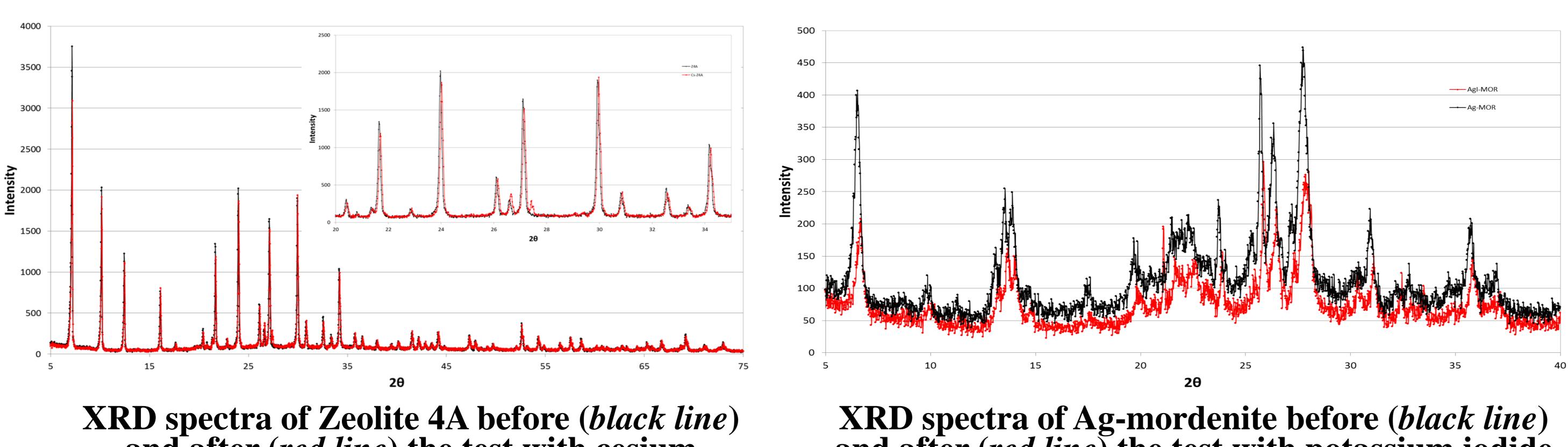
## **Capture of volatile fission products:** present status and future perspectives

G. De Angelis<sup>1</sup>, C. Fedeli<sup>1</sup>, M. Capone<sup>1</sup>, E. Macerata<sup>2</sup>, E. Mossini<sup>2</sup>, M. Mariani<sup>2</sup>,

<sup>1</sup>Unità Tecnica Fissione, Tecnologie e Presidio Nucleare, ENEA Casaccia, Via Anguillarese 301, 00123 S. Maria di Galeria, Roma, Italy

<sup>2</sup>Dipartimento di Energia, Sezione Ingegneria Nucleare, Politecnico di Milano, Via La Masa 34, 20156 Milano, Italy

The performance of some adsorbent materials for retention of volatile fission products has been assessed. Two different zeolites (Zeolite 4A and silver-coated mordenite) have been chosen owing to their known capacity to entrap cesium and iodine, respectively. To this purpose experiments have been made with the pilot plant OGATA (Off-GAs Treatment Apparatus), installed at ENEA laboratories. The tests have been carried out by heating the volatile species at 900°C and passing them through zeolite pellets with the help of argon as carrier gas. The products arising from the interaction with the zeolites have been analyzed mainly by X-rays diffractometry and scanning electron microscopy. The general conclusion from these experiments is that the retention of volatile elements, like cesium and iodine, occurs through a simple adsorption mechanism, which confirms the need to improve the trapping capacity of these materials, especially in view of the final disposal. Moreover, a metal-organic framework (MOF-5), as representative of the new class of materials proposed as adsorbents for volatile radionuclides has been taken into account. Financial support from Italian Ministry for Economic Development (Accordo di Programma: Piano Annuale di Realizzazione 2013) is gratefully acknowledged.

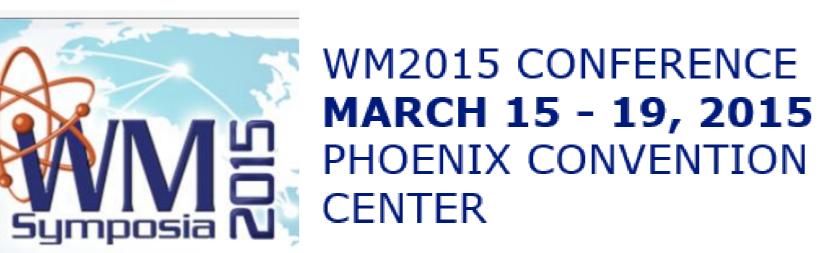


CENTER

and after (*red line*) the test with cesium





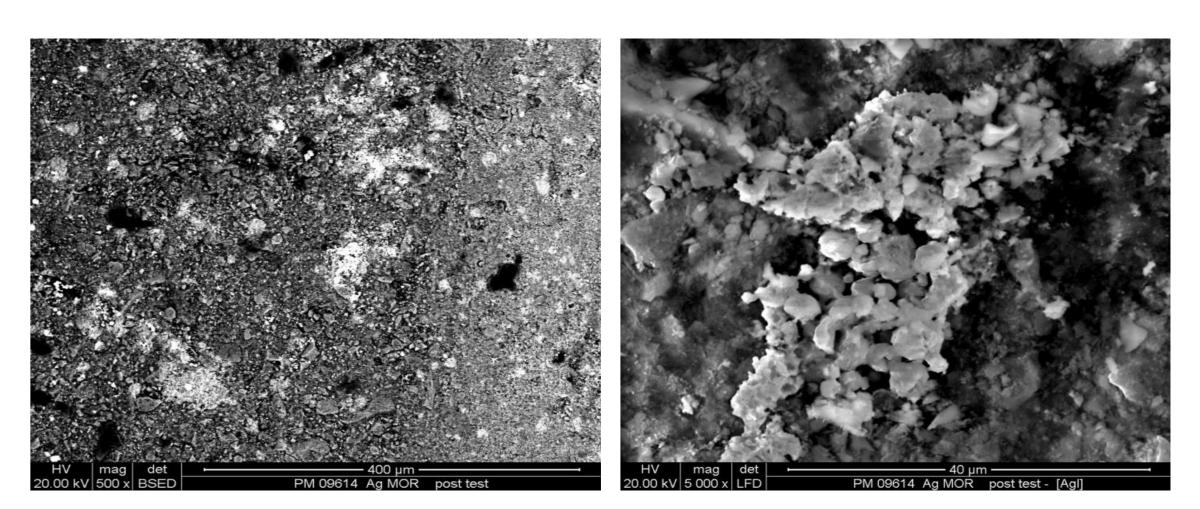


energy and sustainable economic development



and after (*red line*) the test with potassium iodide





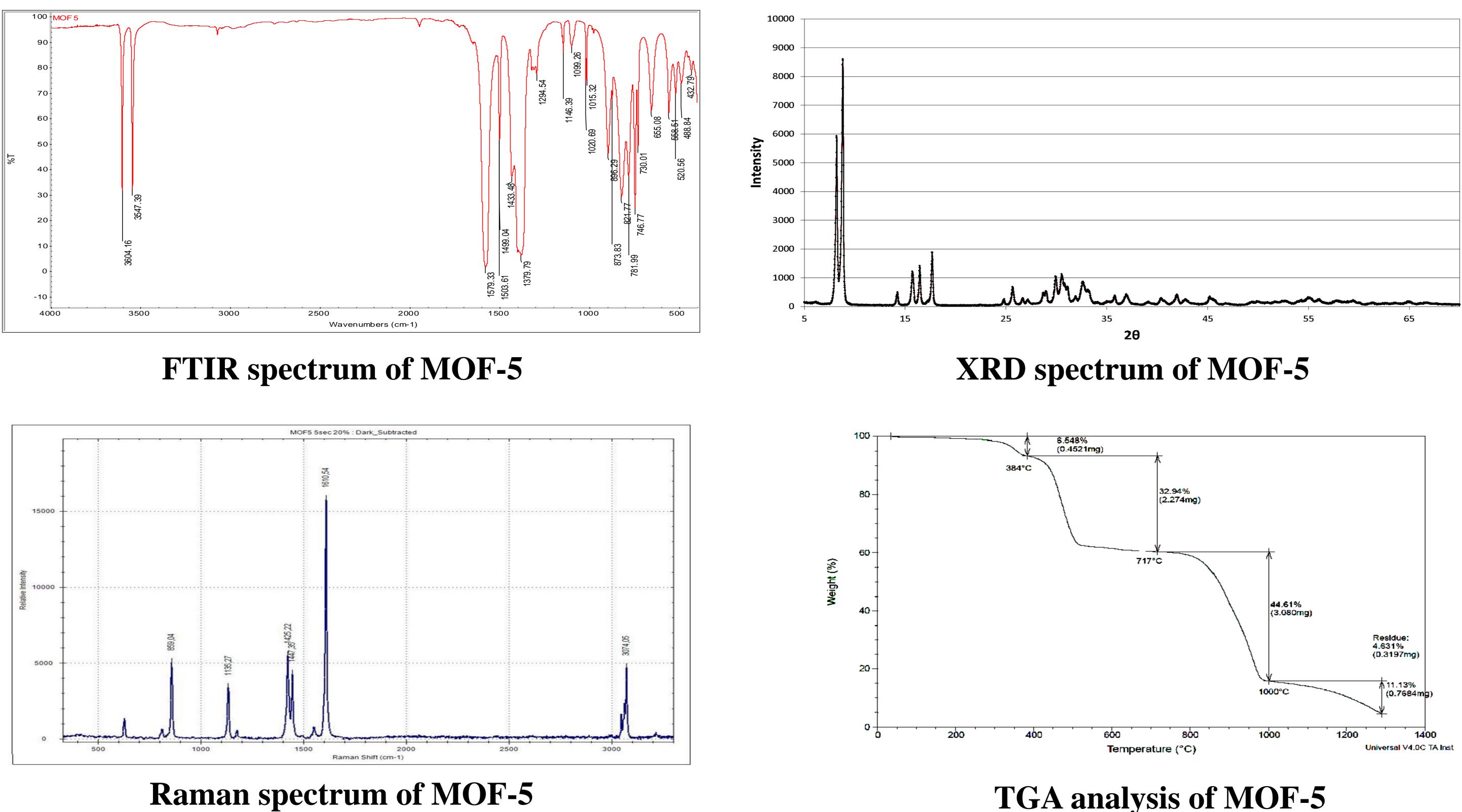
**Micrographs showing the formation of silver** iodide onto the surface of silver-mordenite

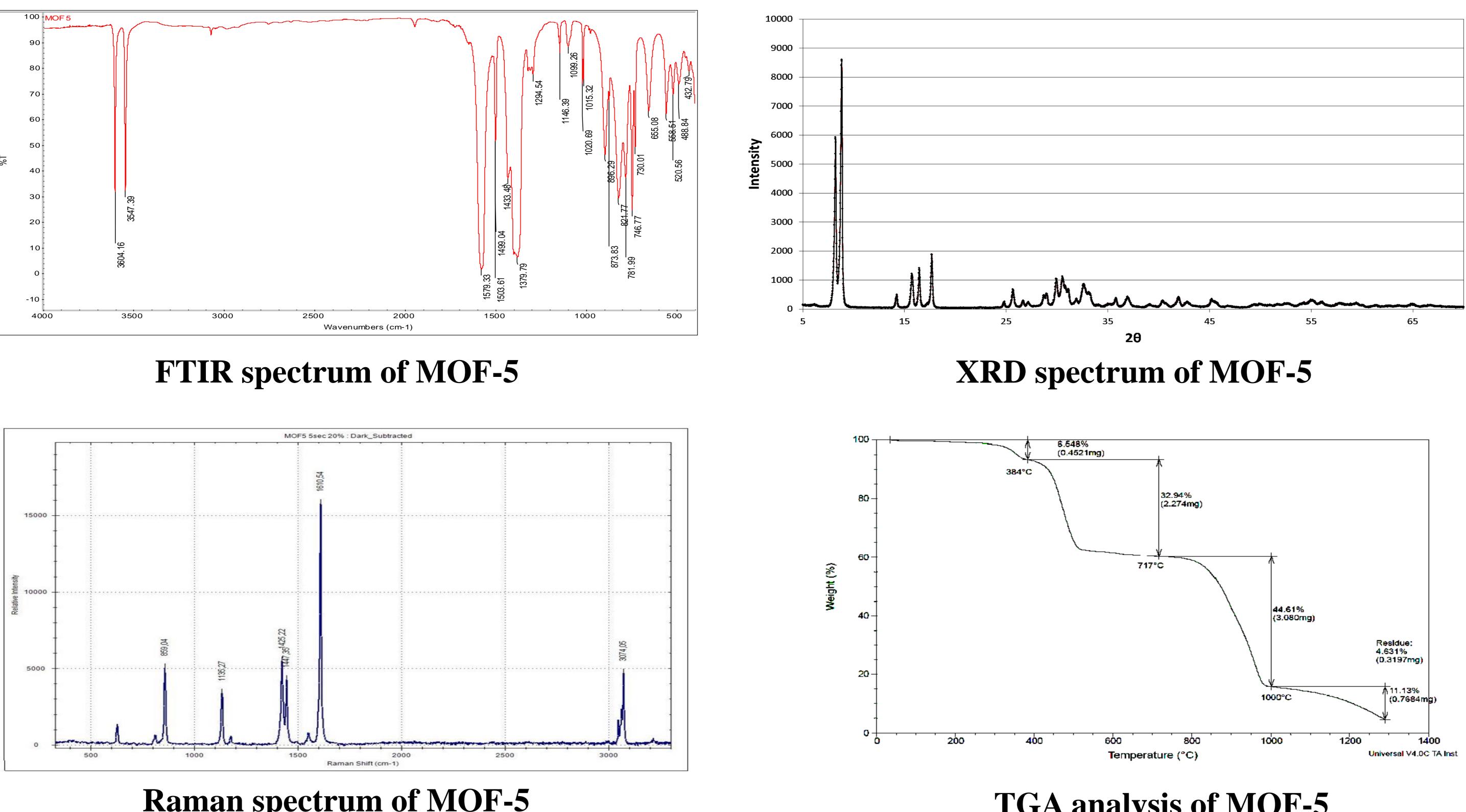
**General view of OGATA pilot plant** 

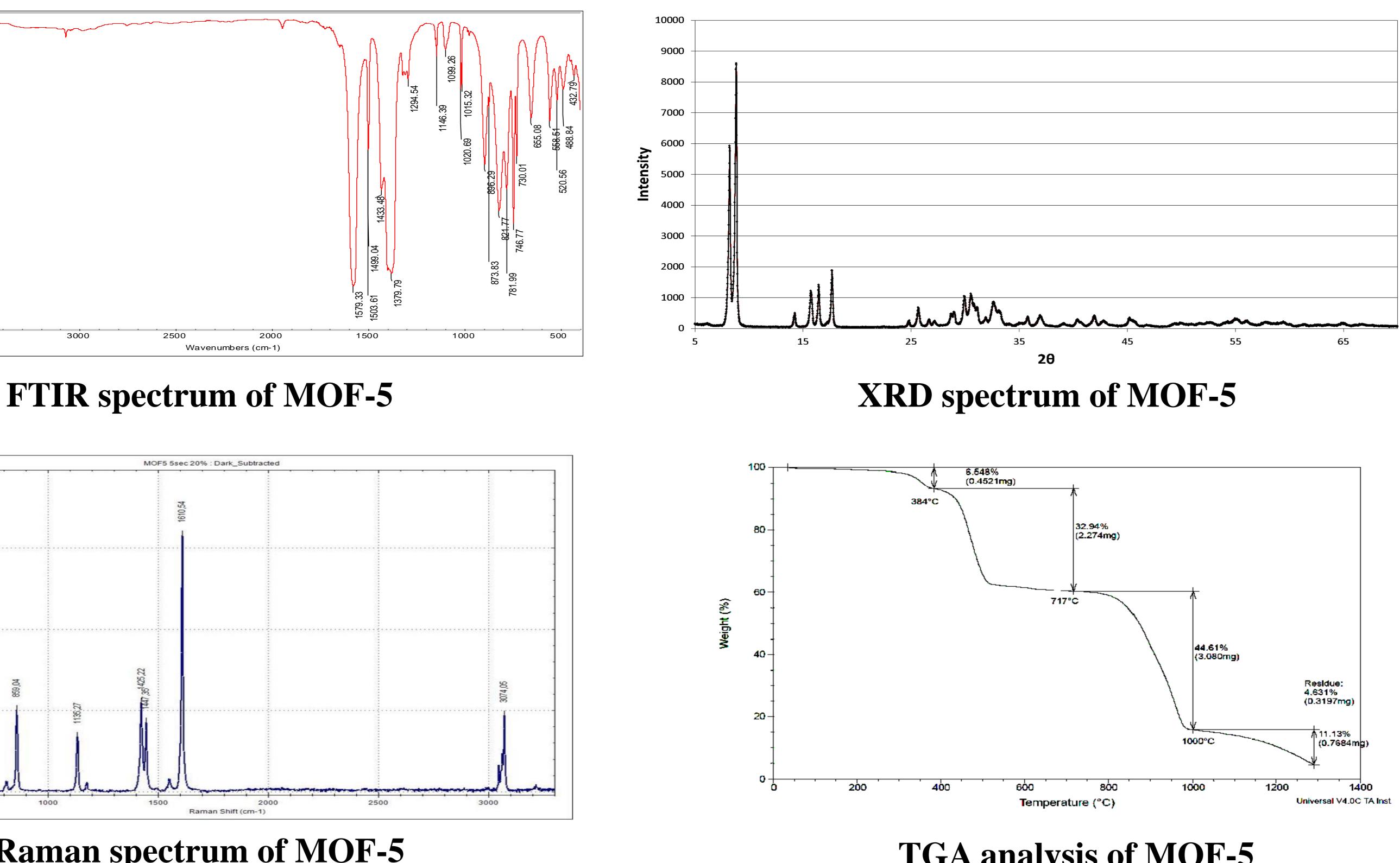
# **Pictures of Ag-mordenite pellets before** (*left*) and after (*right*) the test with KI

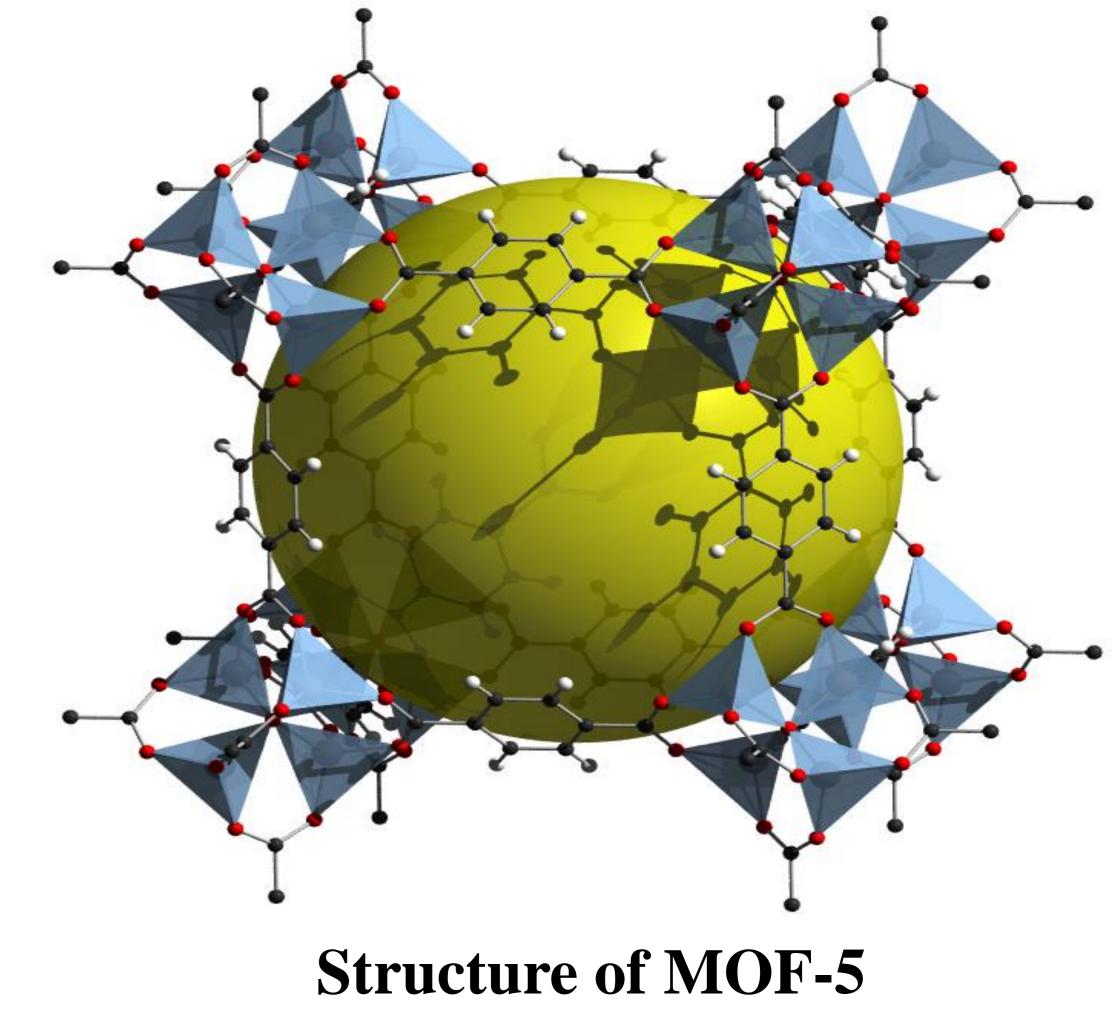
## **Synthesis of MOF-5**

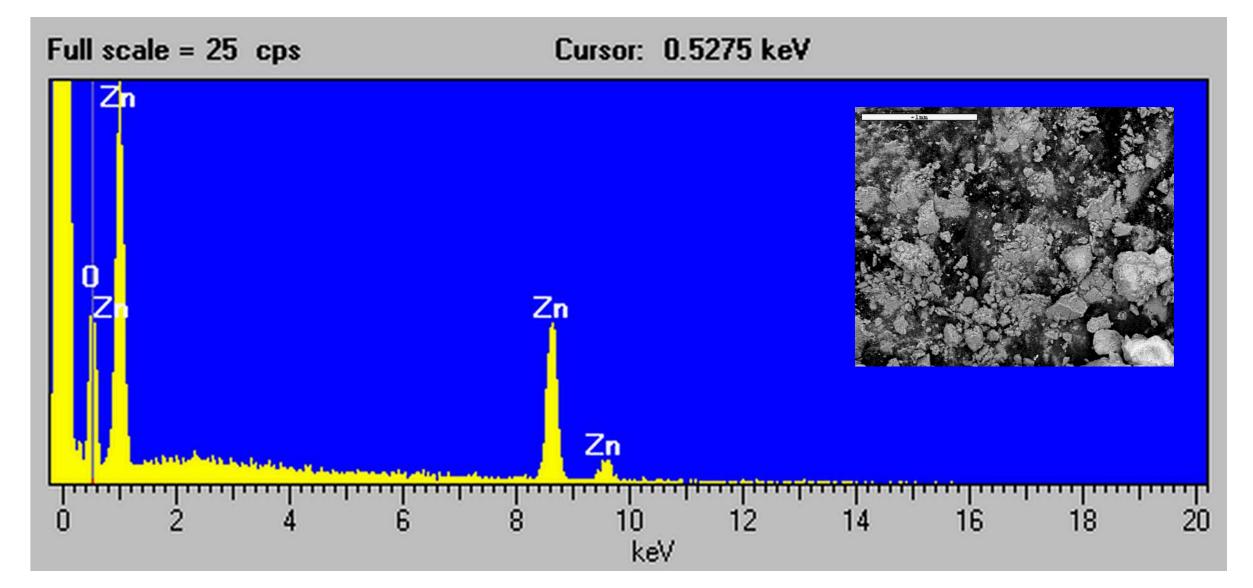
The metal-organic frameworks (MOFs) are a sub-family of the nanoporous crystals, characterized by metal-oxygen cages (vertices) connected by organic bridges. MOFs currently attract intensive interest for their excellent potential for storing and separating gases. Moreover, by changing the organic bridge and/or its functionalization, new MOFs can be designed and synthesized without changing the underlying topology. MOF-5 consists of  $ZnO_4$  inorganic mojety, that acts as secondary building unit, coordinating to benzene 1,4-dicarboxylate, a bidentate ligand that acts as spacers, to form a three dimensional structure. It can be synthesized as follows: about 1.2 g of zinc nitrate hexahydrate and 0.334 g of benzene dicarboxylic acid (BDC) are dissolved in a solution containing 40 mL of dimethyl formamide (DMF) and 5 mL of chlorobenzene under constant stirring. Then 2.2 mL of triethyl amine (TEA) solution and 3–4 drops of Hydrogen Peroxide are added to the mixture. After agitation for 30–45 min a white precipitate is filtered off, washed with DMF and immersed in chloroform for 24 h to exchange DMF solution. Finally, it is dried in vacuum oven at 120 °C for 2 h and stored in vacuum desiccator. All the procedures are performed in a venting hood at ambient temperature (25 °C).















#### **Synthesis steps of MOF-5**



### **SEM-EDS analysis of MOF-5**