

# Uio-68-5-FU as a drug delivery platform for 5-FU

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## Abstract

The drug delivery application of Porous metal–organic frameworks (MOFs) have been investigated due to their unique structures which are built of inorganic nodes and organic ligands. In present study, Uio-68-5-FU was successfully prepared by applied for delivery of 5-fluorouracil (5-FU). Using variety of analytical methods including FTIR, FESEM, EDS, and the prepared nanostructure was characterized. Results revealed the placement of the drug in zeolite is well done and also the in vitro loading and releasing studies, for 5-FU was evaluated. In addition, Based on the in vitro cytotoxicity results, Uio-68-5Fu was able to increase cytotoxicity compared to that of 5-Fu on HT-29 cancerous cells indicating the highlighted role of this drug delivery system.

**Keywords:** Uio-68, MOF, drug delivery, 5-FU, Cytotoxicity, Uio-68-5-FU

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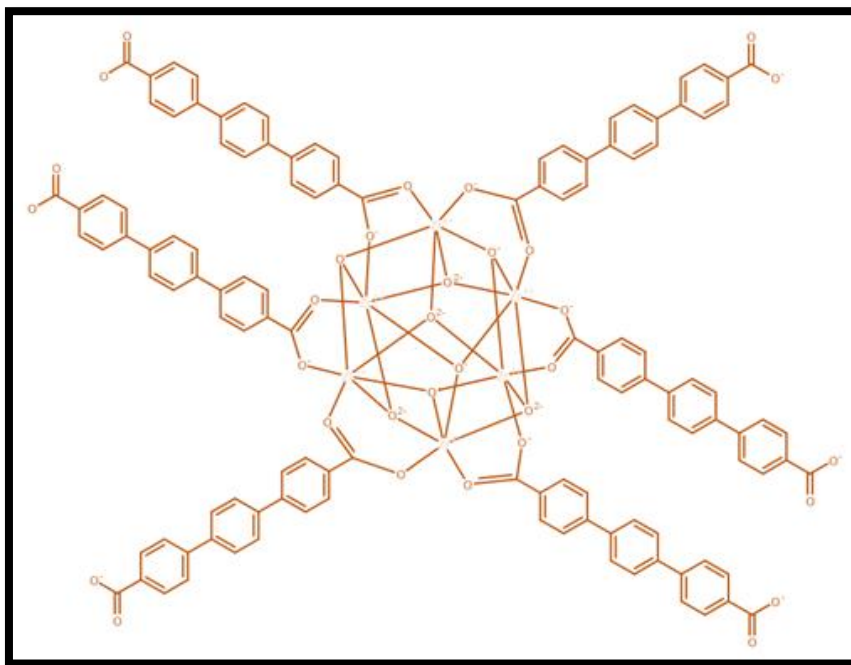
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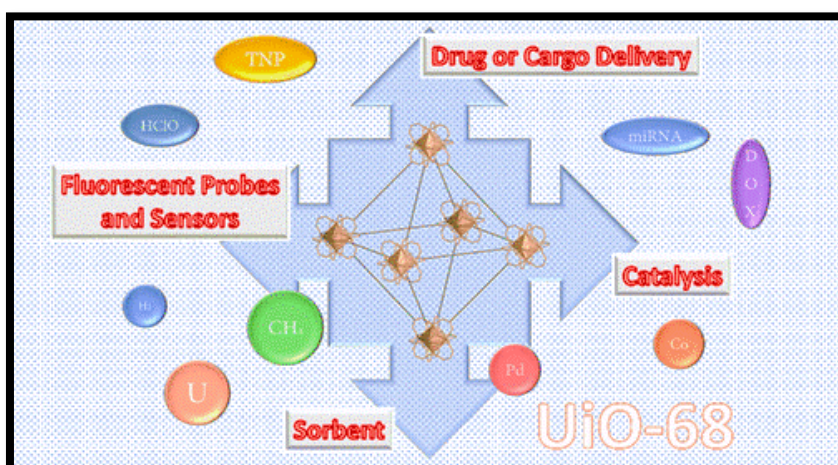
## 1. Introduction

Heavy metal pollution is a global environmental problem, especially in the developing countries, which is caused by increasing industrial and agricultural activities as well as the improper release of metal ions from wastewater and domestic effluents (Boix et al., 2020, Li et al., 2018). These toxic metal ions including Cd (II), As (III and V), Cr (III and VI), Cu (II), Pb (II), and Hg (II) are bioaccumulated readily in humans and animals and believed to be serious threats to the ecological environment and human health (Ma et al., 2019, Zhao et al., 2019). For this purpose, it is much important to effectively remove these toxic pollutants from domestic or industrial wastewater. Many common techniques such as coagulation (Ma et al., 2008), chemical precipitation (Calo et al., 2013), solvent extraction (Cole and Brown, 1959), ion exchange (Crist et al., 1996, Khan et al., 2014), membrane filtration (Fang et al., 2017), electrochemical treatment (Lu et al., 2019) and flotation (Huang et al., 1995) have been used to treat the sewage, while most of them are costly along with having complex operation processes, and these approaches may cause secondary pollution, thus limiting their application (Cai et al., 2017, Ge and Li, 2018). In recent years, adsorption strategies provide a powerful method and have been widely used for the construction of simple and inexpensive platforms to realize high-efficiency removal of heavy metal ions from wastewaters (Sharma and Kumar, 2018, Zhang et al., 2020a, Zhang et al., 2020b, Zhao et al., 2019). In this interesting approach, depending on the strength of the interaction during the adsorption process, it allows an efficient transfer of analytes to the sorbent material. The sorbent is dispersed into the sample solution by magnetic stirring, eddy current or ultrasonic wave, thus increasing the contact area of analyte-sorbent. Therefore, it has become a trend to search for novel materials that can be used not only for adsorption but also for green analysis.



**Fig1. UiO-68 Structure**

In recent decades, metal-organic frameworks (MOFs) as novel organic-inorganic hybrid materials have been an increasing utilization in analytical chemistry. These novel substances are composed of metal ions (clusters or secondary building units (SBUs)) and organic ligands (linkers) through coordination bonds, have attracted wide attention because of their extraordinary properties (high ordered porosity, large surface area, uniform structural cavities, and thermal/chemical stability) (Kumar et al., 2017). Moreover, these properties have emerged in the research of ZIF, MOF, MIL, HKUST, PCN, and UiO, each of which fits the definition of prototype MOFs, implying that various metal and organic ligands can be combined satisfactorily (Winarta et al., 2020). Due to these fascinating properties, the adsorption of target molecules on the surface of MOFs can be effectively promoted. However, most MOFs have relatively lower hydrothermal and chemical stability, and their structures are easy to collapse under high temperatures, high pressure, or acid/alkaline environment. Besides, the structural stability of MOFs decreases as the length of the organic ligand increases, which inhibits the applications of MOFs. Fortunately, the three-dimensional porous nanocomposites constructed from Zr<sup>4+</sup> and carboxylic acids of UiO-MOFs (featuring UiO-66, UiO-67, UiO-68, and UiO-69) have a relatively uniform, large pore size, specific surface area, strong stability as well as active Zr–O clusters, which has been widely studied and used in post-modification and application.



**Fig2. UiO-68 and its applications**

Furthermore, UiO-MOFs have many attractive active binding sites in the organic ligands compared with other MOFs. According to the hard-soft-acid-base (HSAB) theory, some metal ions can be preferred to specific functional ligands (amino, hydroxyl, sulfhydryl, phosphorous, etc.) by ion exchange, electrostatic interaction, or chelation. The specific functional ligands are Lewis bases and have a lone pair of electrons, which can be formed coordination bonds with heavy metal ions (Lewis acids). Therefore, the post-modification of UiO-MOF with specific functional groups to increase the active site for the adsorption of heavy metal ions is a hot topic as one of the promising adsorbents, which is considered as a reliable and advanced solid sorbent in the field of environment.

The purpose of this review was to focus on the potential of UiO series MOFs as the emerging MOFs material for the effective adsorption and separation of heavy metal ions from environmental samples. UiO-MOFs or functionalized UiO-MOFs composites have been widely applied as an efficient and selective adsorbent due to its structural diversity, excellent water stability, good biocompatibility, and other excellent properties (Mukhopadhyay et al., 2019, Yu et al., 2019, Zhao et al., 2019). Moreover, a comprehensive study about UiO-MOFs and their composites including the synthetic methods and the applications in the removal of heavy metal ions were presented in detail. Finally, the adsorption characteristics and mechanism of UiO-MOFs as solid sorbents for heavy metal ions were discussed, including adsorption isotherms equation, adsorption thermodynamics, and kinetics. Scheme 1 was a schematic illustration of the synthesis, characteristics, and application of UiO-MOFs in the environment.

### **Synthetic methods of UiO-MOFs**

The premise for the synthesis of MOFs is experimental property research and practical application. UiO-MOFs are three-dimensional porous materials and constructed by  $Zr^{4+}$  and dicarboxylic acid ligands. Although the ligands of UiO-MOFs have different lengths, UiO-64, UiO-66, UiO-67, UiO-68, UiO-69, and various derivatives have the same reticular structure, the change of the ligand does not affect the thermal stability of UiO-MOFs. Zr(IV)-based MOFs is formed a similar cubic close-packed (CCP) structure with enlarged versions (Cavka et al., 2008, Nguyen et al., 2014), including  $(Zr_6O_4(OH)_4(BDC)_6)$  for UiO-66;  $Zr_6O_4(OH)_4(BPDC)_6$  for UiO-67;  $Zr_6O_4(OH)_4(TPDC)_6$  for UiO-68, and  $Zr_6O_4(OH)_4(2,6-NDC)_6$  for UiO-69 (Xu et al., 2019, Kutzscher et al., 2016). As can be seen,  $Zr_6O_4(OH)_4$  (as SBU) are coordinated with twelve bridge ligands separately to form a three-dimensional structure, which consists of one octahedral center hole cage and eight tetrahedral corner cages. The theoretical values of pore volumes of UiO-MOFs (UiO-66, UiO-67, UiO-68) are 0.45, 1.05, and  $1.82 \text{ cm}^3\text{g}^{-1}$ , and the theoretical values of specific surface area are 1087, 3000, and  $4170 \text{ m}^2\text{g}^{-1}$  (Cavka et al., 2008), respectively. Therefore, due to the strong binding ability of Zr-O bond, the synthesis of UiO-MOFs is gradually concentrated on functionalization or changes in ligand length in order to control the physical/chemical properties, pore size, and morphology of UiO-MOFs. UiO-MOFs and functional UiO-MOFs composites are synthesized based on the node and connector approach and can maintain their topology structures and crystallinity when the organic ligands or the inorganic SBUs are enlarged, replaced, or functionalized. This may depend on the interaction forces such as coordination bond, hydrogen bond,  $\pi-\pi$  interaction, metal bond, and so on. The original pore characteristics of UiO-MOFs are determined by the properties of metal ions and linkers (Qiu and Zhu, 2009). The main structural characterizations can be influenced by the solvent, pH, reaction temperature, and so on. So far, some different synthesis strategies have been reported for UiO-MOFs such as solvent-thermal, microwave, volatilization, diffusion, template synthesis, ultrasonic, and mechanical stirring.

## **2. Results and discussion**

### **Characterization**

The chemical structure of the UiO-68 was characterized with different analytical methods such as XRD, SEM & TEM.

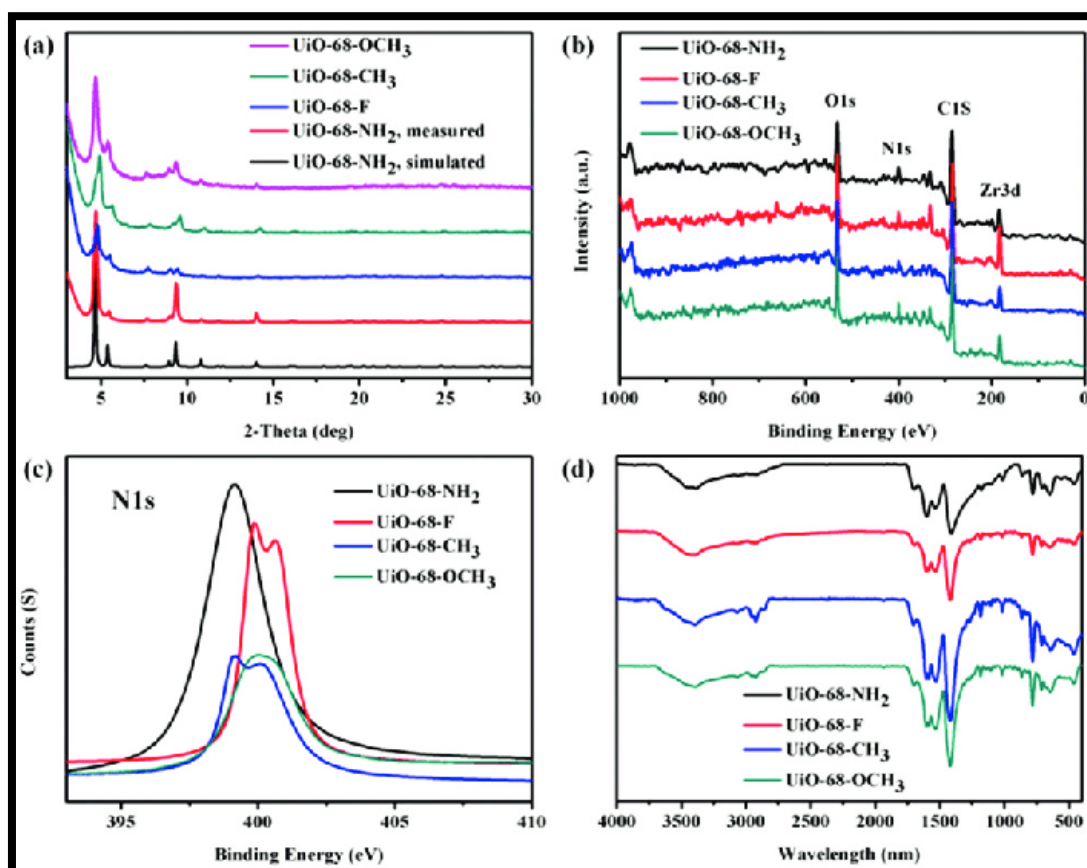


Fig4. (a) PXRD patterns of simulated and measured UiO-68-NH<sub>2</sub>, UiO-68-F, UiO-68-CH<sub>3</sub> and UiO-68-OCH<sub>3</sub>. (b) XPS, (c) XPS of N 1s and (d) IR spectra of UiO-68-NH<sub>2</sub>, UiO-68-F, UiO-68-CH<sub>3</sub> and UiO-68-OCH<sub>3</sub> [36]

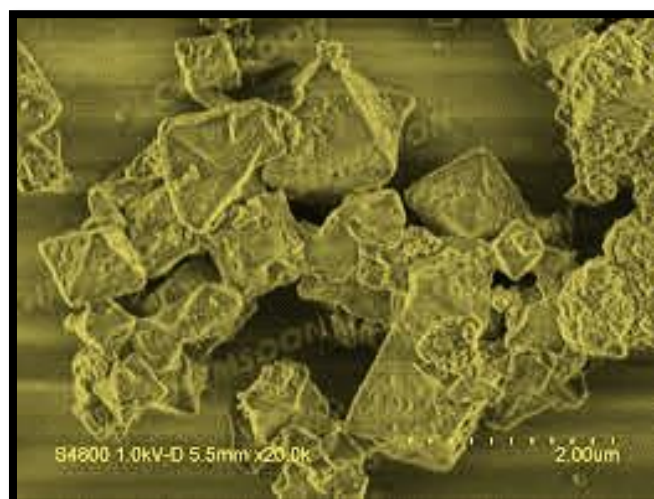
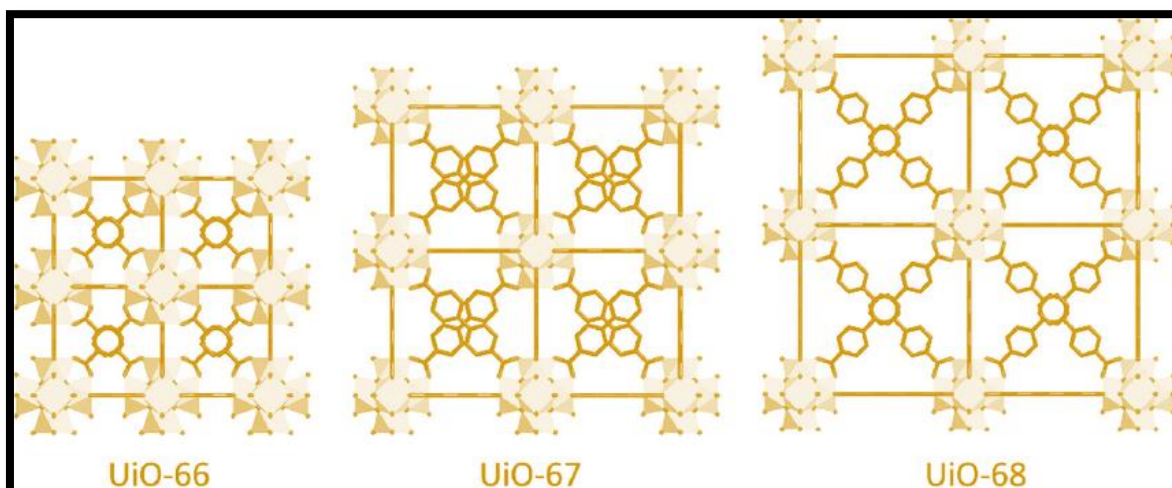


Fig5. TEM of UiO-68



**Fig6. Structures of UiO-66, 67, and 68[37]**

### Conclusions and perspective

In this study, UiO-68 was applied for delivery of 5-Fu. The obtained nanostructure poses spherical morphology with an average diameter of 39-52 nm. Results showed the high loading capacity (90%) and sustained drug release behavior. Moreover, upon exposure by UiO-68-5Fu, higher cytotoxicity than those for UiO-68 and 5-Fu drug against PC3 cells was determined indicating UiO-68-5Fu may could be a promising anticancer drug delivery system in the future.

### Outlook for MOFs

The numerous advantages of MOFs, foremost their high surface area and modular composition, place them at a multidisciplinary crossroads. For good reason, MOFs are one of the most active research fields today, with aspects of their fundamental and applied properties permeating into disciplines as varied as electronics, chemical engineering, and optics. Whereas this Outlook does not attempt to delineate the developments and potential in all these areas, we have introduced some of the exciting prospects related to continued synthetic advances in the field. We further elaborated on three applied areas where MOFs are primed to excel: in challenging gas separations, as porous electrical conductors, and in heterogeneous catalysis. These examples are not exhaustive, but present subtleties that are applicable and relevant to many other applications of MOFs. The challenges and opportunities in these select applications, which span both the traditional and the modern aspects of the field, are illustrative of the continually expanding interest and bright future for MOF chemistry.

### References

- [1]. Tan, Chunxia, et al. "Controlled exchange of achiral linkers with chiral linkers in Zr-based UiO-68 metal-organic framework." *Journal of the American Chemical Society* 140.47 (2018): 16229-16236.
- [2]. Kutzscher, Christel, et al. "Proline functionalized UiO-67 and UiO-68 type metal-organic frameworks showing reversed diastereoselectivity in aldol addition reactions." *Chemistry of Materials* 28.8 (2016): 2573-2580.
- [3]. Li, Yan-An, et al. "UiO-68-ol NMOF-based fluorescent sensor for selective detection of HClO and its application in bioimaging." *Inorganic Chemistry* 56.21 (2017): 13241-13248.
- [4]. Hu, Yu-Hong, et al. "TiO<sub>2</sub>@ UiO-68-CIL: A metal-organic-framework-based bifunctional composite catalyst for a one-pot sequential asymmetric Morita-Baylis-Hillman reaction." *Inorganic Chemistry* 58.8 (2018): 4722-4730.
- [5]. Guo, Fan, et al. "Pt nanoparticles embedded in flowerlike NH<sub>2</sub>-UiO-68 for enhanced photocatalytic carbon dioxide reduction." *Journal of Materials Chemistry A* 7.46 (2019): 26490-26495.
- [6]. O. Zarei, F. Azimian, M. Hamzeh-Mivehroud, J. Shahbazi-Mojarrad, S. Hemmati, S. Dastmalchi. *Medicinal Chemistry Research* 2020, 29, 1438-1448.
- [7]. S. Rojas, T. Devic, P. Horcajada. *Journal of Materials Chemistry B* 2017, 5, 2560-2573.
- [8]. B.-H. Song, X. Ding, Z.-F. Zhang, G.-F. An. *Journal of the Iranian Chemical Society* 2019, 16, 333-340.
- [9]. E. Tawfik, M. Ahamed, A. Almalik, M. Alfaqeeh, A. Alshamsan. *Saudi Pharmaceutical Journal* 2017, 25, 206-213.
- [10]. N.M. Mhaidat, M. Bouklihacene, R.F. Thorne. *Oncology letters* 2014, 8, 699-704.
- [11]. W. Cai, J. Wang, C. Chu, W. Chen, C. Wu, G. Liu. *Advanced Science* 2019, 6, 1801526.
- [12]. Song, Y., Chen, Y., Xu, M., Wei, W., Zhang, Y., Yang, G., Ran, R., Wei, Z., & Zongping, S. (2020). "A Cobalt-Free Multi-Phase Nanocomposite as Near-Ideal Cathode of Intermediate-Temperature Solid Oxide Fuel Cells Developed by Smart Self-Assembly." *Advanced Materials*, 32(8), 1906979. <https://doi.org/10.1002/adma.201906979>
- [13]. Rane, A. V., Kanny, K., Abitha, V.K., & Thomas, S. (2018). "Methods for synthesis of nanoparticles and fabrication of nanocomposites. In S. M. Bhagyaraj, O. S. Oluwafemi & S. Thomas (Eds.), "Synthesis of inorganic nanomaterials (pp. 121-139). Woodhead Publishing. <https://doi.org/10.1016/B978-0-08-101975-7.00005-1>

- [14]. Lopez, N., Janssens, T.V.W., Clausen, B.S., Xu, Y., Mavrikakis, M., Bligaard, T., & Nørskov, J.K. (2004). "On the origin of the catalytic activity of gold nanoparticles for low-temperature CO oxidation". *Journal of Catalysis*, 223(1), 232-235. 10.1016/j.jcat.2004.01.001
- [15]. Hubbe, Martin A., et al. "Cellulosic nanocomposites: a review." *BioResources* 3.3 (2008): 929-980.
- [16]. Kormmann, Xavier, Henrik Lindberg, and Lars A. Berglund. "Synthesis of epoxy-clay nanocomposites: influence of the nature of the clay on structure." *Polymer* 42.4 (2001): 1303-1310.
- [17]. Fornes, T. D., and D. R. Paul. "Modeling properties of nylon 6/clay nanocomposites using composite theories." *polymer* 44.17 (2003): 4993-5013.
- [18]. Wagner, H. D., et al. "Stress-induced fragmentation of multiwall carbon nanotubes in a polymer matrix." *Applied physics letters* 72.2 (1998): 188-190.
- [19]. Gu, Dongdong, et al. "Selective laser melting of TiC/Ti bulk nanocomposites: Influence of nanoscale reinforcement." *Scripta Materialia* 67.2 (2012): 185-188.
- [20]. Fechete, Ioana, Ye Wang, and Jacques C. Védrine. "The past, present and future of heterogeneous catalysis." *Catalysis Today* 189.1 (2012): 2-27.
- [21]. Dąbrowski, A. "Adsorption—from theory to practice." *Advances in colloid and interface science* 93.1-3 (2001): 135-224.
- [22]. Shahabuddin, M., et al. "A review on the production of renewable aviation fuels from the gasification of biomass and residual wastes." *Bioresource Technology* 312 (2020): 123596.
- [23]. Khan, Faisal I., and Aloke Kr Ghoshal. "Removal of volatile organic compounds from polluted air." *Journal of loss prevention in the process industries* 13.6 (2000): 527-545.
- [24]. Rathje, William L., and Cullen Murphy. *Rubbish!: the archaeology of garbage*. University of Arizona Press, 2001.
- [25]. Gorse, Christopher, David Johnston, and Martin Pritchard. *A dictionary of construction, surveying, and civil Engineering*. Oxford University Press, 2012.
- [26]. Malamis, S., and E. Katsou. "A review on zinc and nickel adsorption on natural and modified zeolite, bentonite and vermiculite: examination of process parameters, kinetics and isotherms." *Journal of hazardous materials* 252 (2013): 428-461.
- [27]. VenkataSwaroopDattaDevulapalli, Ryan P. McDonnell, Jonathan P. Ruffley, Priyanka B. Shukla, Tian-Yi Luo, Matheus L. De Souza, Prasenjit Das, Nathaniel L. Rosi, J. Karl Johnson, Eric Borguet. Identifying UiO-67 Metal-Organic Framework Defects and Binding Sites through Ammonia Adsorption. *ChemSusChem* 2022, 15 (1) <https://doi.org/10.1002/cssc.202102217>
- [28]. Amir Hossein Vahabi, Fataneh Norouzi, Esmail Sheibani, Mehdi Rahimi-Nasrabadi. Functionalized Zr-UiO-67 metal-organic frameworks: Structural landscape and application. *Coordination Chemistry Reviews* 2021, 445, 214050. <https://doi.org/10.1016/j.ccr.2021.214050>
- [29]. Mustafa Kõmurcu, Andrea Lazzarini, Gurpreet Kaur, Elisa Borfecchia, Sigurd Øien-Ødegaard, Diego Gianolio, Silvia Bordiga, Karl Petter Lillerud, Unni Olsbye. Co-catalyst free ethene dimerization over Zr-based metal-organic framework (UiO-67) functionalized with Ni and bipyridine. *Catalysis Today* 2021, 369, 193-202. <https://doi.org/10.1016/j.cattod.2020.03.038>
- [30]. Guihao Zhong, Dingxin Liu, Jianyong Zhang. Incorporation of Functional Groups Expands the Applications of UiO-67 for Adsorption, Catalysis and Thiols Detection. *ChemistrySelect* 2018, 3 (25), 7066-7080. <https://doi.org/10.1002/slct.201800840>
- [31]. Yuan, Shuai, et al. "Stable metal-organic frameworks: design, synthesis, and applications." *Advanced Materials* 30.37 (2018): 1704303.
- [32]. Chen, Liyu, et al. "One-step encapsulation of Pd nanoparticles in MOFs via a temperature control program." *Journal of Materials Chemistry A* 3.29 (2015): 15259-15264.
- [33]. Qiu, Lu, et al. "Fluorinated phenylpyridine iridium (III) complex based on metal-organic framework as highly efficient heterogeneous photocatalysts for cross-dehydrogenative coupling reactions." *Journal of Materials Science* 55.22 (2020): 9364-9373.
- [34]. Hong, Yuechao, et al. "Transition metal oxodiperoxo complex modified metal-organic frameworks as catalysts for the selective oxidation of cyclohexane." *Materials* 13.4 (2020): 829.
- [35]. Chen, Rui, et al. "Ruthenium (II) complex incorporated UiO-67 metal-organic framework nanoparticles for enhanced two-photon fluorescence imaging and photodynamic cancer therapy." *ACS Applied Materials & Interfaces* 9.7 (2017): 5699-5708.
- [36]. Wei, Yuan-Ping, et al. "Different functional group modified zirconium frameworks for the photocatalytic reduction of carbon dioxide." *Dalton Transactions* 48.23 (2019): 8221-8226.
- [37]. Yuan, Shuai, et al. "Stable metal-organic frameworks: design, synthesis, and applications." *Advanced Materials* 30.37 (2018): 1704303.