

International Journal Of

Recent Scientific Research

ISSN: 0976-3031 Volume: 7(6) June -2016

BETTER ADSORBING CAPACITY OF CPM-5 IN THE LIGHT OF ITS EQUILIBRIUM KINETICS

Prabin Kumar Sinha and Ajay Kumar Gupta



THE OFFICIAL PUBLICATION OF INTERNATIONAL JOURNAL OF RECENT SCIENTIFIC RESEARCH (IJRSR) http://www.recentscientific.com/ recentscientific@gmail.com



Available Online at http://www.recentscientific.com

International Journal of Recent Scientific Research Vol. 7, Issue, 6, pp. 11942-11944, June, 2016 International Journal of Recent Scientific Research

Research Article

BETTER ADSORBING CAPACITY OF CPM-5 IN THE LIGHT OF ITS EQUILIBRIUM KINETICS

Prabin Kumar Sinha^{1*} and Ajay Kumar Gupta²

^{1,2}University Department of Chemistry, B.R.A Bihar University, Muzaffarpur, Bihar (842001) India

ARTICLE INFO	ABSTRACT
Article History: Received 05 th March, 2016 Received in revised form 08 th April, 2016 Accepted 10 th May, 2016 Published online 28 st June, 2016	The adsorption equilibrium kinetics is one of the important factor in evaluating the stability of the adsorbent for the gas adsorption application, because it controls the time of a fixed bed adsorption system and has an impact on the amount of adsorbent required. The adsorption equilibrium kinetics of CO_2 on CPM-5 was measured at two different temperatures of 298 K and 318 K using Freundlich adsorption isotherm. CPM-5, a MOF is highly porous having high specific surface area, high thermal and chemical stabilities &low densities was found to be better adsorbent for the adsorption of CO_2 gas. The amount of CO2 adsorbed was calculated by weighing CPM-5 before and after experiment.

Copyright © **Prabin Kumar Sinha and Ajay Kumar Gupta., 2016**, this is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution and reproduction in any medium, provided the original work is properly cited.

INTRODUCTION

The release of harmful greenhouse gases (such as CO_2 , CH_4 and N_2O) in to the environment is a growing concern for the world climate change. About 60% of global warming attributed to CO_2 emissions.¹ Fossil fuel power plants are the largest potential source of CO_2 emission. About three-quarters of the increase in atmospheric carbon dioxide is attributed to burning fossil fuels. The current levels of CO_2 concentration in the atmosphere have increased by more than 35% since the industrial revolution.²

In this regards, scientists are trying to develop effective systems for CO_2 removal from fuel gas by combining the high capacity and selectivity, fast kinetics, mild conditions for regeneration, and tolerance to moisture with minimal cost.

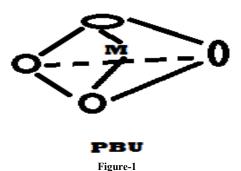
Adsorption in porous adsorbents is considered as an alternative viable approach for CO_2 capturing. Adsorption into solid porous adsorbents is an attractive technology to improve or substitute the current CO_2 adsorption technologies due to their high CO_2 adsorption capacities, simple and easy to control process, low energy consumption, and superior energy efficiency.

Many adsorbents have been developed and studied for CO_2 capturing such as zeolites, activated carbons, modified mesoporous silica; however the common drawbacks of these conventional adsorbents are: high energy consumption for regeneration, low productivity and low CO_2 capacities. New porous materials Metal Organic Frameworks with higher

adsorption capacity and selectivity are needed to improve the CO_2 separation and storage process.

Metal-Organic Frameworks

Metal organic frameworks (MOFs) have emerged as a new class of crystalline porous materials. Metal organic frameworks are crystalline solids consisting of metal ions (normally called nodes) linked by organic ligands (as linkers).³These materials are rigid organic linkers to form one dimensional, two dimensional and three dimensional networks, which is porous. Metal ions generally used for synthesis of MOFs are Zn^{2+} , Ca^{2+} , Cu^+ , Cu^{2+} , $Al^{3+} In^{3+}$ etc. and organic linkers are Phthalic acid, Isophthalic acid, Terephthalic acid, Trimesic acid, Benzene tri benzoate (BTB) etc.



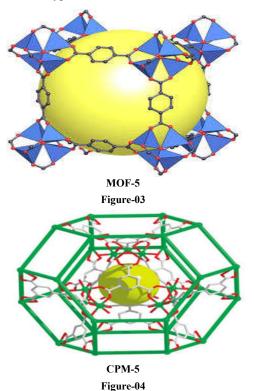
On co-ordination of carboxylates group to a metal centre may results in many different secondary building units (SBUs)⁴ to form one dimensional, two dimensional and three dimensional

University Department of Chemistry, B.R.A Bihar University, Muzaffarpur, Bihar (842001) India

networks. The secondary building units (SBUs) are designed by the four MO_4 primary building units (PBUs). The octahedral geometry of the $[M_4O(O_{12})]$ is designed by the four MO_4 tetrahedral shairing.⁵

rection of the set of

These SBUs connected with an aromatic ring (organic linkers) to design different types of MOFs.⁶



MOFs are renowned materials having remarkable high specific surface area^{7,8}, highly divers' structural chemistry and controlled pore size and shape⁹. The metal organic framework that is studied in the present work is CPM-5, which is a highly porous indium based metal organic framework with a surface area of 2187 m²/g. CPM-5 consists of In₃O clusters as metal centre connected by 1, 3, 5 benzenetricaboxylic acid (Trimesic acid) (H₃BTC) as a linear organic linker. In addition, CPM-5 has unique cage-based porous structures which contribute to a high CO₂ uptake capacity.

Experimental

The adsorption equilibrium kinetics is one of the important

factor in evaluating the stability of the adsorbent for the gas adsorption application, because it controls the time of a fixed bed adsorption system and has an impact on the amount of adsorbent required.

From the Freundlich adsorption isotherm-

$$x/m = K C_i^{1/n}$$

where,

x/m is the adsorbed amount (mol/g), 'Ci' is the equilibrium concentration (mol/dm³)

The constant 'K' and 'n' were estimated from the experimental data of CO_2 adsorption isotherm using the intercept and slope of a Linear Freundlich plot of ln (x/m) VslnCi.

I able -

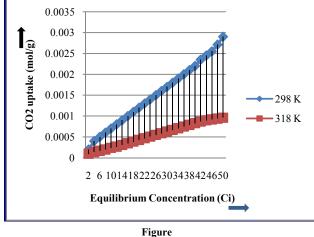
Equilibrium	CO2 uptake	Temperature	
concentration (Ci)	(mol/g)	(K)	
2	0.0002	298	
4	0.0004	298	
6	0.0005	298	
8	0.0006	298	
10	0.0007	298	
12	0.0008	298	
14	0.0009	298	
16	0.0010	298	
18	0.0011	298	
20	0.0012	298	
22	0.0013	298	
24	0.0014	298	
26	0.0015	298	
28	0.0016	298	
30	0.0017	298	
32	0.0018	298	
34	0.0019	298	
36	0.0020	298	
38	0.0021	298	
40	0.0022	298	
42	0.0023	298	
44	0.0024	298	
46	0.0025	298	
48	0.0027	298	
50	0.0029	298	

Table-2

Equilibrium	CO2 uptake	Temperature	
concentration (Ci)	(mol/g)	(K)	
2	0.00010	318	
4	0.00014	318	
6	0.00017	318	
8	0.00020	318	
10	0.00024	318	
12	0.00027	318	
14	0.00031	318	
16	0.00035	318	
18	0.00039	318	
20	0.00043	318	
22	0.00047	318	
24	0.00051	318	
26	0.00055	318	
28	0.00059	318	
30	0.00063	318	
32	0.00067	318	
34	0.00071	318	
36	0.00075	318	
38	0.00079	318	
40	0.00083	318	
42	0.00087	318	
44	0.00090	318	
46	0.00092	318	
48	0.00094	318	
50	0.00096	318	

The adsorption equilibrium kinetics of CO₂ in CPM-5 was measured at two different temperatures of 298 K and 318 K. the amount of CO₂ adsorbed was calculated by weighing CPM-5 before and after experiment.

K is theisothermal equilibrium constant and 'n' are constant for a given adsorbent and adsorbate at a particular temperature.



RESULTS AND DISCUSSION

The CO₂ adsorption equilibrium at temperatures 298 K and 318 K are plotted in the above figure. It was observed that the maximum amount of CO2 adsorbed is 0.0029 mol/g and 0.00096 mol/g at 298 K and 318 K respectively^{10,11}

Finally, CPM-5 showed unique adsorption properties. Therefore, it can be considered as an attractive adsorbent for the separation of CO_2 from fuel gases.

Acknowledgement

The Authors are grateful to Department of Science & Technology (DST), New Delhi to provide financial support and Head, University dept. of Chemistry, B. R. A. Bihar University, Muzaffarpur for providing Lab facilities.

References

- Kiehl, J. T. and K.E. Trenberth, "Earth's Annual 1. Global Mean Energy Budget" Bulletin of the American Metrological Society. 78 (2002)197-208.
- EPA (2008). "Recent Climate Change: Atmosphere 2. Changes". Climate Change Science Program. Nature. 427(2009)138-145.
- 3. Czaja, Alexander U., Trukhan, Natalia, Muller "Industrial applications of Metal Organic Frameworks". Chem. Soc. Rev. 38(5): (2009)1284-1293.
- 4. M. Eddaudi, J. Kim, J.B Wachter, H K Chae, M O'Keeffe and O M Yaghi, J. Am. Chem. Soc.123 (2001)4368-69.
- J L C Rowsell, O M Yaghi, J. Am. Chem. soc. 5. 44(2005)4670-79.
- 6. Yaghi O. M, O'Keeffe, M. Ockwig, N. W. Chae, H. K, M. Kim. J. nature. 423 (2003), 705-708.
- 7. Chae, H.K., Siberio-Perez, D.Y., Kim, J.Y., Eddaudi, M. Matzger et al. nature. 427(2004), 523-527.
- Barrer, D. W. Zeolites molecular Seives(Krieger-8. 1984).
- 9. U. Muller, M. Schubert, F. Teich, H.Puetter, J. Mater.chem. 16(2006) 626-636.
- 10. Zhao Z, Li Z, Lin Y, Adsorption and diffusion of carbon dioxide on metalorganic framework (MOF-5). Industrial Engineering Reviews. 48(2009)10015-10020.
- Saha D, Bao Z, Jia F and Deng S, Adsorption of CO₂, 11. CH₄, N₂O, and N₂ on MOF-5, MOF-177, and zeolite 5A, Environmental Science and Technology.44 (2010)1820-1826.

How to cite this article:

Prabin Kumar Sinha and Ajay Kumar Gupta.2016, Better Adsorbing Capacity of Cpm-5 In The Light of Its Equilibrium Kinetics. Int J Recent Sci Res. 7(6), pp. 11942-11944.

