

Morphological based Mechanism for Adsorption of Hg from Natural Gas using Solid Supported Ionic Liquids

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Results

BET surface area, micropore area, volume and pore size

Adsorbent	BET surface area (m²/g)	t-Plot Micro pore Area (m²/g)	t-Plot micro pore volume (cm³/g)	Pore Size (nm) (average pore diameter)
Fresh Carbon	640.93	549.25	0.22	7.25
[Bmim]CI + KMnO4 coated Carbon	448.85	226.93	0.10	6.37



Introduction

- Fossil Fuels are one of the main anthropogenic source of mercury (Hg) emission
- Hg in natural gas presents predominantly as elemental mercury (Hg⁰)
- Hg in gas plant affects downstream processes and deposits in equipment that poses a health and safety risk for workers involved in maintenance or inspection activities
- To protect the equipment and environment, Hg needs to be removed efficiently.
- Carbon Supported Ionic Liquids have better ability to remove Hg from gas stream than other technologies i-e sulfur or halide impregnated carbon etc.



BET isotherm for [Bmim]CI + KMnO4 coated carbon

Surface morphology of fresh [Bmim]Cl + KMnO4 coated carbon at 5000X







Surface morphology of spent adsorbent ([Bmim]Cl + KMnO4 coated carbon) at 5000X

EDX mapping spent adsorbent ([Bmim]CI + KMnO4 coated carbon)

Elemental analysis of spent adsorbent ([Bmim]CI + KMNO4 coated carbon)





EDX spectra of spent adsorbent ([Bmim]Cl + KMnO4 coated carbon)



Threating effect of Hg in Oil and Gas Industry

Waste incineration · Explosion Hazard Dental amalgam

Anthropogenic mercury sources



Methodology





Synthesis of Carbon Supported Ionic Liquid

Adsorption Experimental Set up

- Surface area, pore size and pore volume of fresh activated carbon and coated (ionic liquid + KMnO4) were evaluated.
- Morphology of adsorbent was studied.
- EDX elemental analysis and mapping were performed for spent adsorbent.

- It was found that the surface area and pore size of adsorbent were reduced due to micropore blockage by the immobilization of ionic liquid.
- The carbon coated with [Bmim]CI & KMnO4 showed promising ability to capture Hg from gas stream. The capacity of Hg removal was found 6.8 mg/g of adsorbent in 72 hours (gas flow rate and Hg⁰ vapor concentration were 60ml/min and 15ppm).
- In the case of spent adsorbent, change in surface morphology was noticed due to the formation of Hg complex (HgCl₂)with [Bmim]Cl.
- EDX elemental analysis and mapping confirmed the presence of Hg on the surface of adsorbent which may be due to chemisorption between Hg and [Bmim]Cl.

References

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