

Synthesis and characterization of new chiral ketopinic acid-derived catalysts immobilized on polystyrene-bound imidazole

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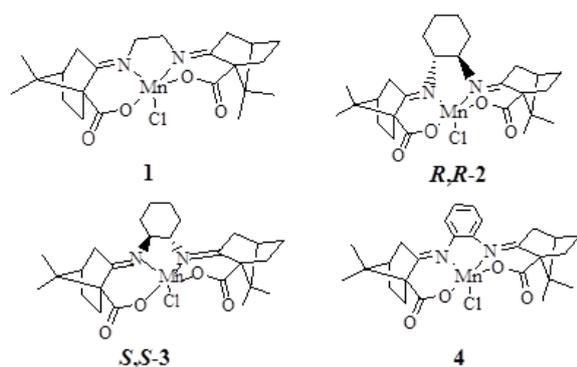
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Abstract. Four new chiral ketopinic acid-derived catalysts were anchored on a polystyrene-bound imidazole via non-covalent bond. The resulting heterogeneous catalysts were successfully characterized using IR, SEM, and TGA analyses.

Keywords: polymer, chiral catalysts, non-covalent bond.

1. Introduction

The ability of new chiral catalysts **1-4** (Scheme 1) derived from ketopinic acid to effect different asymmetric transformations, viz. aldol, Baylis-Hillman, aza-Michael, and reduction reactions was demonstrated [1]. These are widely employed methodologies in the syntheses of natural products and pharmaceutically important compounds [2-12]. However, the fact that these catalysts were homogenous provides the opportunity to explore methods of anchoring them onto a solid support to render them heterogeneous [13]. Heterogeneous catalysis has the potential advantages of easy separation, efficient recycling, and the minimization of metal trace in the product. There are also improved handling and process control issues that could, in principle, result in an overall reduction of costs and waste [14, 15].



Scheme 1. Manganese chiral ketopinic acid-derived catalysts.

2. Experimental

Reagent-grade acetonitrile, ether, ethylacetate, and methanol were used as received from commercial source. All other reagents were purchased from Aldrich and used as received. IR spectral measurements were carried out on a Perkin Elmer spectrum 400 FT-IR spectrometer (ATR). Thermogravimetric analyses of the catalysts were performed on Perkin Elmer 2000 TGA 7. Scanning electron micrographs of the catalysts were taken on a TESCAN VEGAN III XMU Q150 TE.

Synthesis of chiral ketopinic acid-derived catalysts 1-4. Catalysts **1-4** were prepared according to the standard procedure [16, 17].

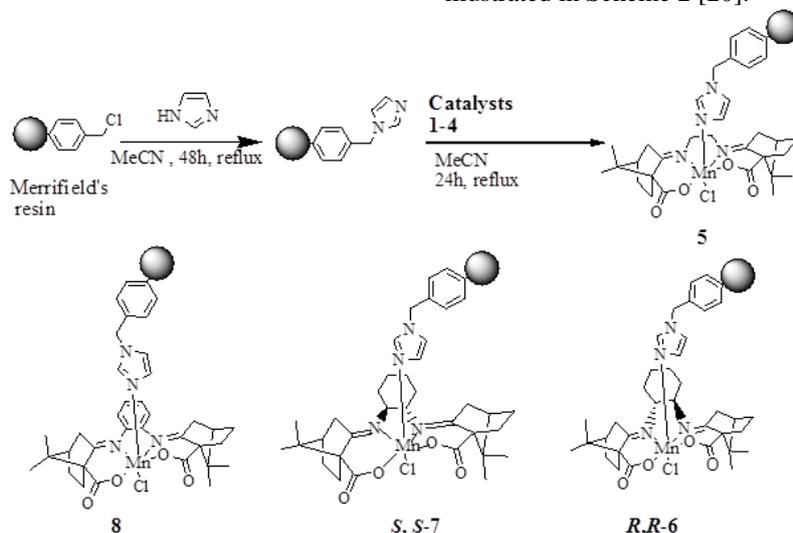
Synthesis of polymer-supported catalysts 5-8.
Polymer-bound imidazole: To a slurry of the Merrifield resin (1 g, 1 mmol) in acetonitrile (6.5 mL) were added imidazole (0.68 g, 10 mmol) and NaI (0.01 g, 0.07 mmol) and the mixture was refluxed with stirring for 48 h. The resin was filtered off and washed successively with acetonitrile (5×5 mL), 10% aq. K₂CO₃/MeOH (1:1), H₂O/MeOH (1:1), MeOH and EtOAc (5×10 mL each). The solid was dried under vacuum overnight to yield polymer-bound imidazole (1.00 g, 99%). IR (neat, cm⁻¹): 3025, 2919, 1492, 1451, 1265. *General procedure for the immobilization of the catalysts:* To a solution of 50 mg each of catalysts **1-4** in acetonitrile (5 mL) was added polymer-bound imidazole (375 mg). The mixture was vigorously stirred at 80 °C for 24 h. After cooling, the brown resin was collected by filtration, washed thoroughly with acetonitrile, methanol, and ether successively. It was then dried in vacuum at room temperature for several hours to afford complexes; **5** (93%), **R,R-6** (89%), **S,S-7** (83%), and **8** (90%) yield. SEM was used for characterization based on morphological changes.

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3. Results and Discussion

The aim was to incorporate catalysts **1-4** onto the Merrifield resin [18] by adopting the non-covalent immobilization approach [19]. A nitrogenous base (imidazole) would be tethered to the Merrifield resin

and subsequently coordinated to the manganese atom of the chiral complex. The polymer-supported imidazole was expected to offer both the heterogeneous scaffold and provide an axial ligand in the corresponding catalytic complexes **5-8** as illustrated in Scheme 2 [20].



Scheme 2. Immobilization of the chiral catalysts **1-4** onto a polystyrene-bound imidazole.

The first step of the synthesis of the polystyrene-supported complexes **5-8** was the treatment of the Merrifield resin with an appropriate quantity of imidazole, in refluxing acetonitrile for 48 h [11]. After cooling, the resin was filtered off and washed with acetonitrile, mixture of aq. $K_2CO_3/MeOH$ and $H_2O/MeOH$ followed by $MeOH$ and then $EtOAc$. The solid was dried under vacuum overnight and afforded the product in 99% yield. Elemental analysis of the isolated polystyrene-supported imidazole (PSI) revealed that the nitrogen content was 2.01% which was equivalent to 51% of the chlorine substituted. Confirmation of the presence of the imidazolyl-methylene bond was explored using IR spectroscopy (Figure 1).

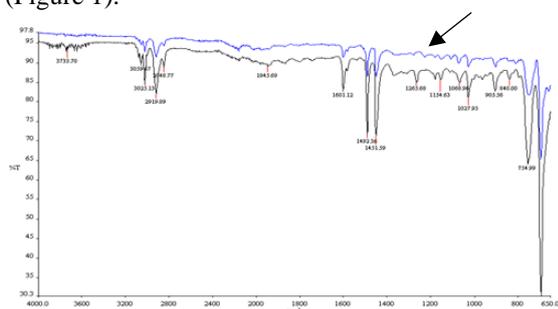


Figure 1. Overlaid IR spectra of Merrifield's resin and the PSI.

Due to the similarity between the Merrifield resin and the PSI in the mid-IR range, the two spectra were overlaid to carefully ascertain the band differences in this region. The lower spectrum is for the Merrifield resin and exhibits a weak band due to the C-Cl bond at 1265.68 cm^{-1} . Upon nucleophilic substitution of

chloride by the imidazole nitrogen, the weak band has largely disappeared (top spectrum).

The supported chiral complex was then prepared by stirring a suspension of the PSI, in solutions of the metal complexes **1-4** in CH_3CN as solvent [11] as shown on Scheme 2. After stirring at an elevated temperature ($80\text{ }^\circ\text{C}$) for 24 h, the mixture was cooled and, in each case, the resins were filtered off, washed sequentially with acetonitrile, methanol, and ether. Organic solvent washings were checked by IR until no unreacted complex was detected, and thus confirming the stability of the coordinative bond, to the PSI. Surface characterization of the supported catalysts was performed using Scanning Electron Microscopy (SEM). The images show some morphological changes on the surface of the polymer resin beads. SEM images of the Merrifield resin and immobilized catalyst **5** are shown in Figures 2 and 3 respectively.

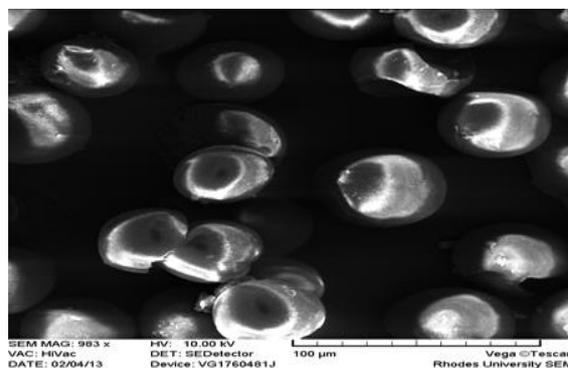


Figure 2. SEM image of Merrifield's resin.

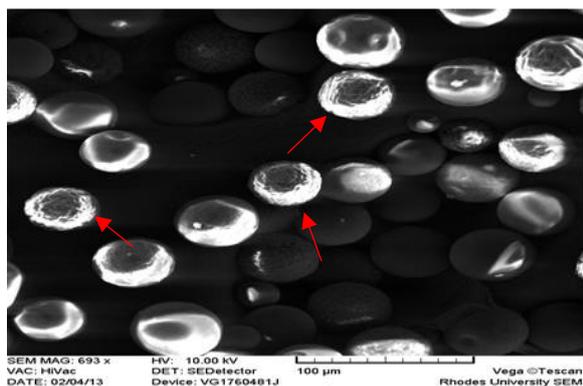


Figure 3. SEM image of the PSI-catalyst **5** with arrows indicating surface changes in the beads.

Finally, thermogravimetric analysis (TGA) was conducted to ascertain the stability of the supported catalysts [19]. The Merrifield resin shows single step degradation at approximately 280 °C (Figure 4). The supported catalysts on the other hand degrade at lower temperatures of approximately 200 °C as illustrated by the representative TGA curve of catalyst **5** (Figure 5). This corresponds to either the dissociation of the coordinative bound manganese complex or a partial scission of the polymeric chain. However, it seems that the supported catalysts are thermally stable up to 200 °C.

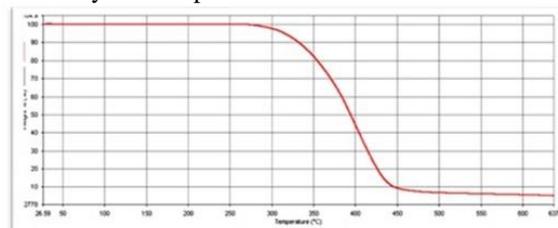


Figure 4. TGA curve of Merrifield's resin.

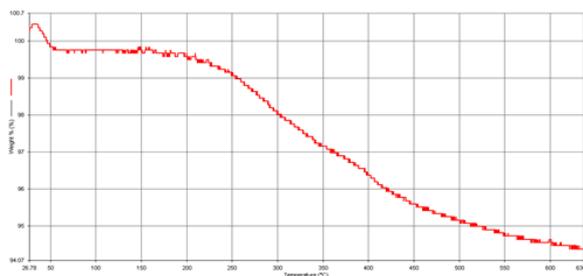


Figure 5. TGA curve of PSI-catalyst **5**.

4. Conclusions

Chiral ketopinic acid-derived catalysts were successfully immobilized onto a polymer support via non-covalent interaction with polymer-bound imidazole. This offers a remarkable advantage because the imidazole also plays the role of an axial ligand in the structure of the catalysts. It is hoped that these heterogeneous catalysts would address some of the limitations that are generally associated with the homogenous catalysts. The application of these

immobilized catalysts in the asymmetric reactions is currently in progress.

Acknowledgements

The financial support of the National Research Foundation and the Tertiary Education Trust Fund, Nigeria for granting fellowship to Yusuf Hassan is gratefully acknowledged.

Conflict of interests

The authors declare that there is no conflict of interests.

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Received: 12.02.2017

Received in revised form: 17.02.2017

Accepted: 24.02.2017