A highly efficient, eco-friendly, room temperature synthesis of bis(indol-3-yl)methanes using the mesoporous titanosilicate Ti-TUD-1: electrophilic substitution reactions of indoles – Part XXIII

Bikash Karmakar,^a Anupam Nayak,^a Biswajit Chowdhury ^{b*} and Julie Banerji^a*

^aCentre of Advanced Studies on Natural Products including Organic Synthesis, Department of Chemistry, University of Calcutta, 92, A.P.C. Road, Kolkata-700009, India ^bDepartment of Applied Chemistry, Indian School of Mines University, Dhanbad-826004, India E-mail: juliebanerji47@gmail.com; biswajit_chem2003@yahoo.com

Abstract

Bis(indol-3-yl)methanes have been synthesized at room temperature in high yields by the reaction of indole or 2-methylindole with aromatic aldehydes using the titanosilicate catalyst Ti-TUD-1 for the first time. The catalyst was recycled and reused five times with reproducible yields. This procedure has the advantage of high yields at room temperature, mildness of reaction conditions, operational simplicity and reusable nature of the catalyst which makes this protocol practically useful for the synthesis of bis(indol-3-yl)methanes.

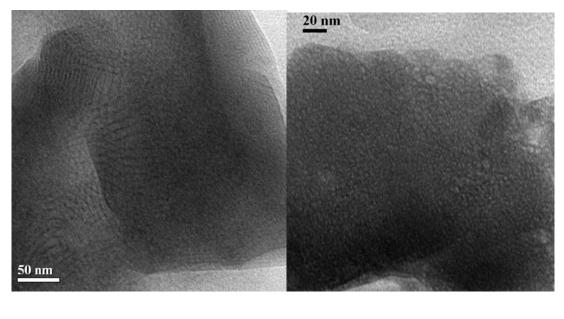
Keywords: Titanosilicate, bis(indol-3-yl)methane, room temperature, recyclable.

Introduction

Ever since the biochemical¹ and clinical² use of various indole derivatives viz., bis(indol-3yl)methane,³ vibrindole A⁴ and streptindole⁵ was reported, attention has been focused on the development of cost-effective, simple operational procedures and high yield routes to bis(indol-3-yl)methanes (BIMs) and substituted bis(indol-3-yl)methanes.⁶ In this communication we report, for the first time, the use of Ti-TUD-1, a mesoporous titanosilicate⁷ catalyst for the synthesis of BIMs from the reaction of indoles with various aldehydes at room temperature. We have exploited the high surface area, larger pores, high Lewis acidic character and reusable nature of the catalyst in synthesizing bis(indol-3-yl)methanes with high chemoselectivity and in quantitative yields following a clean and green pathway.

Results and Discussion

Mesoporous Ti-TUD-1 (Ti/Si mole ratio = 0.01, 0.03, 0.05, named as catalyst x, y, z respectively) catalyst was prepared by non-hydrothermal sol-gel technique and was modified from that followed by T. Maschmeyer *et al.*⁷ It was then characterized by Transmission Electron Microscopy (TEM) and surface area analysis by Brunauer, Emmett and Teller (BET) method. TEM images (Figure 1) of the catalyst clearly show that the surface is highly porous in nature and is of a spongy type. There are mixtures of small and large pores well distributed over the whole surface. The pore sizes range between 5-15 nm. The porous channels formed are also visible in the TEM image. Due to the high porosity its surface area is also found to be high. BET measurement showed that the surface areas range between 380-450 m²/g for the three catalysts. The mesoporosity was also confirmed from X-ray powder diffraction (XRD) studies. The high surface area ensured greater accessibility of the substrates to the active sites of the catalyst.



(A)

(B)

Figure 1. TEM images of the lamellar channels (A) and porous surface (B) of the catalyst

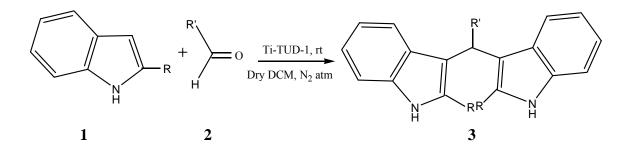
Due to its Lewis acid character, enhanced by the electron pull of the Si atom in the Ti-O-Si bond, and large surface area, Ti-TUD-1 could be effectively used as catalyst for the electrophilic substitution reaction of indoles with aromatic aldehydes for the formation of BIMs (Scheme 1). In order to confirm the efficiency of this catalyst, a comparative study was carried out with a

variety of catalysts along with the Ti-TUD-1 with different Ti loadings and the results are summarized in Table 1. Indole was treated with benzaldehyde as a model reaction in the presence of the catalyst in dry DCM and under N_2 flow. The reaction did not proceed in the absence of any catalyst. Also, with unloaded mesoporous silicate TUD-1 and bulk TiO₂ as catalysts, no product was obtained. Other titanosilicates like Ti-MCM-48 and TS-1 did not prove to be highly efficient in this reaction. Ti-TUD-1 catalysts with different Ti/Si content were studied and it was observed that with a Ti/Si mol ratio 0.03 (catalyst y) provided the best result. The optimization of the conditions was continued using the best catalyst with different solvent variations. We tried several solvents of variable polarity and the observations are shown in Table 2. The reaction proceeded smoothly with the best yield (89%) in dry dichloromethane. Other solvents like toluene, chloroform, acetonitrile and tetrahydrofuran resulted in low to moderate yields. In the absence of solvent, no product was obtained.

Entry	Catalyst	Time (h)	Yield ^b (%)
1	None	24	_
2	TUD-1	24	_
3	TiO ₂	24	_
4	Ti-MCM-48	24	57
5	Ti-TUD-1 (x)	6	85
6	Ti-TUD-1 (y)	6	89
7	Ti-TUD-1 (z)	6	78
8	TS-1	12	48

Table 1. Study of catalyst variation in the synthesis of bis(indol-3-yl)methanes^a

^a reaction conditions: 2 mmol benzaldehyde, 4 mmol indole, 10 mg catalyst, room temperature. ^b isolated yield after purification



Scheme 1. Mesoporous titanosilicate catalyzed synthesis of bis(indol-3-yl)methanes 3.

Entry	Solvent	Time (h)	Yield ^b (%)
1	None	24	_
2	Dichloromethane	6	89
3	Chloroform	12	35
4	Toluene	24	15
5	Acetonitrile	12	52
6	Tetrahydrofuran	12	32

Table 2. Use of Ti-TUD-1 catalyst in various solvents^a

^a reaction conditions: 2 mmol benzaldehyde, 4 mmol indole, 10 mg catalyst, room temperature. ^b isolated yield after purification.

A series of reactions were carried to study the scope and generality of Ti-TUD-1 (y) catalyst using various aromatic aldehydes and two indoles. The results are shown in Table 3. The bis(indol-3-yl)methanes **3a-o** were synthesized utilizing indole **1a** and 2-methylindole **1b** with a variety of aromatic aldehydes **2a-l** under the optimum conditions. In all the cases, BIMs were obtained as the sole products. Most of the aldehydes reacted with indoles to produce the bis(indol-3-yl)methanes in high yields. The effect on the yields of electron-withdrawing or electron-donating substituents on the aldehydes is shown in Table 3 (entries 2 and 7). It was observed that the reaction proceeded faster and in better yields with 2-methylindole **1b** than with indole **1a** possibly due to higher electron density at C-3. With indole, the reaction time was between 6-8 hours, while with 2-methylindole the reaction was complete within 45-120 minutes depending upon the electrophiles used. However, the catalyst did not show any activity towards ketones, viz., acetone **2m**, acetophenone **2n** and cyclohexanone **2o**. After prolonged reaction time at room temperature as well as on heating up to 60 °C both starting materials remained unchanged. This can be attributed to the lower electrophilicity of ketones compared to aldehydes.

Entry	R	R'	Product 3	Yield ^b (%)
1	H (1a)	$C_{6}H_{5}$ (2a)	3 a	89
2	H (1a)	$(4-NO_2) C_6 H_4 (2b)$	3 b	94
3	H (1a)	$(4-Me) C_6 H_4 (2c)$	3c	78
4	H (1a)	$(4-Cl) C_6 H_4 (2d)$	3d	84
5	H (1a)	$(3-NO_2) C_6 H_4 (2e)$	3e	91
6	H (1a)	(4-OH) C ₆ H ₄ (2f)	3f	71
7	H (1a)	$(4-NMe_2) C_6H_4 (2g)$	3g	64
8	H (1a)	$(4-OMe) C_6H_4 (2h)$	3h	73
9	H (1a)	(4-OH) (3-OMe) C ₆ H ₃ (2i)	3i	62
10	H (1a)	(3, 4-di OMe) C ₆ H ₃ (2j)	3j	75
11	H (1a)	(3, 4, 5-tri OMe) C ₆ H ₂ (2k)	3k	78
12	Me (1b)	$C_{6}H_{5}$ (2a)	31	91
13	Me (1b)	$(4-NO_2) C_6 H_4 (2b)$	3 m	96
14	Me (1b)	$(4-Cl) C_6 H_4 (2d)$	3n	86
15	Me (1b)	furoyl (2l)	30	88
16	H (1a)	acetone (2m)	_	_
17	H (1a)	cyclohexanone (2n)	_	_
18	H (1a)	acetophenone (20)	_	_

Table 3. Synthesis of bis(indol-3-yl)methanes over mesoporous titanosilicate Ti-TUD-1^a

^a reaction conditions: 2 mmol carbonyl compound, 4 mmol indole, 10 mg catalyst, room temperature.

^b isolated yield after purification.

Ti-TUD-1 is a very stable catalyst and can be recycled. It was reused for five consecutive times after being dried at 100 °C for 2 hours. No appreciable loss in its catalytic activity was observed as is evident from the yields (Figure 2). The high catalytic activity of Ti-TUD-1 is probably due to the high accessibility of the substrate molecules to the catalytic sites of the mesoporous system where Ti⁴⁺ is in tetrahedral coordination and being symmetrically dispersed over the three dimensional surface. Regarding the mechanism, it can be presumed that Ti-O-Si linkages at the surface bind with the aldehyde oxygen and increase the carbonyl activity. Indole then attacks the activated carbonyl to generate an indolenium cation. This intermediate, being unstable, is attacked by a second indole molecule and subsequent loss of water resulting in the formation of bis(indol-3-yl)methane.

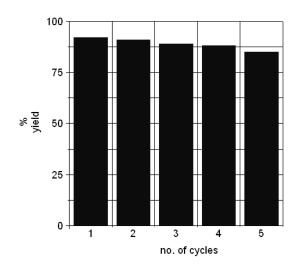


Figure 2. Recycling of the catalyst^a

^a reaction conditions: 2 mmol benzaldehyde, 4 mmol indole, 10 mg catalyst, room temperature, isolated yield.

Experimental Section

All chemicals were of analytical grade and used as such without further purification. The solvents were purified by distillation. All IR spectra were recorded on Perkin Elmer RX-1 FTIR spectrophotometer. ¹H NMR and ¹³C NMR were done on Bruker-Avance Digital 300 MHz and 75.5 MHz Spectrometer. ¹H NMR and ¹³C NMR were done on Bruker-Avance Digital 300 MHz and 75.5 MHz Spectrometer. TMS was used as internal standard. A JEOL JMS 600 mass spectrometer was used to record the HRMS. TEM images were obtained from a Hitachi H-9000 NAR Transmission Electron Microscope at an operating voltage of 100 kV. The sample was prepared by placing one drop dispersed solution of the catalyst in acetone solvent on a carbon coated copper grid followed by drying in air. Melting points (uncorrected) were determined on a Köfler Block apparatus. Analytical TLC was performed using E. Merck aluminium-backed silica gel plates coated with silica gel G and monitored under UV light (254 and 360 nm) and also by exposing to iodine. Indole, aldehydes and ketones were purchased from Merck.

Typical procedure for the preparation of modified Ti-TUD-1 catalyst

A modified sol-gel method was used to prepare mesoporous titanosilicate catalyst. Tetraethyl orthosilicate (TEOS) (Acros 99.5%) and titanium (IV) butoxide (Acros 99%) were used as the precursor for silica and titanium respectively. In the typical procedure, titanium butoxide was added slowly to the TEOS solution. A mixture of triethanolamine (TEA) (Acros 99%) and deionised water was added to a stirring solution and then triethyl ammonium hydroxide

(TEAOH) (E-Merck) (20% aqueous solution) was added dropwise to maintain pH 9. The final molar ratio of the reagents was 1.0 SiO₂: 0.03 TiO₂: 3.0 TEA: 0.3 TEAOH: 15 H₂O. The solution was aged for 24 hours at room temperature. It was then dried at 115 °C for 16 hours and finally calcined at 700 °C for 12 hours at the rate of 2 °C/minute under the flow of air.

Typical procedure for the synthesis of bis(indol-3-yl)methanes

A mixture of benzaldehyde (98 mg, 2 mmol) and indole (200 mg, 4 mmol) were taken in a 50 mL round-bottom flask containing 5 mL dry dichloromethane. Ti-TUD-1 catalyst (10 mg) was then added and the reaction mixture stirred continuously at room temperature under nitrogen atmosphere till the full consumption of starting materials was observed (TLC). After the completion of reaction the catalyst was filtered carefully. The residue obtained was washed with ethyl acetate, dried at 100 °C and reused as catalyst. The filtrate was evaporated under reduced pressure to give the crude product which was purified by column chromatography using neutral alumina and appropriate ethyl acetate/petroleum ether mixtures as eluent. All the isolated products were known. They were characterized by melting point measurement, IR spectroscopy, ¹H NMR, ¹³C NMR and HRMS spectrometry and elemental analysis (C, H, N) and were compared to their authentic data.⁸

Acknowledgements

J.B. and A.N. thank UGC (New Delhi) for providing the funds. BC also thanks Fast Track Young Scientist Scheme under DST (New Delhi) for financial assistance. BK thanks Mr. P. Ghosh for Instrumnetal analysis at Calcutta University and IIT, Kharagpur for TEM analysis.

References

- 1. (a) Lo, K. K.-W.; Tsang, K. H.-K.; Hui, W.-K.; Zhu, N. Chem. Commun. 2003, 2704. (b) Ramirez, A.; Garcia-Rubio, S. Curr. Med. Chem. 2003, 10, 1891.
- (a) Le, H. T.; Schaldach, C. M.; Firestone, G. L.; Bjeldanes, L. F. *J. Biol. Chem.* 2003, 278, 21136.
 (b) Dalessandri, K. M.; Firestone, G. L.; Fitch, M. D.; Bradlow, H. L.; Bjeldanes, L. F. *Nutr. Cancer.* 2004, *50*, 161.
- (a) Maciejewska, D.; Rasztawicka, M.; Wolska, I.; Anuszewska, E.; Gruber, B. *Eur. J. Med. Chem.* 2009, 44, 4136. (b) Karthik, M.; Tripathi, A. K.; Gupta, N. M.; Palanichamy, M.; Murugeson, V. *Catal. Commun.* 2004, 5, 371. (c) Gribble, G. W. *J. Chem. Soc., Perkin Trans 1*, 2000, 1045.
- 4. Deb, M. L.; Bhuyan, P. J. Synthesis 2008, 2891.

- (a) Bartoli, G.; Bosco, M.; Foglia, G.; Giuliani, A.; Marcantoni, E.; Sambri, L. Synthesis 2004, 895. (b) Chakrabarty, M.; Ghosh, N.; Basak, R.; Harigaya, Y. Synth. Commun. 2004, 34, 421.
- 6. (a) Pindur, U.; Adam, R. J. Heterocycl. Chem. 1988, 25, 1. (b) Rainer, J. D.; Smith, A. B. Tetrahedron Lett. 2000, 41, 9419. (c) Bennasar, M.-L.; Vidal, B.; Bosch, J. J. Org. Chem. **1997**, 62, 3597. (d) Amat, M.; Hadida, S.; Pshenichnyi, G.; Bosch, J. J. Org. Chem. **1997**, 62, 3158. (e) Azizian, J.; Teimouri, M. R.; Mohammadizadeh, M. R. Catal. Commun. 2007, 8, 1117. (f) Ramesh, R.; Banerjee, J.; Pal, R.; Das, B. Adv. Synth. Catal. 2003, 345, 557. (g) Chakrabarty, M.; Basak, R.; Harigaya, Y.; Ghosh, N. Tetrahedron Lett. 2002, 43, 4075. (h) Maiti, A. K.; Bhattacharyya, P. J. Chem Res. (S) 1997, 425. (i) Li, J.; Zhou, M.; Li, B.; Zhang, G. Synth. Commun. 2004, 34, 275. (j) Ji, S. -J.; Zhou, M. -F.; Gu, D. -G.; Wang, S. -Y.; Loh, T. -P. Synlett 2003, 13, 2077. (k) Ji, S. -J.; Zhou, M. -F.; Gu, D. -G.; Ziang, Z. -Q.; Loh, T. -P. Eur J. Org. Chem. 2004, 1584. (1) Yadav, J. S.; Reddy, B. V. S.; Sunitha, S. Adv. Synth. Catal. 2003, 345, 349. (m) Bandgar, B. P.; Shaikh, K. A. Tetrahedron Lett. 2003, 44, 1959. (n) Yadav, J. S.; Reddy, B. V. S.; Murthy, Ch. V. S. R.; Kumar, G. M.; Madan, Ch. Synthesis 2001, 783. (o) Firouzabadi, H.; Iranpoor, N.; Javari, A. A. J. Mol. Catal. A: Chemical 2006, 244, 168. (p) Chatterjee, A.; Manna, S.; Banerji, J.; Pascard, C.; Prange, T.; Shoolery, J.; James, N. J. Chem. Soc., Perkin Trans. 1 1980, 553. (g) Chen, D.; Yu, L.; Wang, P. G. Tetrahedron Lett. 1996, 37, 4467. (r) Wang, L.; Han, J.; Tian, H.; Sheng, J.; Fan, Z.; Tang, X. Synlett 2005, 2, 337. (s) Ramesh, C.; Ravindranath, N.; Das, B. J. Chem Res. (S) 2003, 72. (t) Zolfigol, M.; Salehi, P.; Shiri, M. Phosphorous, Sulfur, Silicon Rel. *Elem.* 2004, 179, 2273. (u) Bavanthula, R; Reddy, P. N; Kumar, B. S; Neeladri, S; Reddy, Y; Reddy, T. J. Chem Res. 2005, 222. (v) Seyedi, N.; Saidi, K.; Khabazzadeh, H. Synth. Commun. 2009, 39, 1864. (w) Praveen, C; Wilson Sagayaraj, Y.; Perumal, P. T.; Tetrahedron Lett. 2009, 50, 644. (x) Sabitha, G.; Reddy, N. M.; Prasad, M. N.; Yadav, J. S.; Sivudu, K. S.; Shailaja, D. Lett. Org. Chem. 2008, 5, 300. (y) Nadkarni, S. V.; Gawande, M. B.; Jayaram, R. V.; Nagarkar, J. M. Catal. Commun. 2008, 9, 1728. (z) Khalafi-Nehzad, A.; Parhami, A.; Zare, A.; Zare, A. R. Moosavi; Hasaninejad, A.; Panahi, F. Synthesis, 2008, 617.
- Shan, Z; Gianotti, E; Jansen, J. C; Peters, J. A; Marchese, L; Maschmeyer, T. *Chem. Eur. J.* 2001, 7, 1437.
- (a) Selvam, J. J. P.; Srinivasulu, M.; Suryakiran, N.; Suresh, V.; Malla Reddy, S.; Venkatesarlu, Y. *Synth. Commun.* 2008, *38*, 1760. (b) Sobhani, S.; Safaei, E.; Hasaninejad, A.-R.; Rezazadeh, S. *J. Organomet. Chem.* 2009, *694*, 3027. (c) Sadaphal, S. A.; Shelke, K. F.; Sonar, S. S.; Shingare, M. S. *Eur. J. Chem.* 2008, *6*, 622. (d) Zahran, M.; Abdin, Y.; Salama, H. *Arkivoc* 2008, 256. (e) Teimouri, M. B.; Mivehchi, H. *Synth. Commun.* 2005, *35*, 1835.