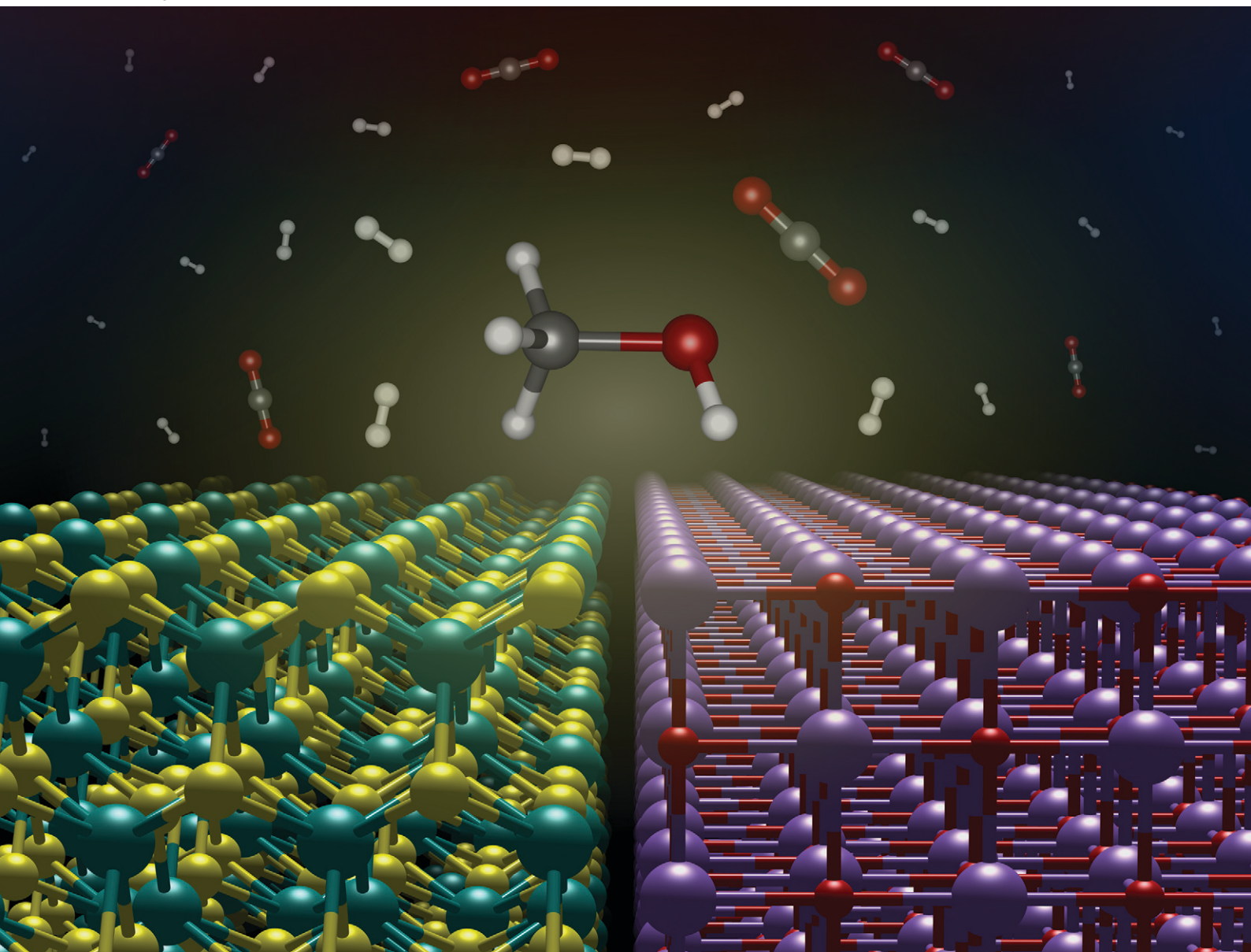


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## PAPER

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# Mn-promoted MoS<sub>2</sub> catalysts for CO<sub>2</sub> hydrogenation: enhanced methanol selectivity due to MoS<sub>2</sub>/MnO<sub>x</sub> interfaces†

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Considering the alarming scenario of climate change, CO<sub>2</sub> hydrogenation to methanol is considered a key process for phasing out fossil fuels by means of CO<sub>2</sub> utilization. In this context, MoS<sub>2</sub> catalysts have recently shown to be promising catalysts for this reaction, especially in the presence of abundant basal-plane sulfur vacancies and due to synergistic mechanisms with other phases. In this work, Mn-promoted MoS<sub>2</sub> prepared by a hydrothermal method presents considerable selectivity for CO<sub>2</sub> hydrogenation to methanol in comparison with pure MoS<sub>2</sub> and other promoters such as K and Co. Interestingly, if CO is used as a carbon source for the reaction, methanol production is remarkably lower, which suggests the absence of a CO intermediate during CO<sub>2</sub> hydrogenation to methanol. After optimization of synthesis parameters, a methanol selectivity of 64% is achieved at a CO<sub>2</sub> conversion of 2.8% under 180 °C. According to material characterization by X-ray Diffraction and X-ray Absorption, the Mn promoter is present mainly in the form of MnO and MnCO<sub>3</sub> phases, with the latter undergoing conversion to MnO upon H<sub>2</sub> pretreatment. However, following exposure to reaction conditions, X-ray photoelectron spectroscopy suggests that higher oxidation states of Mn may be present at the surface, suggesting that the improved catalytic activity for CO<sub>2</sub> hydrogenation to methanol arises from a synergy between MoS<sub>2</sub> and MnO<sub>x</sub> at the catalyst surface.

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## Introduction

With the alarming scenario of anthropogenic climate change due to persistently high emission levels of CO<sub>2</sub> from fossil fuels, Carbon Capture and Utilization (CCU) technologies may play a role in the transition of industry towards a net zero future.<sup>1</sup> Moving away from fossil fuels, CO<sub>2</sub> could be implemented as an alternative carbon feedstock in a variety of industrial processes.<sup>2,3</sup> In this context, the production of methanol has been attracting particular interest, given its current role as a critical building block in the chemical industry and the development of sustainable aviation fuel in the near future.<sup>4,5</sup>

Methanol synthesis from CO-rich syngas derived from natural gas using Cu/ZnO/Al<sub>2</sub>O<sub>3</sub> catalysts has been a well-established industrial process over the past decades.

However, replacing fossil resources by alternative carbon sources such as biomass or captured CO<sub>2</sub> will require the development of new processes, among which the direct CO<sub>2</sub> hydrogenation may play a decisive role. In comparison with the traditional CO-based route, inherent limitations of the CO<sub>2</sub> hydrogenation reaction involve its less exothermic character, in addition to the simultaneous production of water.<sup>6</sup> Nevertheless, reports from the first industrial plant devoted to CO<sub>2</sub> hydrogenation to methanol suggest significant advantages, being less energy-intensive and producing fewer reaction byproducts.<sup>7</sup>

Similarly as in the conventional methanol synthesis process, copper-based catalysts have been widely regarded as the most effective materials for CO<sub>2</sub> hydrogenation. Following extensive research, several structural-property relations have been identified for the Cu/ZnO system.<sup>8–10</sup> Although some copper-based catalysts may produce methanol through the Reverse Water-Gas Shift + CO hydrogenation pathway,<sup>11,12</sup> the Cu/ZnO interaction has been strongly associated with CO<sub>2</sub> hydrogenation *via* formate intermediate, according to theoretical studies and operando characterization.<sup>8,13,14</sup>

In addition to such materials, other recent lines of research involve the search for novel catalysts, since the application of copper-based materials is still limited by the instability of the active phase,<sup>15</sup> low selectivity to methanol<sup>16</sup>

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and surface poisoning in the presence of sulfur-containing gases.<sup>17</sup> In virtue of these issues, non-metallic catalysts such as ZnZrOx (ref. 18) and In<sub>2</sub>O<sub>3</sub> (ref. 19) have attracted growing interest as potential alternatives for this application due to their high stability and selectivity towards methanol.

More recently, similarly selective CO<sub>2</sub> hydrogenation to methanol has also been demonstrated using MoS<sub>2</sub> catalysts. In contrast with the aforementioned materials which optimally operate above 250 °C, MoS<sub>2</sub> can be active at remarkably low temperatures around 180 °C.<sup>20</sup> In addition to the more favorable thermodynamics for methanol formation, these mild operation conditions could offer an economical advantage with respect to other catalysts, in the context of large-scale applications. However, CO<sub>2</sub> hydrogenation to methanol can only be achieved in MoS<sub>2</sub> with specific properties, as under the typical reaction conditions most formulations of pure MoS<sub>2</sub> promote the formation of methane instead of methanol. In fact, only in the presence of basal plane S-vacancies CO<sub>2</sub> was shown to dissociate to CO and O at low temperatures, thus allowing for selective methanol production without further hydrogenation to methane.<sup>20</sup> This suggests that in favor of selective CO<sub>2</sub> hydrogenation to methanol, basal-plane sulfur vacancies should be obtained by preparation methods that induce the formation of sheet-like structures instead of edge-rich nanoparticles.

Moreover, in addition to the MoS<sub>2</sub> morphology, methanol selectivity can arise due to other effects. In a recent study, a MoS<sub>2</sub>/ZnS catalyst produced by a solvothermal route involving a metal-organic framework precursor has shown high selectivity to methanol, as a possible result of ZnS blocking MoS<sub>2</sub> edge sites, thereby inhibiting CH<sub>4</sub> production.<sup>21</sup> Therefore, these results suggest that the investigation of synergistic interactions between MoS<sub>2</sub> and other promoter compounds deserves closer attention.

In earlier works, MoS<sub>2</sub> has been combined with K,<sup>22</sup> Co (ref. 23) and Ni (ref. 24) promoters leading to enhanced CO hydrogenation to higher alcohols at pressures around 100 bar. More recently, these promoters were incorporated into the hydrothermal synthesis of MoS<sub>2</sub>, leading to considerable changes in its catalytic activity for CO and CO<sub>2</sub> hydrogenation, as the selectivity can be shifted from CH<sub>4</sub> to CO under lower reaction pressures of 20 bar.<sup>25</sup>

In this work, K and Co promoted-MoS<sub>2</sub> are compared with Mn-promoted MoS<sub>2</sub> produced by a hydrothermal method. Following catalytic testing for CO<sub>2</sub> hydrogenation, Mn-promoted MoS<sub>2</sub> is further characterized by XRD, EXAFS, SEM, EDX, and XPS methods, seeking to investigate the main active phases that drive methanol production in the material.

## Experimental

### Catalyst synthesis

Mn-promoted MoS<sub>2</sub> was produced by a hydrothermal method, in which 2.2 g of ammonium heptamolybdate tetrahydrate ((NH<sub>4</sub>)<sub>6</sub>Mo<sub>7</sub>O<sub>24</sub>·4H<sub>2</sub>O, Carl Roth, 99%) was

dissolved in approximately 20 mL water, together with the appropriate amounts of thiourea (CH<sub>4</sub>N<sub>2</sub>S, Merck, 99%) and manganese sulfate monohydrate (MnSO<sub>4</sub>·H<sub>2</sub>O, Merck, 99%). After stirring for 30 minutes, the mixture is transferred to an autoclave and kept at 200 °C during 16 h. The precipitate was filtered, washed with water and ethanol, dried under vacuum at room temperature and finally calcined under N<sub>2</sub> for 3 h.

For the comparison of pure MoS<sub>2</sub> with different promoters, Mn(0.5)-MoS<sub>2</sub>, whose nomenclature refers to the nominal Mn/Mo molar ratio of 0.5, was produced using a CH<sub>4</sub>N<sub>2</sub>S:MnSO<sub>4</sub>·H<sub>2</sub>O:(NH<sub>4</sub>)<sub>6</sub>Mo<sub>7</sub>O<sub>24</sub>·4H<sub>2</sub>O molar ratio of 32:3.5:1 and a calcination temperature of 500 °C. The analogous synthesis of K(0.5)-MoS<sub>2</sub>, Co(0.5)-MoS<sub>2</sub> and MoS<sub>2</sub> is reported elsewhere.<sup>25</sup> The optimized Mn(0.3)-MoS<sub>2</sub> was produced using a CH<sub>4</sub>N<sub>2</sub>S:MnSO<sub>4</sub>·H<sub>2</sub>O:(NH<sub>4</sub>)<sub>6</sub>Mo<sub>7</sub>O<sub>24</sub>·4H<sub>2</sub>O molar ratio of 24:2.1:1 and a calcination temperature of 400 °C.

### Material characterization

X-ray powder diffraction (XRD) was performed in a PANalytical Empyrean. A Cu-LLF X-ray tube (45 kV, 40 mA, CuKα λ<sub>1</sub> = 1.5406 Å, λ<sub>2</sub> = 1.5444 Å) and a GaliPIX detector were used in Bragg-Brentano geometry. Scanning Electron Microscopy (SEM) was carried out using a FEI Quanta 250 FEG at a 5 kV voltage. Attached to the same device is an Octane Elite Super detector, which was employed for Energy Dispersive X-Ray (EDX) analysis, carried out at 20 kV.

Near-Ambient-Pressure X-ray Photoelectron Spectroscopy (NAP-XPS) was performed with a near ambient-pressure XPS system from SPECS (AlKα source, Phoibos 150 NAP detector) using a sample stage optimized for catalytic measurements.<sup>26</sup> The sample was mounted on a quartz sample carrier with steel backplate and heated with a laser. Pretreatment was carried out at 0.75 mbar H<sub>2</sub> and 400 °C and reaction conditions were simulated under 1 mbar H<sub>2</sub>:CO<sub>2</sub> = 3:1 at 200 °C. *Ex situ* XPS was carried out using a SPECS u-Focus system (AlKα source, Phoibos 150 WAL detector). XPS data evaluation was carried out with CasaXPS software.<sup>27</sup> The peaks were fitted with Gauss-Lorentz (GL) sum functions and a Shirley background was used. All spectra were calibrated for S 2p<sub>3/2</sub> = 162.0 eV.<sup>28,29</sup>

XAS spectra of the Mn and Mo K-edge were collected at the SuperXAS beamline at PSI. It operated in top-up mode at 2.4 GeV and a ring current of 400 mA. A silicon-coated mirror (which also reduced higher-order harmonics) was used to collimate the polychromatic X-rays from a 2.9 T superbend magnet, which were subsequently monochromatized by a Si(111) channel-cut crystal. The monochromator was rocked at a frequency of 1 Hz resulting in two spectra per second. For Mn K-edge a Si-coated and for Mo K-edge a Rh-coated toroidal mirror focused the beam. The Mn, Mo K-edge absorption spectra were collected in transmission mode using 15 cm long ionization chambers filled with 1 bar of N<sub>2</sub> for Mn and 1 bar of N<sub>2</sub> plus 1 bar of Ar for Mo. A metal foil was measured simultaneously for absolute energy calibration.





Standard background subtraction, interpolation, and averaging were done with the Python-based software ProQEXAFS.<sup>30</sup> Normalization and analysis were performed with Demeter.<sup>31</sup> The samples were measured in pellet form diluted by cellulose for optimized signal. In addition to the catalysts, commercial samples of MnO (ABCR, 99%) and MnCO<sub>3</sub> (ABCR, 95%) were analyzed for comparison.

### Catalytic measurements

The catalytic activity was measured on a “micro effi” system of PID Eng&Tech in a fixed bed plug flow steel reactor. For the measurements, 1 g of the pure catalyst was pretreated at 21 bar H<sub>2</sub> at 400 °C. The reaction was conducted for 8 h at each temperature under 20% CO<sub>2</sub>, 60% H<sub>2</sub> and 20% He at 21 bar, with a total flow of 5 mL<sub>n</sub> min<sup>-1</sup>. The product gas was analyzed by an Inficon Micro GC 3000 using a Plot Q column.

## Results and discussion

Seeking to identify potentially promising promoters for MoS<sub>2</sub> catalysts in CO<sub>2</sub> hydrogenation, samples of M-promoted MoS<sub>2</sub> (M = K, Co or Mn) have been produced by analogous hydrothermal approaches. Subsequently, their catalytic activity for CO<sub>2</sub> hydrogenation was compared with pure MoS<sub>2</sub> obtained by the same method, as shown in Fig. 1a–c.

As expected, pure MoS<sub>2</sub> shows high selectivity for CH<sub>4</sub> between 180 and 320 °C, although the activity for CO production through the reverse water-gas-shift reaction becomes prominent above 220 °C. In contrast, the incorporation of K, Co or Mn precursors clearly lowers CH<sub>4</sub> and increases CO yields with respect to the pure material over the entire temperature range. Furthermore, Co–MoS<sub>2</sub>

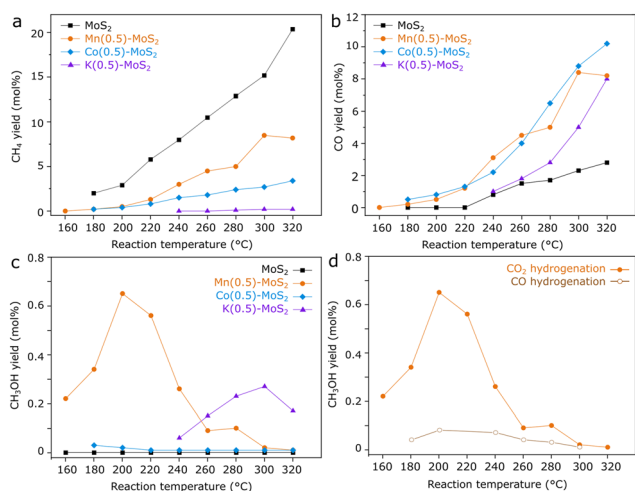
exhibits negligible methanol yield, while in K–MoS<sub>2</sub> some methanol production is observed around 280 °C.

Mn-promoted MoS<sub>2</sub>, however, shows a distinct performance, with the highest methanol yield occurring at much lower temperatures around 200 °C. While pure MoS<sub>2</sub> shows a negligible methanol yield at 180 °C, Mn(0.5)–MoS<sub>2</sub> produces methanol with a selectivity of 45% at a CO<sub>2</sub> conversion of 0.8% under the same conditions. Furthermore, Fig. 1d shows that methanol production is mostly suppressed when CO is alternatively used as the feed gas, demonstrating that the catalyst cannot produce methanol from CO hydrogenation under these conditions. This observation suggests that a reaction mechanism involving CO as an intermediate can be ruled out.

Therefore, it is likely that methanol is produced *via* direct CO<sub>2</sub> hydrogenation *via* formate pathway in the presence of the Mn-promoted MoS<sub>2</sub> catalyst. Although this finding contrasts with the previously demonstrated CO-based pathway observed for pure MoS<sub>2</sub> nanosheets,<sup>20</sup> the coexistence of formate and CO intermediates has been already suggested for the MoS<sub>2</sub>/ZnS catalyst.<sup>21</sup> The divergence in these results support the idea that CO<sub>2</sub> hydrogenation to methanol in MoS<sub>2</sub>-based catalysts may undergo distinct pathways due to structural changes and the addition of promoters to MoS<sub>2</sub>, similarly as observed for copper-based catalysts, for example.<sup>11–14</sup>

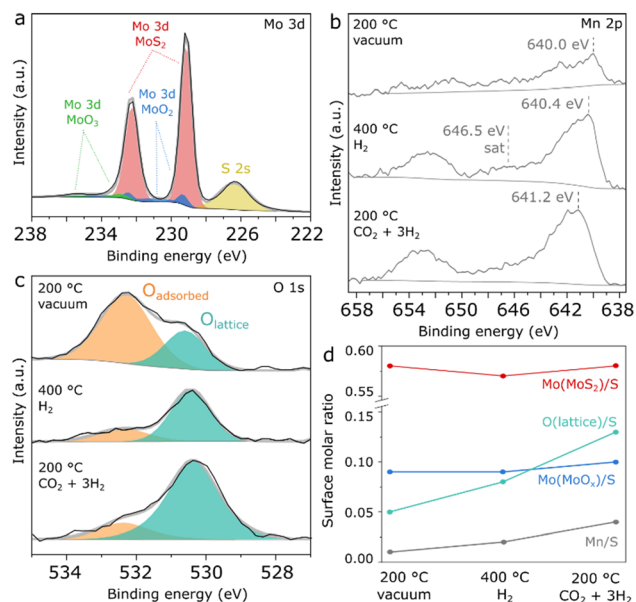
To obtain preliminary insights on the nature of the Mn promoter, characterization of Mn(0.5)–MoS<sub>2</sub> was conducted by XRD and *in situ* NAP-XPS. In Fig. S1,† the XRD pattern of Mn(0.5)–MoS<sub>2</sub> shows that the catalyst crystalline structure consists mostly of MoS<sub>2</sub>,<sup>32</sup> MnO,<sup>33</sup> and MnO<sub>2</sub>.<sup>34</sup> Differently from previously reported K- and Co-promoted MoS<sub>2</sub>,<sup>25</sup> no other sulfides and sulfates are formed in abundance, as MnS (ref. 35) appears with very low crystallinity in the XRD pattern.

In order to obtain more detailed insights into the surface composition of Mn(0.5)–MoS<sub>2</sub> under reaction-relevant conditions, NAP-XPS analysis was carried out firstly under vacuum at 200 °C, followed by H<sub>2</sub> flow at 400 °C and CO<sub>2</sub> + 3H<sub>2</sub> flow at 200 °C. Fig. 2a shows the region comprising Mo 3d and S 2s spectra. A single contribution is observed for sulfur at 226.4 eV, as typically reported for S<sup>2-</sup> in MoS<sub>2</sub>.<sup>28,29</sup> The Mo 3d region, however, shows a more complex profile with three distinct species, which were fitted in accordance with the characteristic doublet separation of 3.14 eV. Coherently with the observation from XRD, the main Mo surface species consist of Mo<sup>4+</sup> associated with MoS<sub>2</sub>, evidenced by the 3d<sub>5/2</sub> peak at 229.1 eV.<sup>28,29</sup> Within a similar energy range, Mo<sup>4+</sup> from MoO<sub>2</sub> is described by two neighboring doublets with different widths, in order to account for the typical asymmetry of MoO<sub>2</sub> peaks arising from the distinctive narrow band metallic character of this compound.<sup>36</sup> Accordingly, this species is fitted with neighboring 3d<sub>5/2</sub> peaks at 229.3 eV and 231.0 eV, and respective 3d<sub>3/2</sub> counterparts following area, width and position constraints reported in a previous study.



**Fig. 1** a) CH<sub>4</sub>, b) CO, c) CH<sub>3</sub>OH yield obtained from catalytic reaction with 1 g MoS<sub>2</sub>, Mn(0.5)-, Co(0.5)-, and K(0.5)-promoted MoS<sub>2</sub> under 1 mL min<sup>-1</sup> CO<sub>2</sub> + 3 mL min<sup>-1</sup> H<sub>2</sub> + 1 mL min<sup>-1</sup> He at 21 bar and d) CH<sub>3</sub>OH yield under CO<sub>2</sub> hydrogenation conditions, compared with an analogous experiment using CO as carbon source.





**Fig. 2** NAP-XPS spectra of Mn(0.5)-MoS<sub>2</sub> showing the regions a) Mo 3d under vacuum at 200 °C, b) Mn 2p and c) O 1s under vacuum at 200 °C, H<sub>2</sub> at 400 °C and CO<sub>2</sub> + 3H<sub>2</sub> at 200 °C and d) summarized quantification of surface molar ratios with respect to S.

Furthermore, a minor contribution from another doublet is verified, with a 3d<sub>5/2</sub> peak at 232.9 eV as a possible result of surface Mo<sup>6+</sup> oxides.<sup>36,37</sup> Correspondingly, MoS<sub>2</sub> produced by an analogous hydrothermal approach without the Mn promoter also presents similar evidences for surface Mo oxides, as seen in Fig. S2.† Since this pure MoS<sub>2</sub> sample has shown negligible methanol selectivity, this finding suggests that MoO<sub>x</sub> should not be responsible for the promoting effect.

As shown in Fig. S3,† the chemical environment of surface Mo and S experiences insignificant changes upon H<sub>2</sub> pretreatment and reaction conditions. In Fig. 2b, the Mn 2p region initially shows a doublet with very low intensity, which suggests a low surface concentration of the Mn promoter. However, under H<sub>2</sub> treatment at 400 °C, the Mn 2p doublet becomes more prominent, exhibiting a 2p<sub>3/2</sub> maximum at 640.4 eV and a satellite feature around 646.5 eV, as possible indicators of Mn<sup>2+</sup> in MnO.<sup>38</sup> Subsequently, under CO<sub>2</sub> + 3H<sub>2</sub> flow at 200 °C the Mn 2p doublet is shifted towards higher binding energy by almost 1 eV, suggesting the formation of higher oxidation states such as Mn<sup>3+</sup> and Mn<sup>4+</sup> at the surface under reaction conditions. However, precise determination and quantification of these phases is challenging in a spectrum with such low intensity due to the complex and asymmetric character of the components associated with manganese oxides.<sup>38</sup>

In Fig. 2c, the O 1s spectrum shows surface oxygen described by two species: the component at approximately 530.6 eV can be associated with lattice oxygen from Mo or Mn oxides, while the one at 532.3 eV typically refers to surface hydroxyl or organic species.<sup>39</sup> Although initially the lattice oxygen component is smaller, it becomes much more

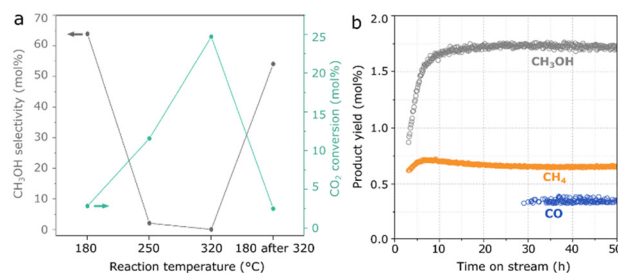
prominent than the adsorbed species under pretreatment and reaction conditions.

As a summary of the NAP-XPS analysis, Fig. 2d shows a quantification based on combined survey spectra and high-resolution Mo 3d and O 1s regions. While the components related to MoS<sub>2</sub> and MoO<sub>x</sub> show negligible changes, the Mn/S surface molar ratio increases concurrently with the O<sub>lattice</sub>/S ratio during H<sub>2</sub> pretreatment and reaction conditions, suggesting the formation of surface Mn oxides at the surface under reaction-relevant conditions. Interestingly, Mn<sup>2+</sup> appears to coexist with higher Mn oxidation states despite the reducing environment created by H<sub>2</sub> and CO<sub>2</sub>, as a possible consequence of oxygen transferred from bulk to surface or due to the dissociation of CO<sub>2</sub> on MnO. Although precise identification of this surface oxidation mechanism is challenging, NAP-XPS analysis suggests that Mn oxides are the key Mn-containing phase under reaction conditions.

Given the promising catalytic activity of Mn-promoted MoS<sub>2</sub>, the hydrothermal synthesis was further optimized in terms of calcination temperature, content of thiourea and content of Mn precursor. As shown in Fig. S4,† the methanol yield is improved by lowering calcination temperature from 500 °C to 400 °C, the MnSO<sub>4</sub>·H<sub>2</sub>O:(NH<sub>4</sub>)<sub>6</sub>Mo<sub>7</sub>O<sub>24</sub>·4H<sub>2</sub>O molar ratio from 3.5:1 to 2.1:1 and CH<sub>4</sub>N<sub>2</sub>S:(NH<sub>4</sub>)<sub>6</sub>Mo<sub>7</sub>O<sub>24</sub>·4H<sub>2</sub>O from 32:1 to 24:1.

In Fig. 3a, as a result of the optimized hydrothermal method, Mn(0.3)-MoS<sub>2</sub> shows a methanol selectivity of 64% with an improved CO<sub>2</sub> conversion of 2.8% at 180 °C. At higher temperatures, the formation of CO and CH<sub>4</sub> becomes dominant, although further cooling to 180 °C does not lead to an expressive decrease in methanol yield and selectivity. Furthermore, the material also exhibits stable catalytic activity at 180 °C during a 50-hour experiment, as shown in Fig. 3b.

To identify the main Mn-containing phases in the optimized catalyst, XRD and EXAFS have been performed in Mn(0.3)-MoS<sub>2</sub> catalyst, following exposure to relevant reaction conditions. In Fig. 4, as-synthesized Mn(0.3)-MoS<sub>2</sub> presents a variety of phases. As expected, the main contribution consists of MoS<sub>2</sub> (ref. 32) with low crystallinity,



**Fig. 3** Methanol selectivity and total CO<sub>2</sub> conversion as a result of a) 8 h catalytic reaction at 180 °C, 250 °C, 320 °C and 180 °C after 320 °C with 1 g Mn(0.3)-MoS<sub>2</sub> under 1 mL min<sup>-1</sup> CO<sub>2</sub> + 3 mL min<sup>-1</sup> H<sub>2</sub> + 1 mL min<sup>-1</sup> He at 21 bar and b) CH<sub>3</sub>OH, CH<sub>4</sub> and CO yields under the same conditions during a 50-hour reaction.

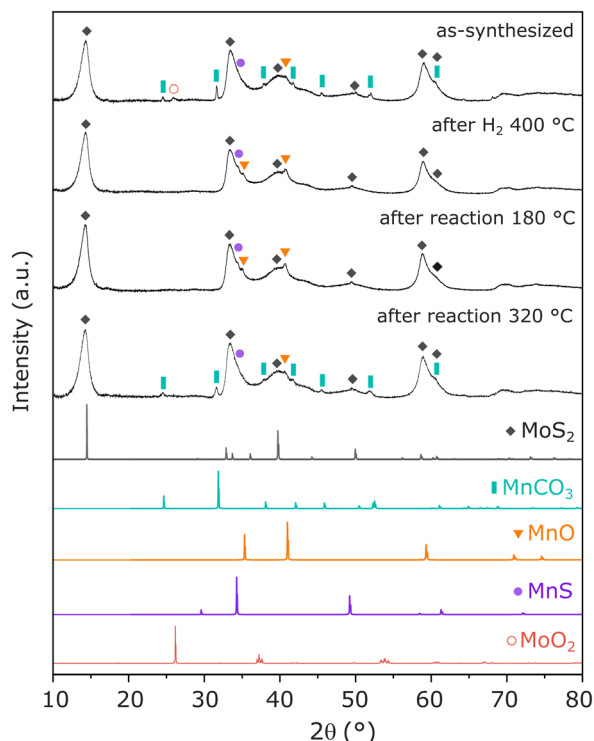


Fig. 4 XRD pattern of Mn(0.3)-MoS<sub>2</sub> following synthesis, H<sub>2</sub> pretreatment at 400 °C and catalytic reaction at 180 °C and 320 °C, in comparison with reference data for MoS<sub>2</sub> (COD-ID 1010993), MnCO<sub>3</sub> (COD-ID 1011228), MnO (COD-ID 1514099), MnS (COD-ID 1011351) and MoO<sub>2</sub> (COD-ID 9009090).

which is a typical outcome from the hydrothermal method. The promoter Mn is mostly present in the form of MnCO<sub>3</sub> (ref. 40) as a result of the reaction between the Mn precursor and thiourea during the hydrothermal synthesis. Additionally, the diffractogram suggests that minor contributions from MoO<sub>2</sub> (ref. 32) and nearly amorphous MnO (ref. 33) and MnS (ref. 35) are also present in the material.

The pretreatment under H<sub>2</sub> at 400 °C significantly alters the catalyst structure: although the patterns related to MoS<sub>2</sub> and MnS do not experience significant changes, MnCO<sub>3</sub> is absent in the pretreated material while a noticeable profile related to MnO (ref. 33) arises. This observation indicates that the pretreatment converts MnCO<sub>3</sub> into MnO, in an analogous manner as previously reported under H<sub>2</sub> at similar temperatures.<sup>41,42</sup> Furthermore, the feature related to MoO<sub>2</sub> disappears, possibly indicating amorphization under reducing conditions. All these features are maintained after the material undergoes catalytic reaction at 180 °C, demonstrated to be the optimal condition for CO<sub>2</sub> hydrogenation to methanol.

Moreover, after reaction at 320 °C, the material exhibits once again the pattern from MnCO<sub>3</sub>, indicating that the higher temperature favors the carbonation reaction of MnO in the presence of the CO<sub>2</sub> + 3H<sub>2</sub> mixture at 21 bar. This phase transformation may be closely associated with the

slight catalyst deactivation after reaction at 320 °C, already shown in Fig. 3a.

Given the observation of MnO following catalytic reaction at the ideal conditions for methanol production, this phase can be pointed out as a likely key feature behind the promoting effect of Mn. Nevertheless, due to the possible effect of amorphous Mn-containing species that could remain undetected by XRD, such as other oxides, sulfides or Mn intercalated within MoS<sub>2</sub> layers, EXAFS analysis was carried out in order to elucidate the coordination environment of Mn atoms in the catalyst.

In Fig. 5a, XANES spectra of the Mo edge show negligible differences between the samples. Fig. 5b and S5† show that the Fourier transform of the Mo K-edge exhibits mainly two features related to Mo-S and Mo-Mo, coherently with MoS<sub>2</sub>.<sup>43</sup> As suggested in the EXAFS fitting in Fig. S5 and Table S1,† the higher prominence of the Mo-S coordination with respect to Mo-Mo may be associated with the low crystallinity of MoS<sub>2</sub>, in line with the XRD patterns and previous EXAFS reports.<sup>44,45</sup> Despite the differences observed for Mn, the Mo K-edges remain unchanged regardless of exposure to pretreatment and reaction conditions. Therefore, the result confirms the high stability of MoS<sub>2</sub> following reaction conditions and rules out major contributions from Mo oxides to the catalyst composition.

As shown in Fig. 5c, XANES spectra from the Mn K-edge region of Mn(0.3)MoS<sub>2</sub> give evidence for considerable changes between as-synthesized and spent catalysts. In Fig. 5d, the corresponding Fourier transforms show 2 main features related to Mn-O and Mn-Mn coordination shells.<sup>46,48</sup> Consistently with the presence of MnO observed by XRD after treatment with H<sub>2</sub> at 400 °C and reaction at 180

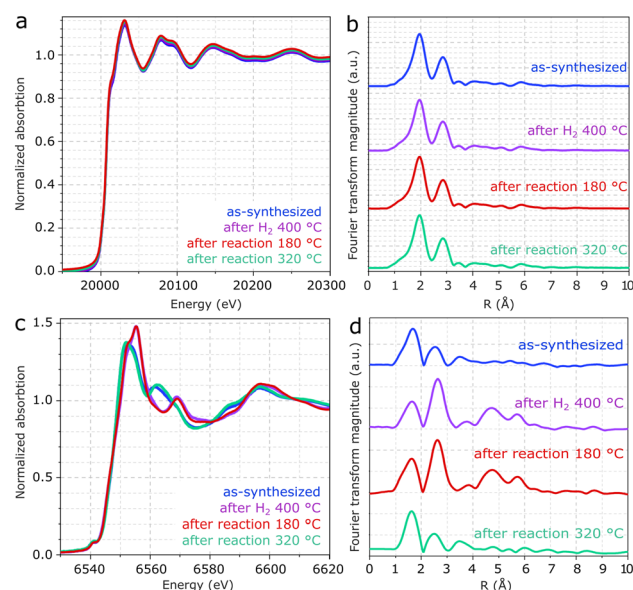


Fig. 5 a and b) XANES spectra and the associated EXAFS analysis of the Mo and c and d) Mn K-edges from Mn(0.3)-MoS<sub>2</sub> after synthesis, H<sub>2</sub> pretreatment at 400 °C and catalytic reaction at 180 °C and 320 °C.





°C, the EXAFS spectrum clearly shows the characteristic profile associated with MnO (ref. 47 and 48) in these two samples, as confirmed by the fitting presented in Fig. S6 and Table S2.<sup>†</sup><sup>43,45</sup>

On the other hand, as-synthesized Mn(0.3)MoS<sub>2</sub> exhibits a similar XANES spectrum from the one observed after 320 °C reaction, as shown in Fig. 5c. Despite some similarity with the typical XANES profile observed in MnCO<sub>3</sub>,<sup>49</sup> the respective EXAFS data shows considerable differences when compared to a MnCO<sub>3</sub> reference in Fig. S7,<sup>†</sup> which raises the possibility of a more complex composition. In fact, Fig. 6 shows that these XANES spectra may be described as a linear combination of MnCO<sub>3</sub> and MnO. A clear indication of this similarity is the distinctive feature of MnO around 6569 eV, which makes the XANES spectra easily distinguishable from those of MnS and other Mn oxides.<sup>50</sup> In view of this finding, as-synthesized Mn(0.3)MoS<sub>2</sub> is considered to present the Mn promoter in the form of polycrystalline MnCO<sub>3</sub> in combination with low-crystallinity MnO, similarly as observed for the material exposed to reaction at 320 °C, since both samples show a weak contribution of the oxide in the XRD patterns in Fig. 4.

Material characterization by XRD and XAS strongly suggests that MoS<sub>2</sub> and MnO are the key components related to CO<sub>2</sub> hydrogenation to methanol in this catalyst. In order to unveil how these phases are present at the catalyst surface, further characterization was performed by SEM and XPS.

Fig. 7a and S8<sup>†</sup> show that the Mn(0.3)MoS<sub>2</sub> surface is mostly composed of thin MoS<sub>2</sub> sheets arranged in a nanoflower morphology, highly similar to MoS<sub>2</sub> produced by an analogous hydrothermal synthesis. This aspect is consistent with the broad XRD pattern observed for MoS<sub>2</sub>, which suggests small crystallites and sparse stacking of MoS<sub>2</sub>

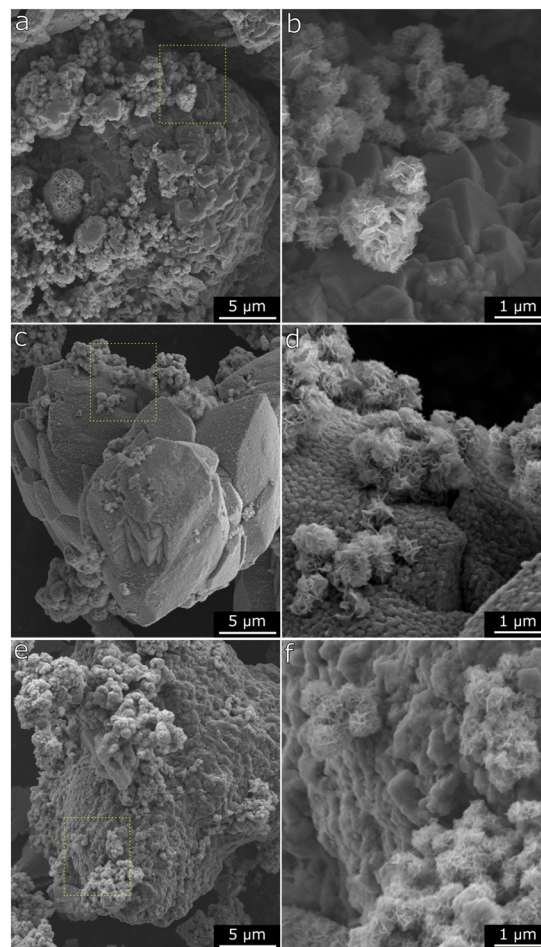


Fig. 7 a and b) SEM micrographs of Mn(0.3)-MoS<sub>2</sub> after synthesis, c and d) after H<sub>2</sub> pretreatment at 400 °C and reaction at 180 °C, e and f) after H<sub>2</sub> pretreatment at 400 °C and reaction at 320 °C.

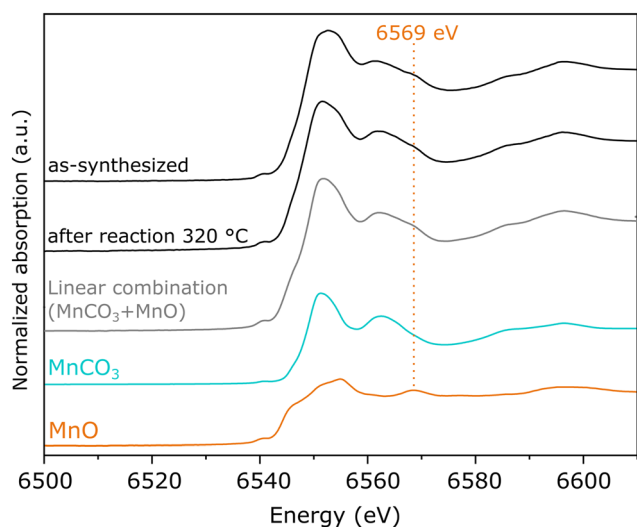
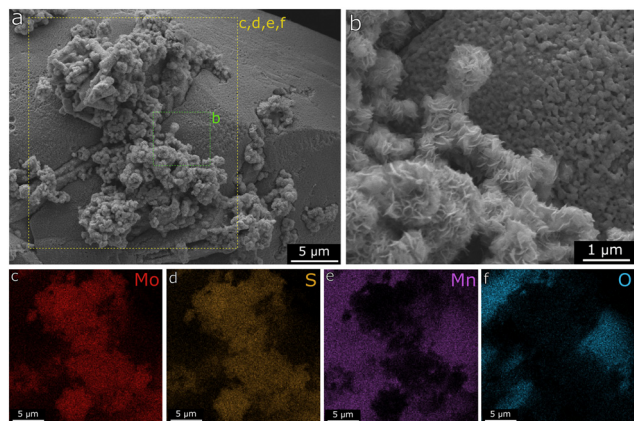


Fig. 6 XANES spectra of the Mn K-edge from Mn(0.3)-MoS<sub>2</sub> after synthesis and catalytic reaction at 320 °C, compared to MnO and MnCO<sub>3</sub> references samples and their linear combination considering equal proportions of each phase.

layers. Moreover, these features coexist with another evident morphology consisting of the micrometer-sized particles shown in more detail in Fig. 6b. After H<sub>2</sub> pretreatment and subsequent catalytic reaction at 180 °C, the nanoflower structure associated with MoS<sub>2</sub> remains unchanged, as shown in Fig. 6c and d. On the other hand, the larger particles are significantly altered, now featuring a rougher surface with abundant pores in the nanometer range. This finding is consistent with the conversion of MnCO<sub>3</sub> into MnO, which releases CO and CO<sub>2</sub>, thus forming the characteristic porous surface. Such morphology change has also been already reported following exposure of MnCO<sub>3</sub> to reducing conditions under similar temperatures.<sup>41,42</sup> As demonstrated by XRD and EXAFS analysis of Mn(0.3)MoS<sub>2</sub>, further increasing reaction temperature to 320 °C induces the partial carbonation of MnO into MnCO<sub>3</sub>. This effect can also be correlated with the SEM data in Fig. 6e and f, where the characteristic porous surface of MnO is again less prominent.

Further evidence for the MoS<sub>2</sub>/MnO system after H<sub>2</sub> pretreatment is shown in Fig. 8, in which the EDX mapping confirms the elemental composition of the nanosheets as





**Fig. 8** a and b) SEM micrograph with the respective EDX element mapping for c) Mo, d) S, e) Mn and f) O atoms for Mn(0.3)-MoS<sub>2</sub>, after H<sub>2</sub> pretreatment at 400 °C.

mostly Mo and S, while the larger particles are rich in Mn and O. A summary of the EDX spectra contained in the evaluated region is displayed in Fig. S9.†

Finally, further insights into the surface composition of Mn(0.3)-MoS<sub>2</sub> are provided by XPS analysis of fresh and used catalysts. In Fig. 9a, the S 2p spectrum shows a unique doublet with the characteristic splitting of 1.2 eV and S 2p<sub>3/2</sub> located at 162.0 eV, as typically reported for sulfides such as MoS<sub>2</sub>. Similarly as observed in the NAP-XPS analysis of Mn(0.5)-MoS<sub>2</sub>, Fig. 9b indicates that MoS<sub>2</sub> (Mo 3d<sub>5/2</sub> at 229.1 eV)<sup>28,29</sup> coexists with surface MoO<sub>2</sub> (Mo 3d<sub>5/2</sub> at 229.3 eV and 231.0 eV) as well as MoO<sub>3</sub> (Mo 3d<sub>5/2</sub> at 232.9 eV),<sup>36,37</sup> although MoS<sub>2</sub> brings again the most prominent contribution.

Accordingly, Fig. 9c shows that the O 1s spectrum indicates two distinct contributions: the minor peak at 532.3 eV can be ascribed to adsorbed oxygen from water or surface hydroxyl species, while the most prominent is located at 530.6 eV, associated with Mo and Mn oxides.<sup>39</sup> The low

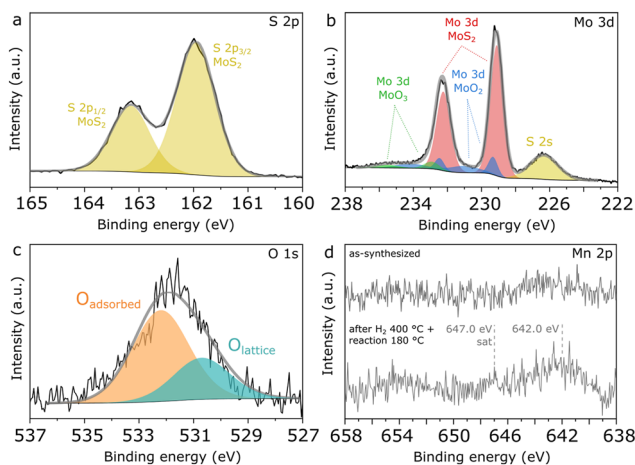
intensity of the peak related to lattice oxygen can be correlated with the minor contributions from oxides in the Mo 3d and Mn 2p spectra, thus confirming that metal oxides consist only of a minor contribution to the total surface composition, with MoS<sub>2</sub> being the dominant species.

In order to verify possible modifications at the surface of Mn(0.3)MoS<sub>2</sub>, XPS analysis was also conducted after H<sub>2</sub> pretreatment at 400 °C and reaction at 180 °C. As shown in Fig. S10,† these conditions do not promote expressive changes in S 2p, Mo 3d and O 1s regions, suggesting that the surface remains rich in stable MoS<sub>2</sub>, in line with the strong similarities in the nanosheet morphology observed by SEM both before and after reaction. On the other hand, Fig. 9d shows that a significant change is observed in the Mn 2p region, as the doublet is only visible in the spent catalyst. This effect may be associated with the phase transition from MnCO<sub>3</sub> to MnO experienced by the catalyst upon H<sub>2</sub> pretreatment, as the higher surface area of the porous MnO could enhance the photoelectron signal related to the Mn species. Even after pretreatment and reaction, the Mn/Mo surface atomic ratio is still approximately 0.1, much lower than the nominal value of 0.3, as a possible outcome of the extensive covering of Mn phases by MoS<sub>2</sub> sheets demonstrated by the SEM data.

Furthermore, in the spent catalyst the Mn 2p spectrum is consistent with concurrent MnO and MnO<sub>x</sub> with higher oxidation states, given the combination of a faint satellite feature related to MnO near 647.0 eV and the 2p<sub>3/2</sub> peak located around 242.0 eV.<sup>38</sup> Since XRD and EXAFS characterization give evidence of MnO as the only Mn oxide phase in the material, these oxidized MnO<sub>x</sub> species are understood to be limited to the catalyst surface.

In previous research, S-vacancies in MoS<sub>2</sub> have been strongly associated with its catalytic activity, and their presence at the surface can be usually verified by XPS. According to quantification based on Mo 3d and S 2s regions shown in Fig. 9b, an approximate S/Mo ratio of 1.7 is calculated considering only the Mo component associated with MoS<sub>2</sub>. This low value may be associated with abundant sulfur vacancies formed during calcination under N<sub>2</sub>, as it does not change significantly after H<sub>2</sub> pretreatment and reaction.

Moreover, given that abundant basal plane sulfur vacancies have been strongly suggested as active sites for CO<sub>2</sub> hydrogenation to methanol,<sup>20</sup> an oxygen chemisorption experiment coupled with *in situ* DRIFTS has been performed in an attempt to distinguish between edge- and basal plane sulfur vacancies. However, as shown in Fig. S11,† the results indicate no visible changes in the vibrational spectrum before and after O<sub>2</sub> flow. Although this suggests a limited concentration of basal plane sulfur vacancies in the catalyst with respect to previous reports, the presence of other surface metal oxides such as MoO<sub>x</sub> and MnO<sub>x</sub> introduces overlapping vibrational bands that may hinder the detection of the typical features related to oxygen chemisorption on sulfur vacancies.



**Fig. 9** High-resolution XPS spectra of Mn(0.3)-MoS<sub>2</sub> showing a) S 2p, b) Mo 3d, c) O 1s and d) Mn 2p regions.





In summary, characterization of Mn(0.3)-MoS<sub>2</sub> upon hydrothermal synthesis indicates that the Mn promoter is initially present as a combination of MnCO<sub>3</sub> and low-crystallinity MnO. Subsequently, H<sub>2</sub> pretreatment fully converts the carbonate into MnO, although higher oxidation states for Mn may be present at the surface. This MoS<sub>2</sub>/MnO<sub>x</sub> character is maintained after reaction at 180 °C. Even though increasing reaction temperature to 320 °C leads to partial carbonation of MnO, only a mild catalyst deactivation is observed.

Despite the correlation of MoS<sub>2</sub>/MnO<sub>x</sub> with catalytic activity, it is challenging to evaluate if MnCO<sub>3</sub> plays any role in catalytic activity, as here this phase always coexists with MnO, as demonstrated by XANES analysis. In light of these findings, CO<sub>2</sub> hydrogenation to methanol may be associated with the synergy between MoS<sub>2</sub> and MnO<sub>x</sub>, while pure MoS<sub>2</sub> obtained by an analogous synthesis approach shows negligible selectivity for methanol.

In view of recent findings, MoS<sub>2</sub> growth in the vicinity of MnO<sub>x</sub> might explain the improved methanol selectivity, since edge-blocking effects could inhibit the production of CH<sub>4</sub> in MoS<sub>2</sub> edges.<sup>21</sup> Therefore, further studies with simpler MoS<sub>2</sub>/MnO<sub>x</sub> systems and detailed characterization of S-vacancies will be important for understanding the interplay between MoS<sub>2</sub> and MnO<sub>x</sub> during CO<sub>2</sub> hydrogenation to methanol.

In comparison with other MoS<sub>2</sub>-based materials presented in Table S3† such as few-layer MoS<sub>2</sub>,<sup>20</sup> MoS<sub>2</sub>/ZnS (ref. 21) and Cu/MoS<sub>2</sub>@SiO<sub>2</sub>,<sup>51</sup> the Mn(0.3)-MoS<sub>2</sub> catalyst has a moderate methanol selectivity at lower or comparable CO<sub>2</sub> conversion levels. Accordingly, as shown in the SEM results, MoS<sub>2</sub> is not well dispersed with the Mn promoter phase, which would explain the notable production of the CH<sub>4</sub> byproduct related to pure MoS<sub>2</sub>. Therefore, this suggests MoS<sub>2</sub>/MnO<sub>x</sub> catalysts could be further improved by employing synthesis methods that enhance the dispersion of these phases and their surface area.

## Conclusions

In summary, this work demonstrates that Mn-promoted MoS<sub>2</sub> presents promising properties as a catalyst for CO<sub>2</sub> hydrogenation to methanol. Even though its catalytic activity is lower than in some other MoS<sub>2</sub>-based catalysts, a sharp increase in methanol selectivity is observed in comparison with pure MoS<sub>2</sub> obtained by an analogous hydrothermal synthesis method. This improvement suggests a promoting effect of Mn, which may be closely associated with the presence of Mn oxides, according to material characterization. More specifically, the optimized catalyst contains MnO as the main Mn phase, although surface characterization indicates that Mn<sup>2+</sup> coexists with higher oxidation states under reaction-relevant conditions. This finding suggests that the limited selectivity to methanol

could be further improved in catalysts with abundant MoS<sub>2</sub>/MnO<sub>x</sub> interfaces.

Furthermore, given the importance of basal plane S-vacancies in MoS<sub>2</sub> catalysts for CO<sub>2</sub> hydrogenation to methanol,<sup>20</sup> it can be speculated that MoS<sub>2</sub>/MnO<sub>x</sub> interfaces facilitate this reaction in Mn-promoted MoS<sub>2</sub>, possibly due the blockage of MoS<sub>2</sub> edge sites, as previously observed for a MoS<sub>2</sub>/ZnS system.<sup>21</sup> Therefore, further characterization focused on S-vacancies is necessary for a deeper understanding of such edge-blocking mechanism between MoS<sub>2</sub> and metal-oxides. Moreover, the evidence for direct CO<sub>2</sub> hydrogenation without a CO intermediate in Mn-promoted MoS<sub>2</sub>, differently as in pure MoS<sub>2</sub> nanosheets,<sup>20</sup> emphasizes the importance of investigating reaction mechanisms with more detail in future work.

## Author contributions

Conceptualization: G. P., K. F. and G. A. S. A. Methodology: G. P. and K. F. Formal analysis: G. A. S. A., G. P. and S. P. Investigation: G. A. S. A., G. P., S. P., T. W., C. R. and R. R. Writing – original draft preparation: G. A. S. A. and G. P. Writing – review & editing: G. A. S. A. and K. F. Supervision: K. F. Funding acquisition: K. F.

## Conflicts of interest

There are no conflicts to declare.

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