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## Reduction-induced alkali ion capture with polyoxometalate encapsulated metal–organic frameworks

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POM@MOFs offer synergistic properties that combine the advantages of both polyoxometalates (POMs) and metal—organic frameworks (MOFs). When combined, POMs are introduced to provide strong acidity, oxygen-rich sites, and redox activity, while MOFs contribute their high surface area, porous cavities, and tunable rigid structure to counteract the high solubility of POMs. This combination provides a heterogenous material that can be used in applications such as, gas storage, catalysts, and proton conductivity.<sup>1</sup>

In this work, various POM@ZIF-8 composites were synthesized for reduction-induced alkali ion capture. ZIF-8 (Zn[C<sub>4</sub>H<sub>5</sub>N<sub>2</sub>]<sub>2</sub>) was chosen for this study for its crystallinity, porosity, chemical, thermal stability, and simple synthesis.<sup>2</sup> As for the POMs, various Keggin-types, including  $[\alpha$ -PMo<sub>12</sub>O<sub>40</sub>]<sup>3-</sup>,  $[\alpha$ -PW<sub>12</sub>O<sub>40</sub>]<sup>3-</sup>, and  $[\alpha$ -SiMo<sub>12</sub>O<sub>40</sub>]<sup>4-</sup>, were introduced into the ZIF-8 framework to compare the effects of differing reducibility on ion capture properties. The ion capturing potential of the synthesized POM@ZIF-8s were initially evaluated in a CsCl methanol solution. With the addition of ascorbic acid as a reducing agent, the metal building blocks of the POMs are reduced. Subsequently, to compensate for the increase in electrons, alkali ions can be captured into the framework of the POM@ZIF-8. Preliminary results suggest that the innate reducibility of the incorporated POMs have a direct impact on the maximum adsorption capacity of certain alkali ions.



Figure 1. Scheme of reduction-induced alkali ion capture in POM@ZIF-8

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