

Green epoxidation of 1,7-octadiene with polymer-supported Mo(VI) catalyst via response surface methodology

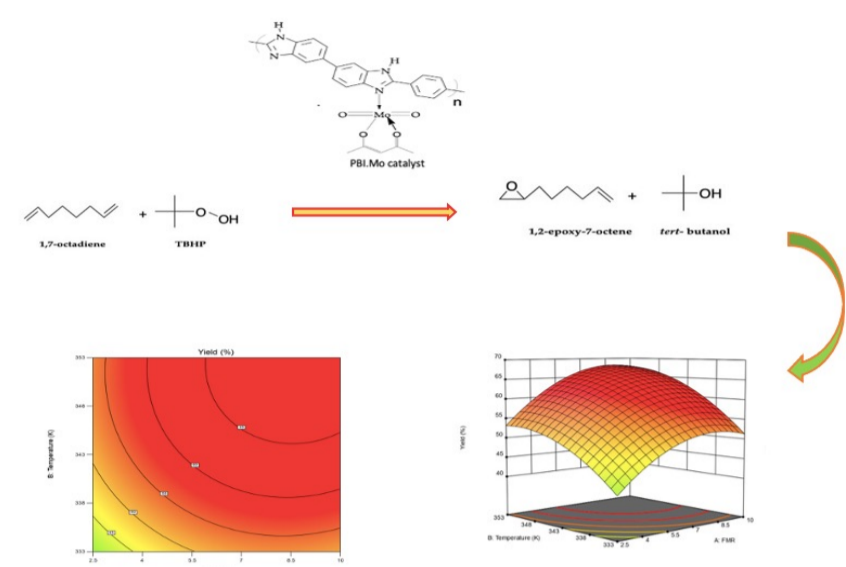
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Motivation



Greener and efficient process

An eco-friendly and efficient technique that can increase production by cutting waste and operating cost.



Important raw materials

Epoxides can be transformed into plasticisers, perfumes, food additives pharmaceutical drugs etc.



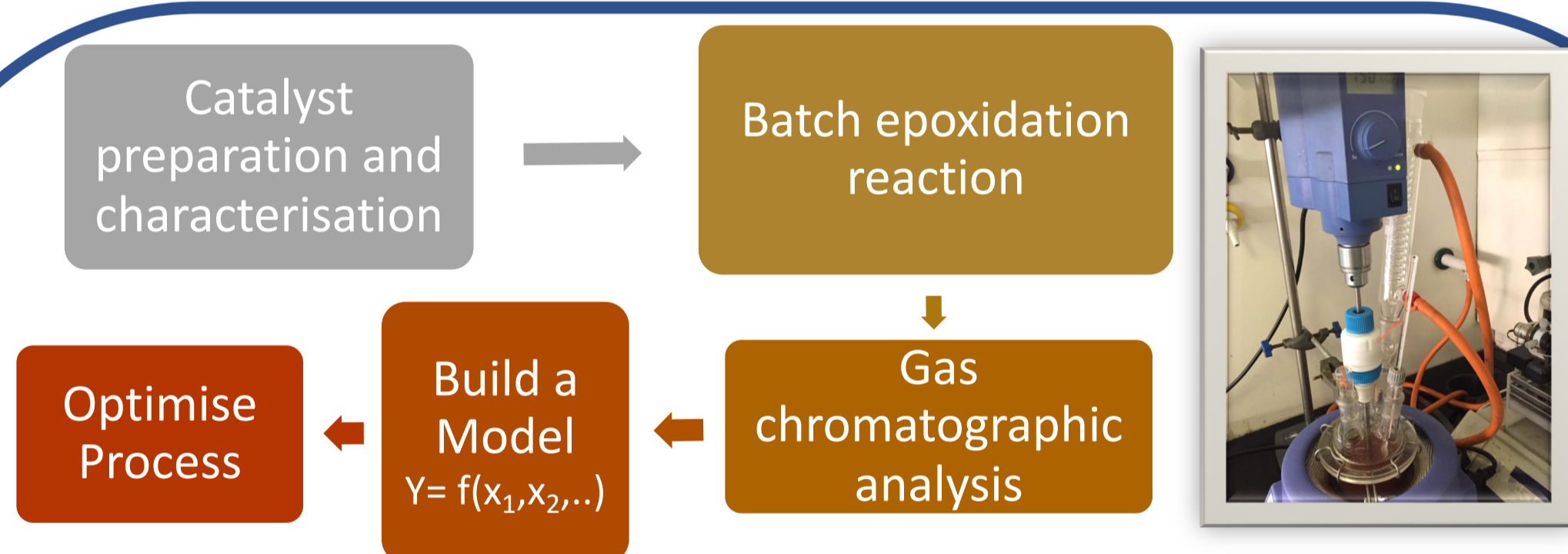
Biochemical and physiological effects

1,2-epoxy-7-octene can act as an antioxidant and found to have anti-cancer properties.

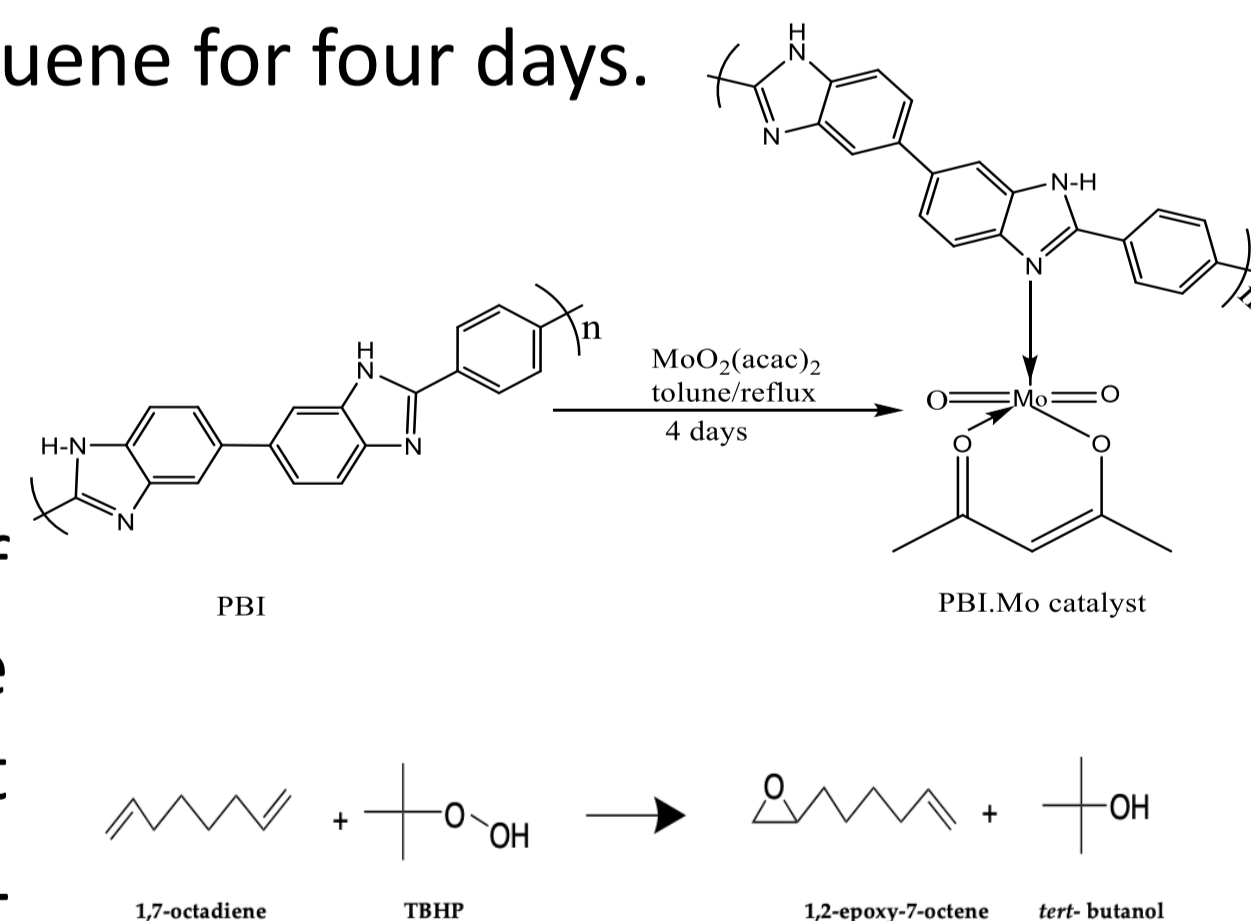
Aim

To synthesise 1,2-epoxy-7-octene using polymer-supported Mo catalyst and evaluate the catalytic efficiency in batch epoxidation reaction via response surface methodology.

Methodology



PBI resin was reacted with $\text{MoO}_2(\text{acac})_2$ in the stoichiometric ratio of 2:1 in toluene for four days.



Batch epoxidation of alkenes with TBHP in the presence of PBI.Mo catalyst was conducted in a 0.25 L jacketed four neck glass reactor.

Applied design of experiments technique to optimise reaction parameters.

Results

Catalyst preparation and characterisation

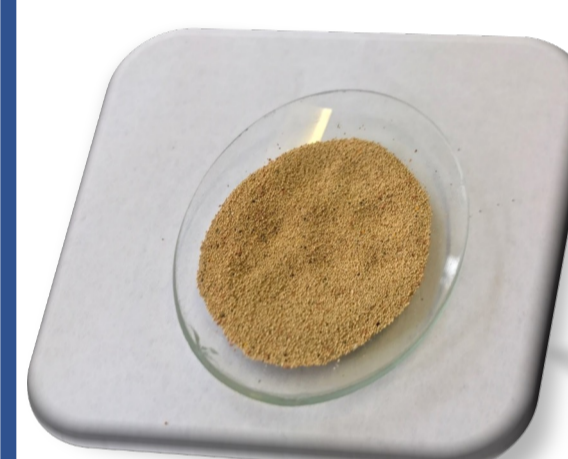
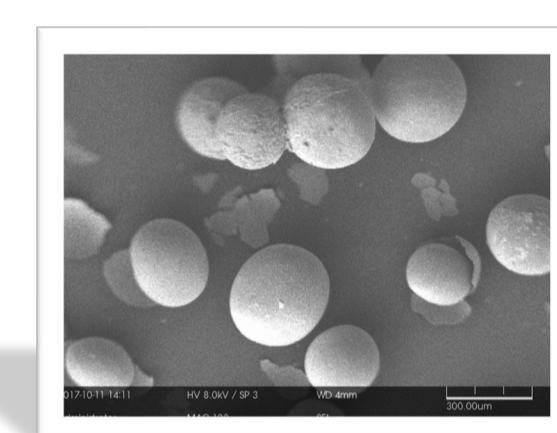
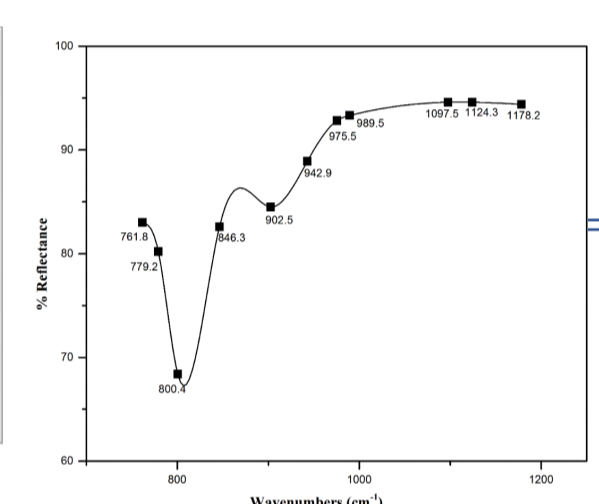


Image of PBI.Mo catalyst

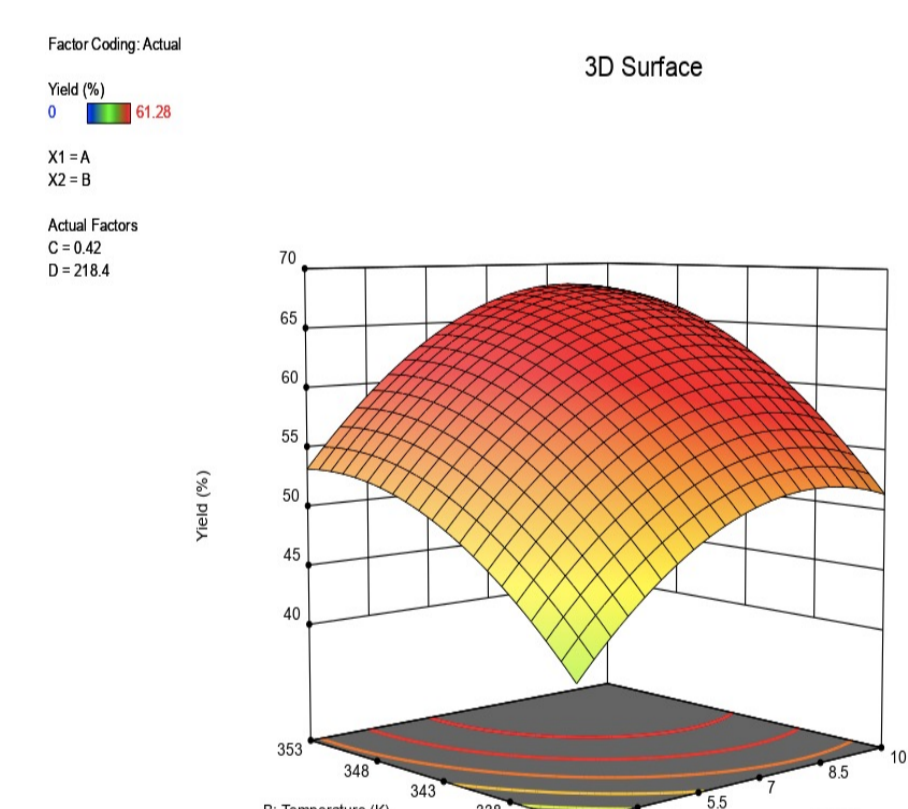


SEM image of PBI.Mo catalyst



FTIR spectra of PBI.Mo catalyst

Optimisation study



3-D graph showing the effect of feed molar ratio and temperature on epoxide yield

The numerical optimisation technique concluded that the maximum yield that can be reached is 66.22% at a feed molar ratio of 7.97:1, reaction temperature 347 K, 0.417 mol% catalyst loading, and reaction time of 218 min.

Conclusions

- PBI.Mo complex could be used as an effective catalyst for a greener and more efficient epoxidation of 1,7-octadiene with TBHP as an oxidising agent.
- Characterisation of PBI.Mo catalyst confirms the presence of Mo(VI) metal centre in the polymer resin.
- The optimisation result has been validated experimentally resulting in an epoxide yield of 64.97% with a relative error of 1.92%.

References

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- M. L. Mohammed, D. Patel, R. Mbeleck, D. Niyogi, D. C. Sherrington and B. Saha, *Appl. Catal., A*, 2013, 466, 142–152.
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