

Immobilized catalysts in synthesis of cyclic carbonates from CO₂

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Abstract – The possibility of obtaining cyclic alkylene carbonates from CO₂ using immobilized catalysts in an one-pot synthesis was investigated. The synthesis proceeded in two stages: oxidation of olefin to epoxide and subsequent reaction of the obtained epoxide with CO₂. The most favorable conditions for conducting of each stage were determined.

Keywords – cyclic carbonates, carbon dioxide, immobilized catalysts

Introduction

Cyclic carbonates have a wide range of applications, among others as plasticisers and solvents, as well as in the synthesis of fine chemicals and polymers [1]. The process of obtaining cyclic alkylene carbonates from CO₂ can be carried out in two main ways: by the reaction of carbon dioxide with epoxides or alkenes [2]. Among the catalysts for these reactions the quaternary onium salts including ionic liquids deserve special attention because they are characterized by high activity.

The direct synthesis method using CO₂ and olefin as substrates is becoming more and more popular. It runs in two stages (Fig. 1). The first one involves the epoxidation reaction of terminal olefins and in the second one *in situ* formed epoxide reacts with CO₂. The advantage of this method is that both process steps can take place in one reaction vessel, so there is no need to isolate and purify the epoxide formed in the reaction. In addition relatively cheap and readily available reagents: alkenes and CO₂, are used in this process. Currently, the key challenge for this technology is the development of new catalysts with high activity and selectivity that would allow the process to run under mild conditions. A particularly interesting aspect is the search for catalysts that can be easily separated after the reaction. This will allow to simplify usually cost-intensive separation operations of the post-reaction mixture.

Results

The paper presents studies on the synthesis of cyclic alkylene carbonates from CO₂ and olefins using immobilized ionic liquids on insoluble carriers as catalysts. The tert-butyl hydroperoxide was used as an oxidant. The influence of selected reaction parameters: temperature, CO₂ pressure, amount of catalyst, molar ratio of reagents on the course of each stage of the synthesis of cyclic carbonates was determined.

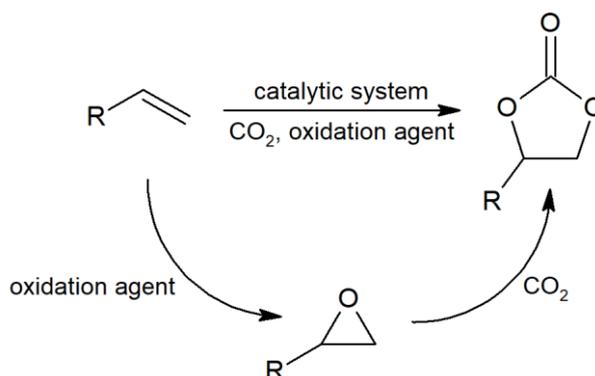


Fig.1. Scheme of synthesis of cyclic carbonates.

The effect of types of inorganic carrier such as silica, as well as the addition cocatalysts, e.g. metal halides, on the activity of catalysts in the synthesis of cyclic carbonates was investigated. It is worth noting that the process did not need a solvent addition.

Conclusion

In this paper the catalysts based on ionic liquids immobilized on insoluble supports for the direct synthesis of cyclic carbonates was investigated. The use of such catalysts allowed easy separation of the post-reaction mixture, easy catalyst separation, e.g. by filtration, and enabled repeated use of the catalyst in the next processes.

Acknowledgments

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References

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