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# Au nanoparticles confined in self-assembled Zn(II) metal–organic cubane cages for light-driven conversion of furfural to 2-methyl furan in biofuel production†

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A polyoxometalate Zn(II) cubane cage with intrinsic confined spaces constructed by the self-assembly of tetrametallic cubane clusters was obtained and utilized as a skeletal support for the encapsulation of gold nanoparticles. AuNP-encapsulated Zn(II) cubane cages (ZnCC@AuNPs) were characterized using FTIR, TGA, BET, SEM, TEM, XPS, and PXRD analysis. The catalytic activity of ZnCC@AuNPs for the hydrogenation of furfural to 2-methyl furan, which is an active precursor for biofuel production, was investigated. The anchored AuNPs generate Au–H species under visible light and catalyse the reaction in 2-propanol at 60 °C, yielding >99% of the product in just 1.5 h. The reaction kinetics and mechanism were confirmed by *in situ* ATR-IR and DFT studies. The parameter optimization suggested exceptional chemical stability and reusability of ZnCC@AuNPs as a catalyst without any agglomeration even after several cycles of use. These intriguing features are suitable for developing sustainably viable catalysts for the biomass refinery industry.

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## 1. Introduction

The self-assembly of metal–organic cages produces porous frameworks that can serve as molecular containers in supramolecular catalysis.<sup>1</sup> These cages can tune the reactivity of molecules by encapsulating the reacting species inside the cavities or by confining the catalytic sites. In particular, in metal–organic cubane cages, the rigidity and strength provided by the organic pillars make them versatile frameworks that can withstand harsh conditions and provide interactive sites for holding various species.<sup>2</sup> This type of metal–organic cage has been reported in the past for catalysing particular reactions like silyl(bis)amido ligand  $[\{\text{Ph}_2\text{Si}(\text{NAr}^*)_2\text{Zn}(\text{HMDS})\}^- \{\text{Na}(\text{THF})_6\}^+]$  ( $\text{Ar}^* = 2,6\text{-diisopropylphenyl}$ ) for enolate formation,<sup>3</sup> Zn cage  $\{[\text{Zn}_4(\text{TBSC})(\mu_4\text{-OH})]_4\text{-}(5\text{-Me-im-1,3-BDC})^+ \text{Cl}^- \}_8$  ( $\text{DMF}$ )<sub>10</sub>( $\text{CH}_3\text{OH}$ )<sub>15</sub>( $\text{H}_2\text{O}$ )<sub>23</sub> for CO<sub>2</sub> conversion reactions,<sup>4</sup>  $[\{\text{Cu}_{12}(\text{TMBTA})_8(\text{DMA})_4(\text{H}_2\text{O})_8\} \cdot 8\text{H}_2\text{O} \cdot \text{X}]$  (where TMBTA is (2,4,6-trimethyl-

benzene)-1,3,5-triacetic acid) for the aerobic oxidation of benzyl alcohol to benzaldehyde,<sup>5</sup> and Cu<sub>3</sub>L<sub>2</sub> copper cage for A<sup>3</sup>-coupling reaction to generate propargylamines.<sup>6</sup> The catalytic activity of these cages can be enhanced by combining them with metal nanoparticles, which have exposed surfaces for generating active species and intermediates. The physicochemical features of metal nanoparticles are different from their parent forms and are incompatible with standard encapsulation procedures. Furthermore, unique surface properties, high surface energy, and a tendency to agglomerate in the solution phase make encapsulation a strenuous task.<sup>7,8</sup> However, limited reports are available on the encapsulation of metal nanoparticles in metal–organic cages due to the non-compatibility of the cavity size.

Plasmonic nanoparticles like Au, Ag, and Pt have emerged as promising materials due to their surface plasmon resonance effect and ability to harness visible light.<sup>9</sup> Cu nanoparticles also exhibit a plasmon resonance effect but they are not as pronounced as noble metals. Moreover, controlling the size and dispersion of copper nanoparticles is comparatively difficult due to their high reactivity. They are more susceptible to oxidation and their stability in aqueous systems is an important concern, especially in air or oxygen-rich environments.<sup>10</sup> Gold nanoparticles can act as catalysts in various oxidation and reduction reactions due to their unique electronic structures. However, they require suitable support to work heterogeneously<sup>11</sup> as they undergo agglomeration and show decreased catalytic

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† Electronic supplementary information (ESI) available: Characterization data of all compounds, elemental mapping, DLS and XPS spectra, PXRD pattern of Zn(II) cubane and ZnCC@AuNPs, HPLC chromatogram, crystallographic data and structural parameters; crystallographic data comprised in CCDC 2298615. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3tc03896c>

activity.<sup>12</sup> Earlier, some COFs, derived by reacting phenothiazine-based trialdehyde with 1,2-cyclohexanediamine<sup>13</sup> and benzene-1,3,5-tricarbaldehyde with 1,2-cyclohexanediamine,<sup>14</sup> and MOFs like Sn–Na MOF<sup>15</sup> and UiO-66 MOF<sup>16</sup> and multiwalled carbon nanotubes<sup>17</sup> have been reported as supports for holding noble metal nanoparticles. Self-assembled cages, macrocycles, polymers, and various other supramolecular structures derived from metallo-ligands constitute a prominent class with confined cavities inside the molecular frameworks and interactive functional sites.<sup>18</sup> Among these, the development of Zn(II) based cages has expanded the applications of inorganic molecules due to their versatile stereochemical properties, capability to form complexes with numerous ligands and catalytic potential.<sup>19</sup> Therefore, Zn(II) cubane cages can be considered as suitable structures to sustain gold nanoparticles for stabilization and improve catalytic efficiency.

Herein, we report a self-assembled Zn(II) metal–organic cubane cage embedded with Au nanoparticles for the selective conversion of furfural to 2-methyl furan. 2-Methyl furan is essential for many scientific and industrial processes such as the synthesis of organic compounds, formation of biofuels, and production of flavours and fragrances.<sup>20</sup> The selective hydrogenation of biomass-derived furfural can produce 2-methyl furan as one of the constituents of various value-added products (like furfuryl alcohol and cyclopentanone). The increasing biofuel demand owing to the depletion of fossil fuels has prompted the upgradation of renewable biomass into chemicals and biofuels for a sustainable future and lignocellulosic biomass can provide affordable and sustainable alternatives.<sup>21</sup> In the past, a variety of catalysts, including CuRe/Al<sub>2</sub>O<sub>3</sub>, Cu<sub>2.5</sub>ZnAl-600, CuFe<sub>2</sub>O<sub>4</sub>, iridium/carbon, Cu–Fe, Cu/ZnO, and Cu–Co/Al<sub>2</sub>O<sub>3</sub>, have been employed for the catalytic hydrogenation of furfural to 2-methylfuran but under harsh conditions and challenging protocols.<sup>22–29</sup> Moreover, the yield of 2-methylfuran is usually low because of the formation of various by-products such as tetrahydrofurfuryl alcohol and furan derivatives. However, the developed Zn(II)-based cubane-like oxocluster with small-sized pockets on self-assembly occupied by gold nanoparticles (AuNPs) provides excellent outputs for converting the biomass into 2-methylfuran in the present work. The catalytic hydrogenation of biomass-derived furfural to 2-methylfuran can be achieved in 1.5 h under mild reaction conditions by using 2-propanol. This is better than the classical hydrogenation methods where the direct addition of H<sub>2</sub> gas is required in the presence of a hydrogenation catalyst. The developed ZnCC@AuNPs composite is a pioneering example of gold nanoparticles supported within a metal cage, proving its practical application in biofuel production for the first time.

## 2. Experimental

### 2.1 Materials and methods

The compounds were synthesized using the Schlenk technique. 5-Bromo-2-hydroxybenzaldehyde, ethanol amine and furfural were procured from Aldrich, USA. Sodium borohydride, sodium

hydroxide, triethylamine, zinc nitrate hexahydrate, and gold chloride trihydrate were procured from SRL, India, in high purity. The solvents methanol, 2-propanol, and toluene (Finar, India) were dried using standard procedures. The reduced Schiff base (rSB) was synthesized using the procedure reported in our previous report.<sup>30</sup>

### 2.2 Instrumentation

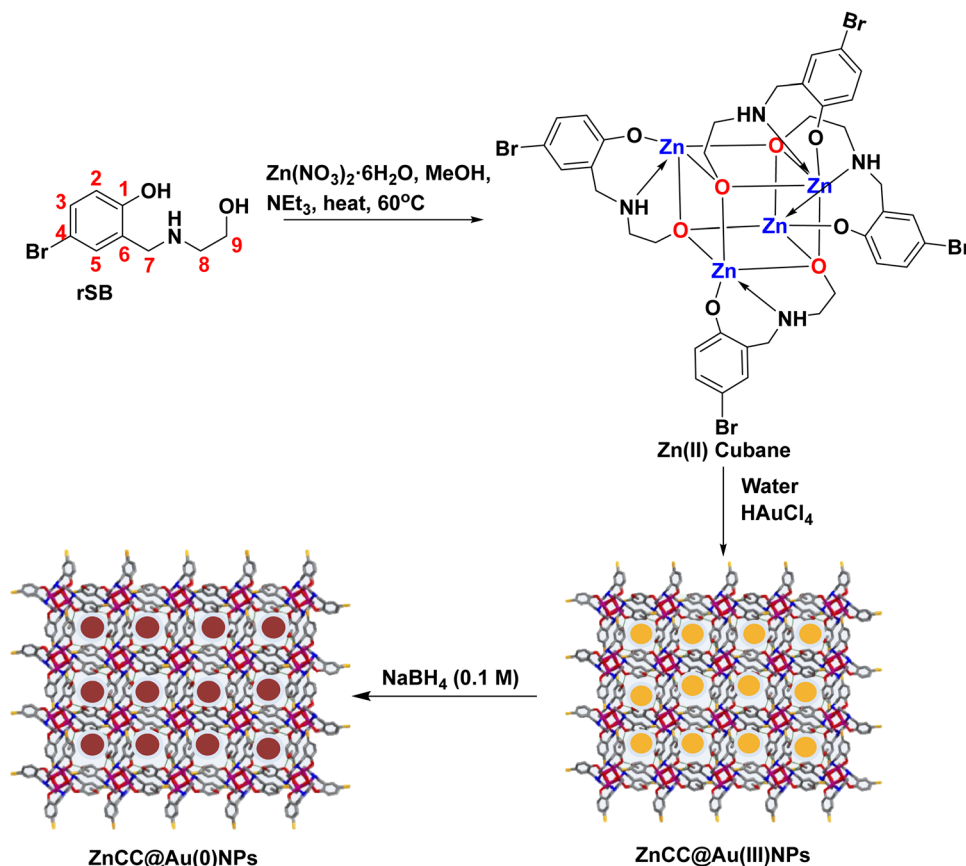
The FT-IR spectra, solution-phase NMR spectra (comprising <sup>1</sup>H and <sup>13</sup>C NMR spectra), and mass spectrometry measurements were conducted using the following instruments: BRUKER alpha Eco-ATR spectrometer, BRUKER AVANCE 500-NMR spectrometer, and VG Analytical (70-S) spectrometer, which was equipped with an ESI source operating at a capillary voltage of 2500 V. The crystallographic data for the single crystal of the compound were collected at 165 K using a Rigaku XTA Lab SuperNova with Mo K $\alpha$  radiation ( $\lambda = 0.71073$ ). The crystal structure was solved using Olex2 software, utilizing the intrinsic phasing method of the ShelXT structure solution program. Subsequently, the structure was refined using the least-squares method, which was integrated into the ShelXL refinement package.<sup>31,32</sup> For structural optimization and frequency calculations, density functional theory (DFT) was employed using Gaussian 16 software.<sup>33</sup> Becke's three-parameter hybrid correlation functional B3LYP was applied, along with the 6-31+G(d,p) and LanL2DZ basis set, for the geometry optimization process.<sup>34</sup> A Waters 2998 System, which consisted of a photodiode array detector and a binary HPLC pump with a Supelco Analytical C18 column (25 cm  $\times$  4.6 mm  $\times$  5  $\mu$ m) was utilized for the analysis.

### 2.3 Synthetic procedures

**2.3.1 Zn(II) cubane cage.** In a double-neck round bottom flask, a solution of rSB (8.19 mmol, 2.00 g) was dissolved in 70 mL dry methanol. Triethylamine (11.18 mmol, 1.13 g) and zinc nitrate hexahydrate (8.11 mmol, 2.41 g) were added and heated to reflux under an inert atmosphere for 4 hours. The volume of the solution was reduced to half by collecting the solvent using a Dean–Stark apparatus. The slow evaporation of the concentrated solution yielded white crystals that were suitable for single-crystal X-ray diffraction.

Yield (crystals): 32.34% (3.20 g, 2.62 mmol). FT-IR (KBr) cm<sup>-1</sup>: 3432 (N–H), 2900–2750 (CH<sub>2</sub>), 1586, 1472 (C–C), 1282 (C–N), 1180 (C–O), 641 (C–Br), 548 (Zn–O). <sup>1</sup>H-NMR (500 MHz, DMSO-d<sub>6</sub>):  $\delta$  (ppm) 2.78 (8H, broad singlet, H<sup>8</sup>), 3.62 (8H, broad singlet, H<sup>9</sup>), 3.80 (8H, s, H<sup>7</sup>), 4.15 (4H, s, –NH), 6.48 (4H, broad singlet, H<sup>2</sup>), 7.11 (8H, broad singlet, H<sup>3</sup>, H<sup>5</sup>); ESI-MS (*m/z*): 647.49 [M/2 + H]<sup>+</sup>.

**2.3.2 Encapsulation of AuNPs in the Zn(II) cubane cage (ZnCC@AuNPs).** Gold nanoparticles (AuNPs) were encapsulated in Zn(II) cubane using a dispersion and reduction method. Zn(II) cubane (50 mg) was ground to a fine powder using a mortar and pestle and dispersed in 10 mL distilled water by sonication. An aqueous solution of HAuCl<sub>4</sub> (10 mg in 2 mL) was added dropwise under vigorous stirring. The vial was covered with aluminium foil and stirred for 6 hours under dark conditions. The precipitates were collected by centrifugation



Scheme 1 Synthesis of rSB, Zn(II) cubane, and ZnCC@AuNPs.

(at a rate of 8000 rpm/10 min), washed with distilled water and redispersed in 3 mL water. A 0.50 mL of freshly prepared NaBH<sub>4</sub> solution (0.1 M) was added slowly, which changed the colour from wine-red to black and the solution was further stirred for 6 hours. Finally, ZnCC@AuNPs were collected by centrifugation (Scheme 1).

**2.3.3 Conversion of furfural to 2-methyl furan.** In a 25 mL round bottom flask, furfural (1.0 g, 10.40 mmol), NaOH (83 mg, 2.07 mmol) and ZnCC@AuNPs (10 mg) were dissolved in 15 mL of 2-propanol. The reaction mixture was stirred under a nitrogen atmosphere for 1.5 h at 60 °C. The contents were centrifuged and the product was separated by filtration. The catalyst was washed with ethanol and dried for the next reaction cycle and the resulting liquid products were analysed using HPLC.

## 2.4 Product analysis

The product was analysed for the formation of 2-methyl furan by HPLC-PDA system using an isocratic method with ACN:MeOH/75:25 (%) as the solvent flowing at a rate of 0.8 mL min<sup>-1</sup>. The following equations were employed to calculate the selectivity (%) of 2-methyl furan and the conversion (%) of furfural:

$$\text{2-methyl furan selectivity (mol\%)} = 100 \times \frac{C_{2\text{-MF}}}{C_{\text{Total products}}}$$

$$\text{Furfural conversion (mol\%)} = 100 \times \left(1 - \frac{C_{\text{Fur}}}{C_{\text{Fur}^{\circ}}}\right)$$

In these equations,  $C_{2\text{-MF}}$  and  $C_{\text{Total products}}$  represent the concentrations of 2-methylfuran and total products, respectively, at a specific reaction time ( $t$ ).  $C_{\text{Fur}^{\circ}}$  denotes the initial concentration of furfural before the reaction.

## 3. Results and discussion

### 3.1 Synthesis and characterization of the Zn(II) cubane cage

A tridentate ligand (rSB) was synthesized by the Schiff base condensation of 5-bromosalicylaldehyde and ethanolamine followed by reduction using NaBH<sub>4</sub> in methanol. The reaction of rSB with zinc nitrate hexahydrate in the presence of triethylamine (as a base) in methanol yielded a Zn(II) cage consisting of Zn<sub>4</sub>O<sub>4</sub> cubanes with coordinated ligand pendants as side bridges (Scheme 1).

The Zn(II) cubane cage was characterized using FTIR, <sup>1</sup>H NMR, and ESI-MS. The FT-IR spectrum of rSB showed a strong band at 3413 cm<sup>-1</sup> due to N-H stretching and a broad band in the region 3300–2960 cm<sup>-1</sup> due to O-H stretching. However, the band in the region 3470–3400 cm<sup>-1</sup> due to N-H stretching was broadened in the case of Zn(II) cubane, and the O-H stretch disappeared showing coordination of the ligand *via* NH and deprotonated -OH. Besides, a strong band at 624 cm<sup>-1</sup> and a weak band at 479 cm<sup>-1</sup> were observed due to C-Br stretching and Zn-O stretching, respectively. The <sup>1</sup>H-NMR spectrum of rSB in DMSO-*d*<sub>6</sub> showed the appearance of a singlet at 3.80 ppm due to the -CH<sub>2</sub> protons (H<sup>7</sup>). The NH



**Fig. 1** (a) Ortep view of Zn(II) cubane, (b) framework of the  $Zn_4O_4$  cubane cage, (c) wireframe representation of interactions between crystallized cubane units, (d) crystal packing of Zn(II) cubane showing 2-D sheet-like view, (e) space-fill representation of Zn(II) cubane (for non-hydrogen atoms, the radius of hydrogen atoms was eliminated for clarity).

proton appearing at 5.92 ppm merged with the signal of the alcoholic OH. In Zn cubane, triplets of  $H^8$  and  $H^9$  merged into a broad singlet at 2.78 and 3.62 ppm, respectively. The NH signal was seen as a singlet at 4.15 ppm and aromatic signals showing doublet ( $H^2$ ), doublet of doublet ( $H^3$ ) and doublet ( $H^5$ ) in rSB were transformed into singlets at 6.48 ppm for  $H^2$  and 7.11 ppm for  $H^3$  and  $H^5$ . The ESI-MS for Zn cubane exhibited a  $[M/2 + H]^+$  peak at  $m/z$  647.49 (see ESI,† Fig. S1–S3).

Zn(II) cubane was obtained in crystalline form by cooling the reaction mixture. The structure of Zn(II) cubane was elucidated by single-crystal X-ray diffraction. Table S1 (ESI†) provides a summary of the crystallographic data and structure refinement parameters, while Table S2 (ESI†) presents selected measurements of the bond lengths and bond angles. The Zn(II) cubane crystallized in the tetragonal space group  $I4_1/a$ . It consists of a distorted cubane-like cavity formed by joining the Zn and O atoms of four asymmetric units on the alternate corners of the cube. Each Zn atom was coordinated to five atoms by a nitrogen atom, aryl oxygen, and alkoxide oxygen atom of one ligand and alkoxide oxygen donors of two other ligands. Overall, the Zn(II) atoms showed distorted trigonal bipyramidal geometry, where all four ligands were coordinated in  $\mu^3$ - $\eta^1$ : $\eta^1$ : $\eta^3$  bridging mode with alkoxide oxygen coordinated in  $\eta^3$  mode with Zn(II) centres. The Zn–N, Zn–O1 and Zn–O2 bond lengths were in the usual ranges of 1.479, 1.965 and 2.428 Å, respectively (Fig. 1). The Zn(II) cubane was crystallized as a methanol solvate, which created hydrogen-bonded self-assembly with other cubane units generating voids ranging from 1.6 to 1.8 nm. This type of hydrogen-bonded cage is produced by the solvation of building blocks due to the prevalent force of hydrogen bonds.<sup>1</sup>

### 3.2 Synthesis and characterization of ZnCC@AuNPs

The encapsulation of nanoparticles is important for catalytic applications because it enhances surface activity and stability

and prevents aggregation. However, the encapsulation of gold nanoparticles in a protective Zn(II) cage is challenging as it requires Au-supportive interactions. Further, the generation of the appropriate cavity for encapsulation depends upon the geometry of the cage, the arrangement of metal ions and the choice of ligands. Schiff base ligands serve as excellent supporting molecules for such materials because they can be derived with the desired number of functionalities through a highly efficient and straightforward synthesis.<sup>35,36</sup> Reduced Schiff bases are more flexible than Schiff base complexes due to the absence of  $\pi$ -conjugation, which may allow them to attain any geometry.<sup>37</sup> Here, the flexibility and multidenticity of the ligand led to the construction of the Zn(II) cubane cage with a small-sized A-type cavity. The cubane units self-assembled to form a framework equipped with B-type cavities. The appropriate size of the B-type cavities facilitated the encapsulation of Au nanoparticles. To gain insight into the size, shape, composition, encapsulation, and surface properties of ZnCC@AuNPs, various studies like FTIR, TGA,  $^1H$  NMR, BET, SEM, TEM, XPS, and PXRD analyses were conducted.

The FT-IR spectrum of ZnCC@AuNPs was found to be similar to that of Zn(II) cubane showing a strong vibrational band in the region  $3500$ – $3200$   $cm^{-1}$  for N–H stretching and O–H stretching. Additional bands consisting of strong split vibrational bands centred at  $1600$  and  $1637$   $cm^{-1}$  due to C=C stretching, a strong band at  $621$   $cm^{-1}$  due to C–Br and a weak band at  $480$   $cm^{-1}$  due to Zn–O stretching were also observed in the region similar to that of Zn(II) cubane (Fig. 2a). Two weak vibrational bands were also observed at  $1474$   $cm^{-1}$  and  $1386$   $cm^{-1}$  for C–H bending and C–N stretching, respectively, for both Zn(II) cubane and ZnCC@AuNPs. This revealed that the incorporation of AuNPs barely changed the structural framework and chemical composition of Zn(II) cubane.

The thermal stabilities of Zn(II) cubane and ZnCC@AuNPs were studied using thermogravimetric analysis (Fig. 2b). Weight losses of 11.73% and 7.36% were observed in the first step in the thermograms of Zn(II) cubane and ZnCC@AuNPs, respectively, up to  $200$   $^{\circ}C$ , which is generally due to the loss of absorbed water or solvent (methanol). Weight losses of 30.36% and 45.98% were observed in the second step when the temperature was increased from  $200$  to  $800$   $^{\circ}C$ , which may be attributed to the decomposition of the organic framework of Zn(II) cubane and ZnCC@AuNPs, respectively. The third weight loss of 14.69% can be related to the formation of the metal residue ZnO when the temperature was increased above  $800$   $^{\circ}C$ .

The  $^1H$  NMR spectra of Zn(II) cubane and ZnCC@AuNPs were compared. The  $^1H$  NMR spectrum of Zn(II) cubane showed a significant shift in the N–H proton and broadening of alkyl and aromatic protons due to the complexation of Zn(II) with the ligand. The broadening may be attributed to the limited mobility with rapid spin relaxation of skeletal protons in the complex.<sup>38,39</sup> The  $^1H$  NMR spectra of ZnCC@AuNPs showed a shift in the N–H proton signal from 4.15 ppm to 3.81 ppm, thus validating the increase in the electron density due to the binding of AuNPs. Although this suggested the presence of AuNPs inside the cage skeleton, the solvent effect cannot be ignored at the same time (see ESI,† Fig. S4).



Fig. 2 (a) A comparison of FT-IR spectra of Zn(II) cubane and ZnCC@AuNPs. (b) TGA thermogram of Zn(II) cubane and ZnCC@AuNPs.

Nitrogen gas adsorption–desorption isotherms were used to study the adsorption properties of Zn(II) cubane and ZnCC@AuNPs. The BET and the BJH models were used to compute the specific surface area and pore size distribution of Zn(II) cubane and ZnCC@AuNPs. The adsorption–desorption isotherms exhibited an H3 hysteresis loop suggesting the porous nature of the material. The surface area and pore volume of the Zn(II) cubane were calculated to be  $19.107 \text{ m}^2 \text{ g}^{-1}$  and  $4.390 \text{ cm}^3 \text{ g}^{-1}$ , respectively. However, after the incorporation of gold, the surface area and pore volume decreased to  $2.321 \text{ m}^2 \text{ g}^{-1}$  and  $0.5066 \text{ cm}^3 \text{ g}^{-1}$ , respectively (see ESI,† Fig. S5a and b). The  $\text{N}_2$  sorption isotherm of ZnCC@AuNPs exhibited a gradual decrease in both surface area and pore volume, suggesting the blockage of cavities due to the encapsulation of Au nanoparticles in the vacant spaces. The BJH results of Zn(II) cubane and ZnCC@AuNPs showed main peaks at 7.14 nm and 3.65 nm, respectively, for the pore size (see ESI,† Fig. S5c and d). The significant reduction in the pore size of the Zn(II) cubane also confirmed the occupancy of cavities by the Au nanoparticles.

Field emission scanning electron microscopy (FE-SEM) images of Zn(II) cubane revealed a flaky material exhibiting a rough morphology (Fig. 3a–c). However, the infusion of

spherical particles was visible in the flakes of ZnCC@AuNPs, confirming the presence of Au nanoparticles (Fig. 3d–f). However, Au nanoparticles were observed over the surface of the flakes, which may be attributed to the agglomeration of larger particles ( $> 1.60 \text{ nm}$ ) during sample preparation, preventing their entry into the pores. The elemental mapping confirmed that the composition of the cubane cage consisted of C, H, N, Br, and Zn in the Zn(II) cubane and the inclusion of Au in the ZnCC@AuNPs cage. This revealed a symmetrical distribution of each composite on the surface of Zn(II) cubane and ZnCC@AuNPs (see ESI,† Fig. S6).

The size and morphology of the synthesized ZnCC@AuNPs were studied by transmission electron microscopy (TEM). The size distribution calculated from the TEM images revealed the formation of Au nanoparticles with an average particle diameter of 1.55 nm (Fig. 4a). The single-crystal X-ray diffraction data revealed the presence of Zn(II) cubane units with cavity 'A' of length 0.3 nm. These units further unite to form self-assembly that creates voids 'B' with a length of 1.651 nm. It can be inferred from the size distribution studies that only void B can provide appropriate space to accommodate Au nanoparticles and the spaces inside the cubane units (cavity A) are not approachable for particles of 1.55 nm in diameter

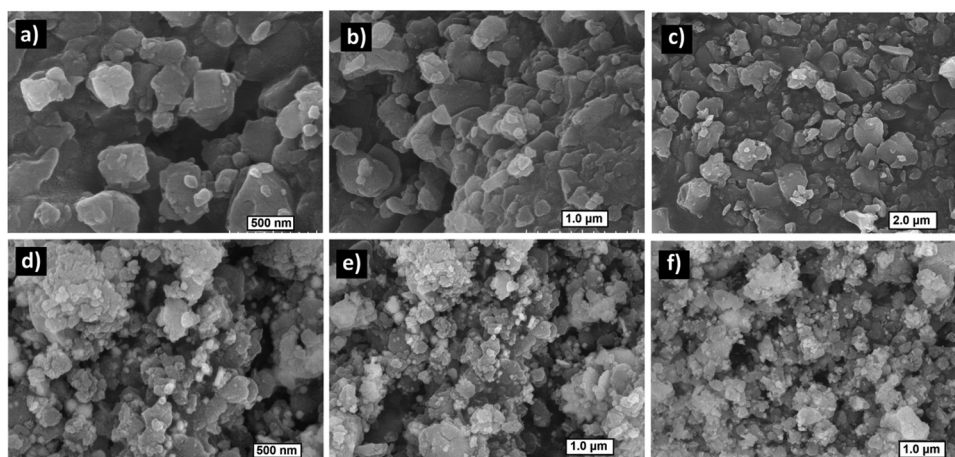


Fig. 3 (a)–(c) FE-SEM images of Zn cubane and (d)–(f) SEM images of ZnCC@AuNPs.

(Fig. 4b).<sup>14</sup> The dynamic light scattering (DLS) study supported the formation of Au nanoparticles in a narrow size distribution range with a mean particle size of 1.2 nm (see ESI,† Fig. S7).

The mass loading % of  $\text{HAuCl}_4$  was optimized by varying the concentration of  $\text{HAuCl}_4$ . This resulted in the formation of nanoparticles with a size range of 1.4–2.1 nm and 20–25 nm for concentrations of 16.48 mM and 25.0 mM, respectively (Fig. 4c and d). The increase in the concentration of  $\text{HAuCl}_4$  resulted in increased particle size due to the production of quasi-spherical nanoparticles with a much larger size distribution of 20–25 nm (Fig. 4e). Energy dispersive X-ray analysis (EDAX) measurements of  $\text{ZnCC@AuNPs}$  were carried out to illustrate the presence of Au nanoparticles in the  $\text{Zn(II)}$  cubane. The EDAX spectrum shows the coexistence of C, N, O, Br, Zn and Au in the material (Fig. 4f).

X-ray photoelectron spectroscopy (XPS) studies of  $\text{ZnCC@AuNPs}$  revealed the presence of Au in the zero oxidation state as

designated by its characteristic binding energy values of 87.7 eV and 83.9 eV corresponding to two distinct spin-orbit pairs  $4f_{5/2}$  and  $4f_{7/2}$ , respectively (Fig. 5a). The peak area ratio for spin-orbit pairs  $\text{Au(0)} 4f_{5/2}:4f_{7/2}$  is 1:0.5. Also, there is one weak spin-orbit pair at 89.9 eV and 84.4 eV corresponding to the  $\text{Au(III)}$  state. The peak area ratio for spin-orbit pairs  $\text{Au(III)} 4f_{5/2}, 4f_{7/2}$  is 0.32 and 0.16 compared to  $\text{Au(0)} 4f_{5/2}$ . The Zn 2p in Zn cubane and  $\text{ZnCC@AuNPs}$  showed intense peaks at binding energies of around 1045 eV and 1021 eV corresponding to two distinct spin-orbit pairs  $2p_{1/2}$  and  $2p_{3/2}$ , respectively (Fig. 5b and c). Additionally, the XPS spectra of C 1s, O 1s, Br 3d and N 1s of  $\text{Zn(II)}$  cubane and  $\text{ZnCC@AuNPs}$  were found as expected (given in the ESI†) (see Fig. S8 and S9).

PXRD analysis of  $\text{Zn(II)}$  cubane was performed to determine the purity and crystallinity of the bulk material. The PXRD pattern of the  $\text{Zn(II)}$  cubane obtained experimentally showed similarity with the predicted pattern obtained from Mercury

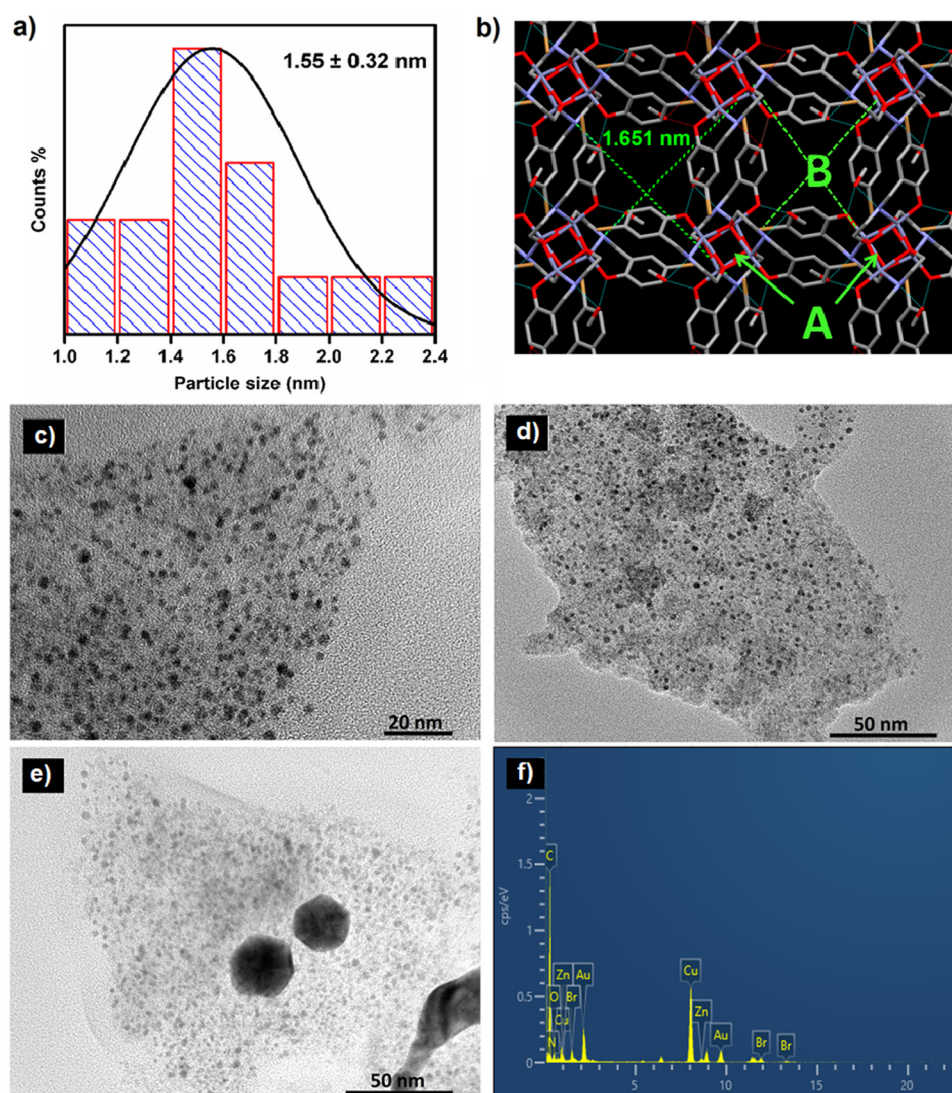


Fig. 4 (a) Particle size distribution of AuNPs, (b) crystal image of the Zn cubane template showing void 'B' of size 1.651 nm, (c)–(e) HR-TEM images of  $\text{ZnCC@AuNPs}$  at different mass loadings, and (f) elemental distribution analysis of  $\text{ZnCC@AuNPs}$ .

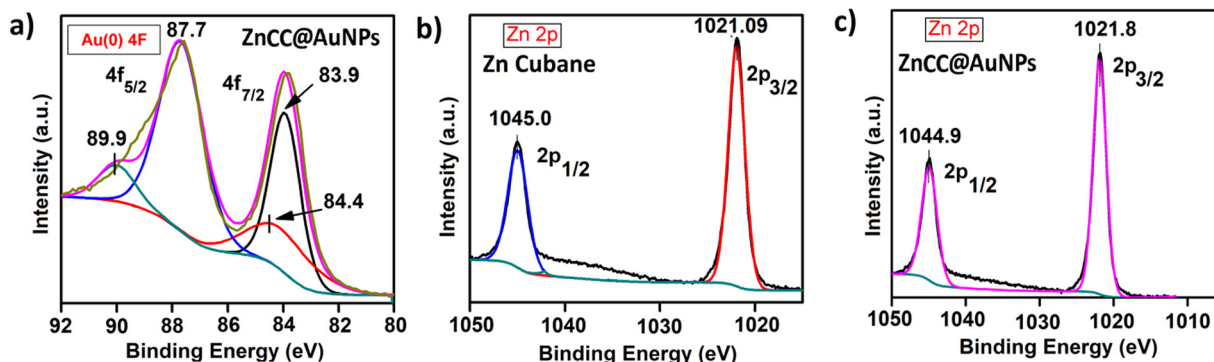


Fig. 5 XPS of (a) Au(0) 4F of ZnCC@AuNPs, (b) Zn 2p of Zn cubane, and (c) Zn 2p of ZnCC@AuNPs.

3.8, demonstrating the rational crystalline phase purity of the bulk sample (see ESI,† Fig. S10a). Similarly, the ZnCC@AuNPs showed strong diffraction peaks at  $38.14^\circ$  and  $44.29^\circ$ , which correspond to the (111) and (200) lattice planes, respectively. These lattice planes are typical of Au with a face-centred cubic (fcc) structure (see ESI,† Fig. S10b).

The UV absorption spectrum of ZnCC@AuNPs exhibited a new band centred at 530 nm due to localized surface plasmon resonance (LSPR) (see ESI,† Fig. S11b). A Tauc plot was employed to determine the band gap energy of ZnCC@AuNPs, using the relationship  $(\alpha h\nu)^2 = h\nu - E_g$ , where  $\alpha$  and  $E_g$  represent the absorption coefficient and the energy gap of the semiconductor, respectively. The resulting linear fit of  $(\alpha h\nu)^2$  versus photon energy ( $h\nu$ ) provided the band gap values of 1.93 and 3.17 eV for ZnCC@AuNPs due to AuNPs and Zn(II), respectively (Fig. 6a). To understand the enhanced photocatalytic activity of ZnCC@AuNPs compared to that of Zn(II) cubane, electrochemical impedance spectroscopy (EIS) was conducted. The Nyquist plot revealed a significantly smaller arc for ZnCC@AuNPs than for Zn(II) cubane, indicating a lower charge-transfer resistance and faster interfacial charge-transfer rate (Fig. 6b). Photoluminescence investigations demonstrated a similar characteristic band for Zn(II) cubane and ZnCC@AuNPs. However, the emission band intensity centred at 450 nm was notably lower for ZnCC@AuNPs, suggesting reduced electron/hole pair recombination and superior photocatalytic activity (Fig. 6c). The unique optical properties of gold nanoparticles,

specifically the surface plasmon effect in gold-doped materials, contribute to their interaction with fluorescence.<sup>40</sup> Comparable photoluminescence responses were observed for Au–ZnO nanocomposites with ZnO nanorods.<sup>41</sup> The association of Zn(II) cubane with Au ions in ZnCC@AuNPs resulted in improved LSPR, enhanced charge transfer, and consequently superior photocatalytic efficiency.

### 3.3 Catalytic conversion of furfural

The conversion of furfural to 2-methylfuran is an important reaction in the biofuel industry. The selectivity and yield of the reaction depend on different variables like the effect of the metal, catalyst loading, reaction temperature and solvent. The solvent plays a unique catalytic role in the hydrogenation of furfural which cannot be ignored because the reaction is in the liquid phase. Previous reports confirmed that nonpolar solvents (*i.e.* toluene) lower the rate of the hydrogenation reaction and water renders the decarbonylation of furfural alcohol to 2-methylfuran. Therefore, the use of polar solvents like methanol (MeOH), ethanol (EtOH) and isopropanol (*i*-PrOH) is a wise selection as they donate hydrogen for furfural hydrogenation. In addition, the hydrogenation of furfural to 2-methylfuran in the presence of toluene and water was not catalysed by the developed ZnCC@AuNP catalyst (Table 1). However, 100% conversion of furfural was observed in *i*-PrOH followed by 30% in methanol and 28% in ethanol. The variation in the catalyst amount from 5 mg to 12 mg suggested that 10 mg of

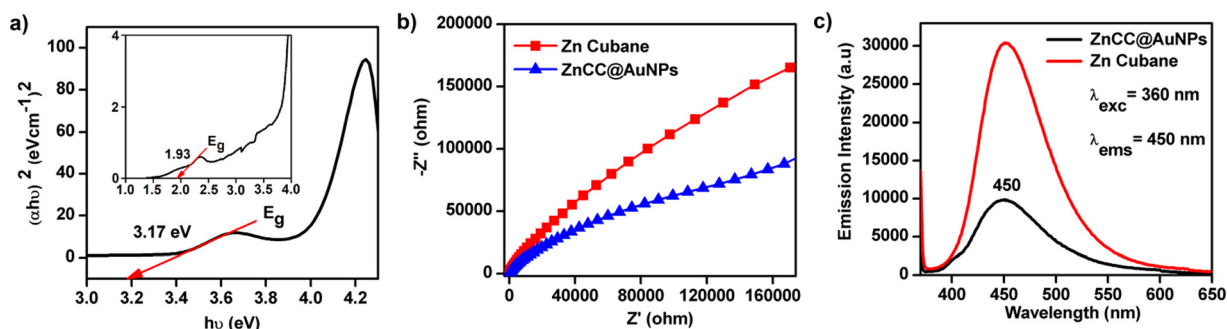


Fig. 6 (a) Tauc plot of ZnCC@AuNPs showing the band gap  $E_g$  (inset: magnified view of the Tauc plot showing a weak band due to LSPR), (b) EIS of Zn cubane and ZnCC@AuNPs, and (c) photoluminescence response of Zn(II) cubane and ZnCC@AuNPs.

Table 1 Optimization reaction with the ZnCC@AuNP catalyst



S. no.	Catalyst amount (mg)	Solvent	Temperature (°C)	Time (min)	Conversion (%)	Selectivity (%)	
						2-MF	FR
1.	0	i-PrOH	60	90	36	>99	—
2.	5	i-PrOH	60	90	64	>99	—
3.	8	i-PrOH	60	90	94	>99	—
4.	10	i-PrOH	30	60	62	>99	—
5.	10	i-PrOH	60	60	100	97	—
6.	10	i-PrOH	60	90	100	>99	—
7.	12	i-PrOH	60	120	100	98.40	1.59
8.	10	i-PrOH	100	120	100	97.90	2.10
9.	10	i-PrOH	120	120	100	97.30	2.70
10.	10	i-PrOH	150	120	100	97.10	2.90
11.	12	i-PrOH	150	120	100	96.90	3.10
12.	10	H <sub>2</sub> O	60	90	0	NR	—
13.	10	Tol	60	90	0	NR	—
14.	10	MeOH	60	90	30	>99	—
15.	10	EtOH	60	90	28	>99	—
16.	10	i-PrOH	60	90	40 <sup>a</sup>	>99	—

Reaction conditions: visible light, furfural (10.40 mmol), NaOH (2.07 mmol), solvent (15 mL). <sup>a</sup> Reaction conditions: dark, furfural (10.40 mmol), NaOH (2.07 mmol), solvent (15 mL). \*2-MF = 2-methylfuran, FR = furan, i-PrOH = 2-propanol, H<sub>2</sub>O = water, Tol = toluene, MeOH = methanol, EtOH = ethanol, NR = no result.

ZnCC@AuNPs was sufficient for 100% conversion of the chosen amounts of furfural (1.0 g, 10.40 mmol) into 2-methylfuran. By decreasing the amount of ZnCC@AuNPs from 10 mg to 5 mg, the conversion rate decreases from 100% to 64%. It can be inferred that the lower amounts of ZnCC@AuNPs could not generate sufficient H<sup>+</sup> for the complete hydrogenation of furfural to 2-methylfuran.

The impact of temperature on the hydrogenation process was examined within a temperature range of 30 °C to 150 °C while keeping the other reaction conditions the same. The temperature change resulted in variations in the product yield due to the formation of two products 2-methylfuran and furan. When the reaction was carried out at 60 °C, 100% conversion of furfural with >99% of 2-methylfuran was achieved. However, by increasing both the amount of catalyst and the temperature, the furan yield increased from 1.59% to 3.10% (see ESI,† Fig. S12). This increase can be attributed to the increase in the surface area of gold, which facilitates the decarbonylation process. These findings indicate that the hydrogenation of the aldehyde group proceeded effectively under relatively mild conditions with 60 °C being identified as the optimal temperature. Further, the reaction was carried out in the dark in the presence of ZnCC@AuNPs under identical reaction conditions and only the conversion of 40% 2-methylfuran was detected. A summary of the results is presented in Table 1. The formation of 2-methylfuran was analyzed using <sup>1</sup>H and <sup>13</sup>C NMR spectroscopy (see ESI,† Fig. S13 and S14), as well as the HPLC-PDA method where a peak was detected at a retention time of 3.42 min (see ESI,† Fig. S15). A time-scaled UV-visible spectrophotometric experiment (0–90 min) was

performed to monitor the progress of the catalytic transformation reaction under the optimized conditions (see ESI,† Fig. S16). The intensity of the absorption band at 274 nm corresponding to furfural showed a significant decrease with time, indicating the consumption of furfural during the conversion. Consequently, a new absorption band appeared at 326 nm, indicating the formation of 2-methylfuran. Table S3 (ESI†) summarises a literature review of several catalysts employed in the catalytic hydrogenation of furfural to 2-methylfuran. The prepared ZnCC@AuNP catalyst provided the mildest reaction conditions for the complete conversion of furfural into 2-methylfuran.

### 3.4 Heterogeneity, recyclability studies and leaching tests

The reusability of ZnCC@AuNPs as a catalyst for the hydrogenation of furfural to 2-methylfuran was examined under optimized conditions. After each reaction cycle, the recovered catalyst powder was washed with ethanol and then dried. The selectivity of the catalyst for the formation of 2-methylfuran was maintained for up to five consecutive reaction runs (see ESI,† Fig. S17). However, the catalytic hydrogenation activity gradually decreased, with a decrease in the conversion % from 99.9% to 88.6% in the fifth run. This decrease is due to a minor loss of the catalyst during the recovery procedure, *i.e.* separation and washing processes. A loss of 1.6 mg was observed over five catalytic cycles, which reduced the conversion.

### 3.5 Mechanism

The exact route of catalysis was investigated based on literature reports, *in situ* ATR-IR kinetic studies, and computational

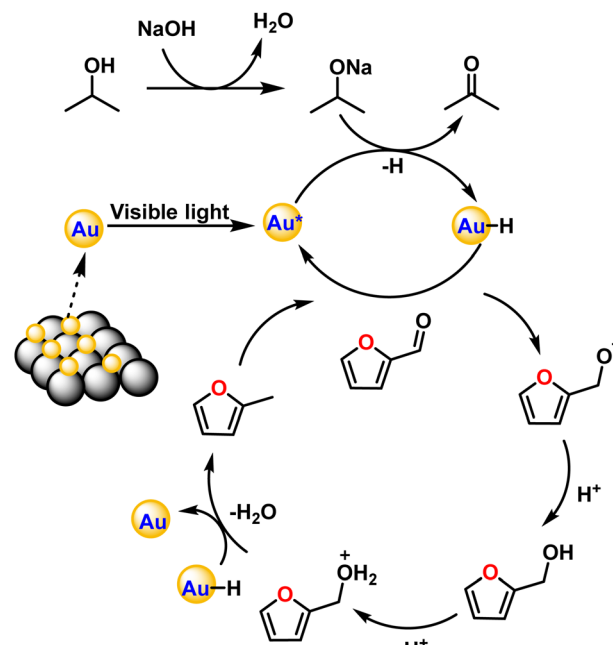


Fig. 7 A plausible mechanism pathway for the hydrogenation of furfural to 2-methylfuran.



studies. First, NaOH abstracts hydrogen from 2-propanol to form sodium isopropoxide. Consecutively, light is absorbed by AuNPs due surface plasmon resonance effect which excites electrons from the valence band to the conduction band. This electron transfers to isopropoxide, which loses  $H^-$  and results in the formation of Au–H species leaving acetone as an oxidized product.<sup>42</sup> The surface hydride is transferred to furfural yielding furfuryl alcohol as an intermediate after the supply of  $H^+$  ions. Further, the transfer of hydrides leads to the formation of 2-methyl furan and water as a side product (Fig. 7). The C–C bond cleavage is usually observed as a side reaction during the hydrogenation of furfural and the decarbonylation reaction is frequently observed as a side reaction on the surface of metal catalysts. Decarbonylation involves adsorption of species on the metal surface, where both the furan ring and the formyl group strongly interact with the metal atoms resulting in the formation of furan and the side product carbon monoxide.

ATR-IR spectra were recorded *in situ* during the catalytic hydrogenation of furfural under optimized conditions at regular intervals of 30 min. The IR spectrum of furfural exhibited a strong absorption band at  $1690\text{ cm}^{-1}$  due to the carbonyl group (C=O). The split bands at  $2812$  and  $2855\text{ cm}^{-1}$  due to C–H stretching and C–H bending at  $1392\text{ cm}^{-1}$  are characteristic

vibrations of the aldehyde group. The bands at  $1464$  and  $1566\text{ cm}^{-1}$  are ascribed to the C=C bond and that at  $743\text{ cm}^{-1}$  to the C–H bond of the furan ring. At 0 min, after the addition of *i*-PrOH and the catalyst, there was no change in the observed spectrum of furfural. After 30 min, the C=O band (at  $1690\text{ cm}^{-1}$ ) and two double C–H bands (at  $2812$  and  $2855\text{ cm}^{-1}$ ) disappeared while new bands at  $2966$  and  $3344\text{ cm}^{-1}$  appeared for C–H stretching and O–H, respectively, confirming the presence of furfuryl alcohol (FAL). Also, the intensity of the C=C stretching vibrations (at  $1464$  and  $1566\text{ cm}^{-1}$ ) of furfural weakened due to the loss of conjugation between C=O and C=C. After 60 min, the O–H band disappeared completely and only three bands, *i.e.* furan ring C–H bending at  $773\text{ cm}^{-1}$ , C–O stretching at  $1218\text{ cm}^{-1}$  and C–H stretching at  $2923\text{ cm}^{-1}$  were observed, confirming the formation of 2-methylfuran (2-MF). After 90 min, the C–H band was predominant, which clearly explains the complete conversion of furfural to 2-methylfuran (Fig. 8a).

DFT studies were performed according to a plausible mechanism from the literature reports and ATR-IR studies. It is inferred that the Au–H bond transfers hydride to furfural, ultimately resulting in the formation of the intermediate furfuryl alcohol. The hydrogenation of furfural to furfuryl alcohol is an exothermic reaction with a loss of energy equivalent to  $-15.59\text{ eV}$  (Fig. 8b). It is important to recognize that the conversion of furfuryl alcohol to 2-methylfuran through hydrogenation is an exothermic process, releasing energy of  $-40.32\text{ eV}$ . The calculated energy loss was significantly lower than that of furfuryl alcohol. This difference was attributed to the inherent stability of the furanic ring structure, which facilitated the breaking of the C–OH bond in furfuryl alcohol, leading to the formation of 2-methylfuran.



Fig. 8 (a) *In situ* ATR-IR spectra of furfural (FUR) conversion to 2-methylfuran (2-MF) with furfuryl alcohol (FAL) as an intermediate at  $60\text{ }^\circ\text{C}$  for 90 min in *i*-PrOH at regular intervals of 30 min and (b) energy profile diagram of catalytic conversion of furfural (FUR) to 2-methylfuran (2-MF).

## 4. Conclusion

A hydrogen-bonded  $Zn_4O_4$  cubane cage assembly that could encapsulate gold nanoparticles was obtained. It catalyzes the hydrogenolysis of bio-mass derived furfural to 2-methylfuran under moderate conditions, which is classified as a value-added product. The ZnCC@AuNP catalyst facilitated the hydrogenation of furfural to 2-methylfuran at  $60\text{ }^\circ\text{C}$  using 2-propanol as a solvent in just 1.5 h of a reaction time. Under visible light, absorption by AuNPs excited electrons in the conduction band and transferred them to isopropanol. This generated Au–H species on the Au-NP surface and reduced the isopropanol to acetone. High selectivity with a product yield  $>99\%$  of 2-methylfuran was achieved in the reaction. Furthermore, the ZnCC@AuNP catalyst displayed satisfactory structural stability and catalytic stability during the recycling reaction tests.

## Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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