

## Zirconium(IV) Chloride - Catalysed Reaction of Indoles: An Expeditious Synthesis of Bis(indolyl)methanes

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Zirconium(IV) chloride is found to be an efficient catalyst for the electrophilic addition reaction of indole with aldehydes/ketones to afford the corresponding bis(indolyl)methanes in good yields. The remarkable features of this new procedure are high conversions, shorter reaction times, cleaner reaction profiles and simple experimental and work-up procedures.

**Key Words :** Zirconium(IV) chloride, Aldehydes, Ketones, Bis(indolyl)methanes, Indole

### Introduction

Development of bis(indolyl)alkane synthesis has been of considerable interest in organic synthesis because of their wide occurrence in various natural products possessing biological activity<sup>1</sup> and usefulness for drug design.<sup>2</sup> Bis(indolyl)methanes are most active cruciferous substances for promoting beneficial estrogen metabolism and inducing apoptosis in human cancer cells.<sup>3</sup> Consequently, numerous methods have been reported for the synthesis of bis(indolyl)methanes.<sup>4</sup> Of these methods, the acid catalysed electrophilic addition reaction of indole with aldehydes is one of the most simple and straightforward approaches for the synthesis of bis(indolyl)methanes. A variety of reagents such as acetic acid,<sup>5</sup> InCl<sub>3</sub>,<sup>6</sup> In(OTf)<sub>3</sub>,<sup>7</sup> InF<sub>3</sub>,<sup>8</sup> Dy(OTf)<sub>3</sub>,<sup>9</sup> Ln(OTf)<sub>3</sub>,<sup>10</sup> LiClO<sub>4</sub>,<sup>11</sup> FeCl<sub>3</sub>,<sup>12</sup> I<sub>2</sub>,<sup>13</sup> NBS,<sup>14</sup> KHSO<sub>4</sub>,<sup>15</sup> NaHSO<sub>4</sub>·SiO<sub>2</sub>,<sup>16</sup> PPh<sub>3</sub>·HClO<sub>4</sub> (TPP),<sup>17</sup> CAN,<sup>18</sup> zeolites,<sup>19</sup> clay,<sup>20</sup> H<sub>3</sub>PMO<sub>12</sub>O<sub>40</sub>·xH<sub>2</sub>O,<sup>21</sup> and 1-butyl-3-methylimidazolium tetrafluoroborate or 1-butyl-3-methylimidazolium hexafluorophosphate ionic liquids<sup>22</sup> have been employed to accomplish this transformation. Recently, rare-earth perfluorooxanoates [RE(PFO)<sub>3</sub>],<sup>23</sup> trichloro-1,3,5-triazine,<sup>24</sup> hexamethylenetetramine-bromine,<sup>25</sup> ion-exchange resin<sup>26</sup> and ionic liquids in conjugation with In(OTf)<sub>3</sub> or FeCl<sub>3</sub>·6H<sub>2</sub>O<sup>27</sup> were also found to promote this reaction. However, some of the reported methods have the following drawbacks: for example, use of expensive reagents,<sup>7,9,10</sup> longer reaction times,<sup>5,6,8,22</sup> low yields of products<sup>5,20b</sup> and use of an additional microwave oven.<sup>12</sup> Because of their wide range of biological, industrial and synthetic applications, the preparation of bis(indolyl)methanes has received renewed interest

of researchers for the discovery of improved protocols and still awaits further development of milder and high-yielding processes.

Zirconium(IV) chloride is an efficient Lewis acid catalyst used in various transformations, such as, electrophilic amination of activated arenes,<sup>28</sup> transthiocetylation of acetals,<sup>29</sup> deoxygenation of heterocyclic-N-oxides,<sup>30</sup> reduction of nitro compounds,<sup>31</sup> conversion of carbonyl compounds to 1,3-oxathiolanes<sup>32</sup> and in Biginelli reaction for the synthesis of 3,4-dihydropyrimidin-2(1*H*)-ones.<sup>33</sup> However, there are no examples of the use of zirconium(IV) chloride as a catalyst for the synthesis of bis(indolyl)methanes.

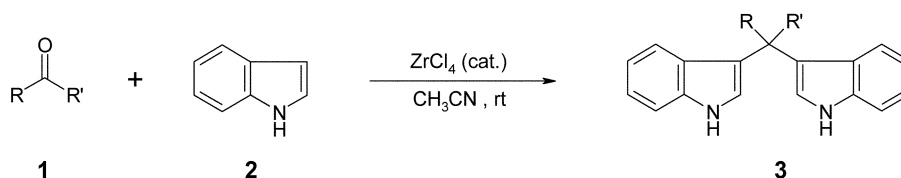
### Results and Discussion

Herein, we wish to disclose a novel protocol for the rapid synthesis of variety of biologically significant bis(indolyl)methanes using a catalytic amount of zirconium(IV)

**Table 1.** Screening of reaction conditions for the synthesis of **3a**

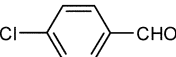
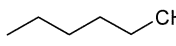
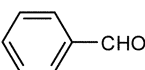
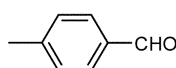
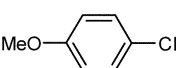
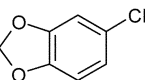
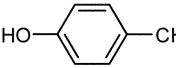
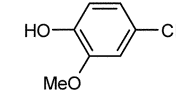
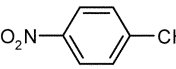
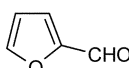
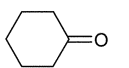
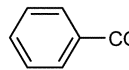
Entry	Solvent	Catalyst	Time (hr.)	Yield (%) <sup>a</sup>
1	CH <sub>3</sub> CN	5% ZrCl <sub>4</sub>	0.35	92
2	AcOEt	5% ZrCl <sub>4</sub>	2	86
3	CH <sub>2</sub> Cl <sub>2</sub>	5% ZrCl <sub>4</sub>	3	84
4	MeOH	5% ZrCl <sub>4</sub>	3	78
5	EtOH	5% ZrCl <sub>4</sub>	2	76
6	THF	5% ZrCl <sub>4</sub>	5	74
7	DMF	5% ZrCl <sub>4</sub>	7	55
8	CH <sub>3</sub> CN	1% ZrCl <sub>4</sub>	0.75	85

<sup>a</sup>isolated yields



**Scheme 1**

**Table 2.** ZrCl<sub>4</sub>-Promoted Synthesis of Bis(indolyl)methane derivatives

Entry	Substrate	Time (min)	Yield %	M.P. (°C)	
				Found	Reported
a		20	92	76-77	77-81 <sup>26</sup>
b		65	86	68-70	70-72 <sup>26</sup>
c		35	91	123-125	125-127 <sup>20b</sup>
d		40	95	96-98	94-96 <sup>27b</sup>
e		45	90	190-192	187-189 <sup>27b</sup>
f		35	88	100-102	97-99 <sup>11</sup>
g		40	86	124-125	122-124 <sup>18</sup>
h		45	92	111-112	110-112 <sup>19b</sup>
i		15	90	221-223	220-222 <sup>26</sup>
j		40	89	320-323	322 <sup>25</sup>
k		310	71	118-120	118-120 <sup>11</sup>
l		325	56	189-190	190-192 <sup>18</sup>

chloride under extremely mild conditions (Scheme 1). Firstly, 4-chlorobenzaldehyde (**1a**) was chosen as a model for the reaction with indole. Compound **1a** was treated with 2.0 equivalent of indole in the presence of 5 mol % ZrCl<sub>4</sub> (based on the amount of indole) at room temperature in various solvents (Table 1, entries 1-7). In CH<sub>3</sub>CN, the reac-

tion completed within 20 minutes to give bis(indolyl)methane **3a** in 92% yield. The reactions in CH<sub>2</sub>Cl<sub>2</sub>, AcOEt, MeOH, EtOH, DMF and THF required longer reaction times and unreacted **1a** and indole remained. Lower catalyst loading can be used with only a marginal drop in reaction rate (Table 1, entry 8).

Having established the reaction conditions, various aldehydes **1** were reacted with indole **2** to investigate the reaction scope, and several representative examples are summarized in Table 2. Aromatic, aliphatic (entry **b**) and heterocyclic aldehyde (entry **j**) underwent smooth transformation to the corresponding bis(indolyl)methanes in high to excellent yield in relatively shorter reaction times. It has been observed that the electronic properties of the aromatic ring have an effect on the rate of this electrophilic addition reaction. The rate is accelerated if an electron withdrawing group is present on the aromatic ring. The reaction of ketones with indole took longer time when compared with aldehydes, and unreacted ketone and indole remained. It is also found that the reaction of terephthalaldehyde **4** with 4 equivalent of indole proceeded rapidly to give *p*-di(bis-indolylmethane)benzene **5** in similar conditions in high yield (Scheme 2).

In summary, the electrophilic addition reaction of indole with aldehydes was successfully carried out in the presence of catalytic amount of ZrCl<sub>4</sub> at room temperature. This method offers several significant advantages, such as, high conversions, easy handling, cleaner reaction profile and shorter reaction times, which makes it a useful and attractive process for the rapid synthesis of substituted bis(indolyl)methanes.

### Experimental Section

Melting points are uncorrected. <sup>1</sup>H-NMR and <sup>13</sup>C-NMR spectra were recorded on Varian Gemini 200 MHz spectrometer. Chemical shifts are reported in  $\delta$  units (ppm) relative to TMS as internal standard. Electron spray ionization mass spectra (ES-MS) were recorded on Water-Micromass Quattro-II spectrometer. IR spectra were recorded on Varian spectrometer. All the reagents used were of AR grade and were used without further purification. Column chromatography employed silica gel of 60-120 mesh.

**General Procedure.** A mixture of indole (2 mmol), aldehyde or ketone (1 mmol) and ZrCl<sub>4</sub> (0.1 mmol) in acetonitrile was stirred at room temperature for the appropriate time (Table 2). The progress of the reaction was monitored

**Scheme 2**

by TLC. After completion of the reaction, as indicated by TLC, the reaction mixture was quenched with water and extracted with ethyl acetate (3 × 10 mL). The combined organic layers were dried (Na<sub>2</sub>SO<sub>4</sub>) and concentrated in vacuo to afford the crude compound. The crude compounds were purified by silica gel column chromatography using ethyl acetate/hexane as eluent, to afford the desired compound in pure form. All the synthesized compounds were characterized by I.R., <sup>1</sup>H-NMR, <sup>13</sup>C-NMR, Mass (ES-MS), elemental analysis and melting point.

**4-Chlorophenyl-3,3'-bis(indolyl)methane (3a).** Yield 92%; m.p. 76-77 °C; <sup>1</sup>H-NMR (200 MHz, CDCl<sub>3</sub>): 5.86 (s, 1H, Ar-CH), 6.65 (s, 2H), 7.02 (t, 2H, *J* = 7.6 Hz), 7.18 (t, 2H, *J* = 7.6 Hz), 7.26-7.38 (m, 8H), 7.93 (br s, 2H, NH); <sup>13</sup>C-NMR (50 MHz, CDCl<sub>3</sub>): 46.5, 100.6, 110.1, 118.0, 119.1, 121.2, 127.0, 127.8, 129.5, 130.3, 134.2, 135.2, 135.6; ES-MS *m/z* 357 (M+H, 100%); Analysis Calcd. for C<sub>23</sub>H<sub>17</sub>ClN<sub>2</sub>: C, 77.41; H, 4.80; N, 7.85. Found: C, 77.55; H, 4.73; N, 7.90.

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