sea salt per liter of double distilled water and filtered. Sea water was placed in a small tank and shrimp eggs (Artemia Salina) (1 mg) were added to the large compartment of the tank, which was darkened by covering it with aluminum foil. The illuminated compartment attracted shrimps larvae through perforation in the dam. It was allowed to stand for 24 h at 25°C for the shrimps to hatch and mature. Diffe-rent concentrations (150,100 and 150 µg/ml) of the test samples were prepared in dimethyl sulfoxide (DMSO). Three (3) replicates were prepared for each concentration making a total of 24 vials. After 2 days when shrimp larvae matured, 5 ml sea water and 12 shrimps per vial were added, allowed to stand for 24 h under illumination. After 24 h, the number of surviving shrimps were counted and recorded. Data were analyzed with a Finney computer program to determine the LD₅₀ values (Solis et al., 1993; Finney 1979).

Antibacterial activity

The antibacterial activity was determined by using the agar well diffusion method (Rahman et al., 2001). The wells were dug in the media with a sterile borer and eight-hour-old bacterial inoculums containing ca 10⁵- 10⁶ colony-forming units (CFU)/ml was spread on the surface of the nutrient agar using a sterile cotton swab. The recommended concentration of the test sample (2 mg/ml in DMSO) was introduced in the respective wells. Other wells containing DMSO and the reference antibacterial drug served as negative and positive controls, respectively. The plates were incubated immediately at 37°C for 20 h. The activity was determined by using the diameter of the inhibition zone (in mm) showing complete inhibition. Growth inhibition was calculated with reference to the positive control.
RESULTS AND DISCUSSION

Synthesis of Schiff bases

Schiff bases have been synthesized by reacting aromatic aldehydes and primary aromatic amine under acidic conditions using ethanol as solvent. This is depicted in Figure 1 and Table 1.

Spectroscopic characterization

Several Schiff bases have been synthesized with structures as characterized by NMR, mass and IR data (see experimental section). IR spectroscopy is a constructive tool in structural determination of Schiff base compounds (Rehman et al., 2005). The FTIR spectrum of Schiff base showed stretching frequencies at 1580, 3078 and 1610 cm\(^{-1}\) which are characteristics of C=N, C-H and C=C (Aro), respectively. The \(^1\)H NMR spectrum showed signals at 2.3, 3.9 and 8.3 ppm (s) which are characteristics of methyl, methoxy and N=CH groups, respectively present in the structure. The signals of aromatic protons appeared at the range of 7.1 - 7.2 ppm (m). The main peaks in \(^{13}\)C-NMR spectra are successfully assigned and these results are supported by literature (Rehman et al., 2009). The mass spectrum showed molecular ion peak at m/z which is the molecular weight of the product. The remaining fragments appeared at m/z 270, 118 and 91, respectively.

Biological activities

Brine shrimp bio-assay

Bioactive compounds are often toxic to shrimp larvae, hence in vivo lethality to shrimp larvae can be used as simple and convenient preliminary monitor for new bioactive synthetic products (Wikipedia, 2006). This is rapid, inexpensive, in-house general bioassay that may serve as an intermediate test before further in vivo animal experiments on large scale.

All the synthesized Schiff bases were investigated for their cytotoxic property by brine shrimp lethality bioassay. Most of them showed significant activity against brine shrimp nauplii. The LD\(_{50}\) values of the synthesized bases are presented in Table 2.

The Schiff base 4 was found to be highly active with LD\(_{50}\) value of 0.053 \(\mu\)l/ml. The compounds 1, 2 and 6 also showed good activity with LD\(_{50}\) of 45.02, 25.52 and 1.26 \(\mu\)l/ml, respectively; the rest of the synthesized Schiff bases also exhibited significant activity. The activity order is as 4 < 6 < 2< 1 < 7 < 3 < 8 < 5.

Antibacterial studies

The resulting synthesized Schiff bases were checked against various microorganisms such as Salmonella typhi, Schigella flexenari, Escherichia coli, Staphylococcus aureus and Bacillus subtills in order to establish their

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Table 1. Synthesis of Schiff bases.

<table>
<thead>
<tr>
<th>Schiff base</th>
<th>R</th>
<th>R'</th>
<th>Mol. formula</th>
<th>Mol. wt</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>3- OCH(_3)</td>
<td>4-CH(_3)</td>
<td>C(<em>{15})H(</em>{15})NO</td>
<td>225</td>
</tr>
<tr>
<td>2</td>
<td>3, 4- OCH(_3)</td>
<td>4-CH(_3)</td>
<td>C(<em>{16})H(</em>{17})NO(_2)</td>
<td>255</td>
</tr>
<tr>
<td>3</td>
<td>3, 4, 5-OCH(_3)</td>
<td>4-CH(_3)</td>
<td>C(<em>{17})H(</em>{19})NO(_3)</td>
<td>285</td>
</tr>
<tr>
<td>4</td>
<td>3-OCH(_3)_3, 4-OH</td>
<td>4-CH(_3)</td>
<td>C(<em>{15})H(</em>{15})NO</td>
<td>241</td>
</tr>
<tr>
<td>5</td>
<td>4-F</td>
<td>4-CH(_3)</td>
<td>C(<em>{14})H(</em>{12})FNO</td>
<td>213</td>
</tr>
<tr>
<td>6</td>
<td>4-Cl</td>
<td>4-CH(_3)</td>
<td>C(<em>{14})H(</em>{12})Cl N</td>
<td>229</td>
</tr>
<tr>
<td>7</td>
<td>4-Br</td>
<td>4-CH(_3)</td>
<td>C(<em>{14})H(</em>{12})BrN</td>
<td>273</td>
</tr>
<tr>
<td>8</td>
<td>4-I</td>
<td>4-CH(_3)</td>
<td>C(<em>{14})H(</em>{12})I N</td>
<td>321</td>
</tr>
</tbody>
</table>
bioactivities. In order to compare the results obtained, the impinem was used as a standard drug. The result obtained showed that synthesized bases have significant activity but lower than the standard drug. The result indicate that the base 1, 2 and 4 showed promising activity because it is proposed that oxygen of methoxy group may bind with bacteria and hence play an important role in enhancing the activity. Among these three, compound 4 has greater activity because of the hydroxyl group, which enhances the biological activity (Rehman et. al., 2005); base 6 was found to be of moderate activity. However Schiff base 3, 5, 7 and 8 showed somewhat less inhibition than others. The anti-bacterial activity results of these bases are presented in Table 3.

**Minimum inhibitory concentration (MIC)**

The MIC values for Schiff bases against *B. subtilis*, *S. aureus*, *E. coli*, *S. typhi* and *P. aeruginosa* were 18, 20, 22, 18 and 2 µg/ml for Schiff base 1; 16, 14, 18 and 16 µg/ml for Schiff base 2; 40, 34, 36, 40 and 38 µg/ml for Schiff base 3; 9, 9, 12, 8 and 8 µg/ml for Schiff base 4; 60, 58, 64, 60 and 64 µg/ml for Schiff base 5; 32, 28, 26, 30 and 32 µg/ml for Schiff base 6; 68, 72, 64, 64 and 68 µg/ml for Schiff base 7; 128, 120, 124, 120 and 128 µg/ml for Schiff base 8, respectively (Table 4). The values of MIC showed the following order: 4 > 2 > 1 > 6 > 3 > 5 > 7 > 8, among these Schiff bases, the Schiff base 4 exhibited excellent MIC result as it is proposed that the

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### Table 2. Brine shrimp bioassay of Schiff bases.

<table>
<thead>
<tr>
<th>Schiff base</th>
<th>% Age of mortality at doses</th>
<th>LD₅₀</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>50 (µg/ml)</td>
<td>100 (µg/ml)</td>
</tr>
<tr>
<td>1</td>
<td>4</td>
<td>9</td>
</tr>
<tr>
<td>2</td>
<td>75</td>
<td>80</td>
</tr>
<tr>
<td>3</td>
<td>8</td>
<td>7</td>
</tr>
<tr>
<td>4</td>
<td>85</td>
<td>15</td>
</tr>
<tr>
<td>5</td>
<td>34</td>
<td>52</td>
</tr>
<tr>
<td>6</td>
<td>37</td>
<td>95</td>
</tr>
<tr>
<td>7</td>
<td>10</td>
<td>20</td>
</tr>
</tbody>
</table>

### Table 3. Anti bacterial activity results of Schiff bases.

<table>
<thead>
<tr>
<th>Bacteria</th>
<th>High concentration samples</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1</td>
</tr>
<tr>
<td><strong>Bacillus subtilis</strong></td>
<td>++</td>
</tr>
<tr>
<td><strong>Staphylococcus aureus</strong></td>
<td>+++</td>
</tr>
<tr>
<td><strong>Escherichia coli</strong></td>
<td>+</td>
</tr>
<tr>
<td><strong>Schigella flexenari</strong></td>
<td>n.a</td>
</tr>
<tr>
<td><strong>Pseudomonas aeruginosa</strong></td>
<td>++</td>
</tr>
<tr>
<td><strong>Salmonella typhi</strong></td>
<td>++</td>
</tr>
</tbody>
</table>

* Impinium; ++++, highly active; ++, good activity; +, moderate activity; +, significant activity; n.a, no activity; n.c, not checked.

### Table 4. Minimum inhibitory concentration (MIC) values for the Schiff bases.

<table>
<thead>
<tr>
<th>Microorganism</th>
<th>MIC (µg/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1</td>
</tr>
<tr>
<td><em>Bacillus subtilis</em></td>
<td>18</td>
</tr>
<tr>
<td><em>Staphylococcus aureus</em></td>
<td>20</td>
</tr>
<tr>
<td><em>Escherichia coli</em></td>
<td>22</td>
</tr>
<tr>
<td><em>Salmonella typhi</em></td>
<td>18</td>
</tr>
<tr>
<td><em>Pseudomonas aeruginosa</em></td>
<td>22</td>
</tr>
</tbody>
</table>
oxygen of methoxy may bind with the bacteria leading to excellent activity.

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REFERENCES


