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# Structural dynamics of an iron molybdate catalyst under redox cycling conditions studied with *in situ* multi edge XAS and XRD†

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The structural dynamics and phase transformations of an iron molybdate catalyst with excess molybdenum trioxide (Mo/Fe = 2.0) were studied during redox cycling of the catalyst using *in situ* multi-edge X-ray absorption spectroscopy (XAS) at the Mo K-edge (transmission mode) and Fe K-edge (fluorescence mode) in combination with X-ray diffraction (XRD). X-ray absorption near edge structure (XANES) and extended X-ray absorption fine structure (EXAFS) analysis showed that heating under reducing conditions with methanol up to 400 °C produced MoO<sub>2</sub> and FeMoO<sub>4</sub>. Linear combination fitting (LCF) analysis showed that iron was reduced completely, while molybdenum remained partly oxidized (60% as Mo(VI)). Complementary *in situ* XRD also supported the phase transformation due to reduction of Fe<sub>2</sub>(MoO<sub>4</sub>)<sub>3</sub> and MoO<sub>3</sub> to FeMoO<sub>4</sub> and MoO<sub>2</sub>. Subsequent heating under oxidative conditions from 200 to 500 °C transformed the catalyst into its initial state via Fe<sub>2</sub>O<sub>3</sub> and extra MoO<sub>3</sub> as intermediate phases. This underlines the segregation and iron enrichment during redox cycling. MoO<sub>3</sub> volatilization, observed under industrial reaction conditions of a methanol and oxygen containing atmosphere, causes this segregation to be permanent. Complete regeneration could only be achieved at 500 °C, which is significantly higher than industrial reaction temperatures. Overall, multi edge *in situ* XAS along with complementary XRD was found to be an ideal tool for tracing the different amorphous and crystalline phases present during redox cycling of the catalyst.

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## 1. Introduction

Iron molybdate with an excess of molybdenum oxide (denoted FeMo throughout the paper) is an industrially important

catalyst for the selective oxidation of methanol to formaldehyde, which has led to a number of fundamental studies concerning its structure, activity, mechanism, stability and deactivation.<sup>1–6</sup> To improve the performance of a catalyst on a knowledge-based approach, it is critical to understand the structure and structural dynamics of the catalyst under reaction conditions. Commercial iron molybdate catalysts are typically formulated with an excess of molybdenum(VI)oxide, since Mo is known to volatilize from the reactor inlet and hot spot<sup>7</sup> during the reaction, leaving an unselective Fe rich surface.<sup>8</sup> The FeMo system contains multiple distinct phases including bulk  $\alpha$ -MoO<sub>3</sub> and Fe<sub>2</sub>(MoO<sub>4</sub>)<sub>3</sub>, as well as a surface layer of MoO<sub>x</sub> on the Fe<sub>2</sub>(MoO<sub>4</sub>)<sub>3</sub> surface.<sup>9</sup> Hence, it is difficult to assess the nature of the active sites and role of Fe<sub>2</sub>(MoO<sub>4</sub>)<sub>3</sub>, specifically whether it merely acts as a support for the active MoO<sub>x</sub> or whether it has its own inherent activity. The catalytic behavior of FeMo catalysts was observed to be mainly dependent on their Mo/Fe atomic ratio.<sup>2,10</sup> The most active phase composition of the bulk iron molybdate catalyst was proposed to have the stoichiometric Mo/Fe ratio of 1.5<sup>3</sup> and the function of excess MoO<sub>3</sub> is speculated to maintain the active phase of the catalyst

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† Electronic supplementary information (ESI) available: Overview of the experimental procedure, XANES spectrum of the initial state of the FeMo catalyst and the structural parameters obtained from EXAFS fitting, EXAFS fitting curves at the Mo K-edge for the FeMo catalyst during different stages of redox cycling, structural parameters obtained from EXAFS fitting for intermediate phases during TPR and TPO, PCA and LCF results at the Fe K-edge during TPR, MS traces of MeOH, HCHO and H<sub>2</sub>O during TPR and isothermal period, *in situ* XANES spectra at the Mo K- and Fe K-edge and XRD during cooling down to 200 °C after TPR and switching to oxidizing conditions, complete LCF results at the Mo K-edge during TPO and PCA results at the Fe K-edge during TPO. See DOI: 10.1039/d0cp01506g



at this ratio. The activity and selectivity of bulk iron molybdate catalysts for methanol oxidation was examined as a function of Mo/Fe ratios from 0.02 to 4,<sup>2</sup> revealing that Mo/Fe ratios greater than 1.5 are required for high catalytic activity and selectivity to formaldehyde. The presence of an octahedral Mo oxide overlayer has been reported,<sup>1</sup> which corresponds to one monolayer. Various groups have proposed that this overlayer structure is the fundamental, active surface for the selective oxidation of methanol to formaldehyde.<sup>1,9,11</sup> The presence of two terminal oxygen atoms double bonded to Mo in such coordination allows the reacting methanol molecules to be bonded simultaneously at two points to the surface. The formation of several molybdenum oxy-methoxy compounds was also reported<sup>12</sup> and it was argued that hydrogen abstraction of the methanol hydroxyl group produces methoxy species that are intermediates in formaldehyde formation. The role of Fe in iron molybdate catalysts would be to favor the transfer of O<sub>2</sub> and H<sub>2</sub>O between the surface and the gas phase and to facilitate re-oxidation of reduced Mo.<sup>11</sup> Furthermore, oxygen atoms from terminal Mo=O groups were argued to be crucial for the activity and selectivity of MoO<sub>3</sub>.<sup>13–16</sup> Several samples revealed a co-existence of various MoO<sub>x</sub> phases and similar qualitative phase compositions existed for different Mo/Fe ratios which does not allow to draw any conclusions about a correlation between a certain MoO<sub>3</sub> phase and its catalytic behavior.

In order to predict and model the performance of catalysts, their electronic and structural properties need to be studied at an atomic scale as they strongly influence the catalytic properties. Under dynamic conditions (temperature, pressure, gas atmosphere, *etc.*) catalysts can be chemically transformed, giving rise to changes in their initial surface composition correlated to bulk structure, morphology and particle size. Thus, complementary techniques have been used for detailed studies of the structural, electronic and chemical properties of different types of catalysts under these conditions. The combination of X-ray absorption spectroscopy (XAS) with complementary techniques can provide *in situ* and *operando* chemical, structural, and electronic insights into the catalysts and other involved species to holistically understand the catalytic process. XAS coupled with X-ray diffraction (XRD) is beneficial since the nature of crystallite phases can be determined by XRD whereas XAS is useful for determining the oxidation state and coordination geometry of the metal atoms. In order to evaluate the potential of the multiple-technique approach and usefulness of different *in situ/operando* reactors at the synchrotron facilities, the behavior of an Fe<sub>2</sub>(MoO<sub>4</sub>)<sub>3</sub> catalyst under redox conditions was discussed in the literature.<sup>17–19</sup> During reduction in MeOH, due to continuous removal of oxygen, pseudo-molybdate and a MoC type phase were observed, leading to the breakdown of the catalyst to separate oxides (low Mo/Fe ratio of 1.5).<sup>17</sup> The rate of re-oxidation was found to increase with increasing Mo concentration. Along with other observations, this clarified the role of MoO<sub>3</sub> excess in the as-prepared catalyst, which is to maintain the stability of the active phase and to replenish any evaporated molybdenum.<sup>18</sup>

In a recent study<sup>20</sup> on the same catalytic system, we have shown that during its evolution with time on stream for

selective oxidation of methanol to formaldehyde, the FeMo catalyst undergoes strong structural changes due to evaporation of MoO<sub>3</sub>. In this study we now probe the structural changes of Mo and Fe related crystalline and amorphous phases during redox cycling using *in situ* multi-edge XAS at the Mo K- and Fe K-edges combined with complementary XRD. Using redox cycling in methanol and under oxidizing conditions insight into the structural dynamics can be received. The combination of multi-edge XAS and XRD allows for investigating the effect of over-reduction followed by re-oxidation of the catalyst, corresponding to one cycle of the selective oxidation of methanol to formaldehyde. For this purpose we exploited the advanced beamline set-ups available at the synchrotron radiation sources which allow obtaining high quality bulk structural data by XAS and XRD from catalysts under dynamic conditions when simultaneously using multiple techniques.

## 2. Experimental

The catalyst was prepared by hydrothermal synthesis with special interest paid to the precise adjustment of the atomic molybdenum to iron ratio.<sup>7</sup> The Mo/Fe ratio was 2.0, which resulted in an active and selective catalyst in a broad temperature range, typically 250–400 °C, at a feed concentration of 5 vol% CH<sub>3</sub>OH with about 10 vol% O<sub>2</sub>.<sup>7</sup> The catalyst was calcined at 535 °C in air, resulting in a BET surface area of 4.7 m<sup>2</sup> g<sup>−1</sup> and Rietveld refined particles size of 133 nm for the MoO<sub>3</sub> phase and 375 nm for the Fe<sub>2</sub>(MoO<sub>4</sub>)<sub>3</sub> phase. The catalyst was crushed and ground with silica in a 1/5 catalyst/silica weight ratio. This mixture was pressed as a pellet, crushed and sieved to 100–150 μm. Combined XAS/XRD studies during temperature programmed experiments were conducted at the SNBL beamline BM31 at ESRF, Grenoble, France.<sup>21</sup> Fig. 1 shows the schematics of the advance experimental setup used for the *in situ* multi-edge XAS/XRD experiments. As optics requirements for high quality XAS and XRD are quite different, the beamline is equipped with a channel-cut monochromator (CCM) for a high-resolution powder diffraction and a second double crystal monochromator (DCM) for XAS (interchanged within 10 s).<sup>21</sup> Measurements were performed by means of a fixed-bed micro-reactor based on quartz glass capillaries<sup>22</sup> for mimicking optimal heat and mass transfer.<sup>23,24</sup> The catalyst in the form of powder was loaded in a glass capillary with inner diameter of 1.5 mm with the total bed length inside the capillary of 1 cm and the catalyst was fixed from both sides by quartz wool. XAS was performed at the Fe K-edge (7.112 keV) in fluorescence mode and at the Mo K-edge (20.000 keV) in transmission mode quasi-simultaneously by using the fast edge changing facility at the beamline. A molybdenum foil was measured simultaneously as a reference compound for energy calibration. The measurements were conducted in a loop that consisted of two Mo K-edge scans (4 min each), three Fe K-edge scans (5 min each) and one XRD pattern (1 min). Switching between Fe K- and Mo K-energies took approximately 30 s and switching from XAS to XRD monochromator and detector took 60 s.



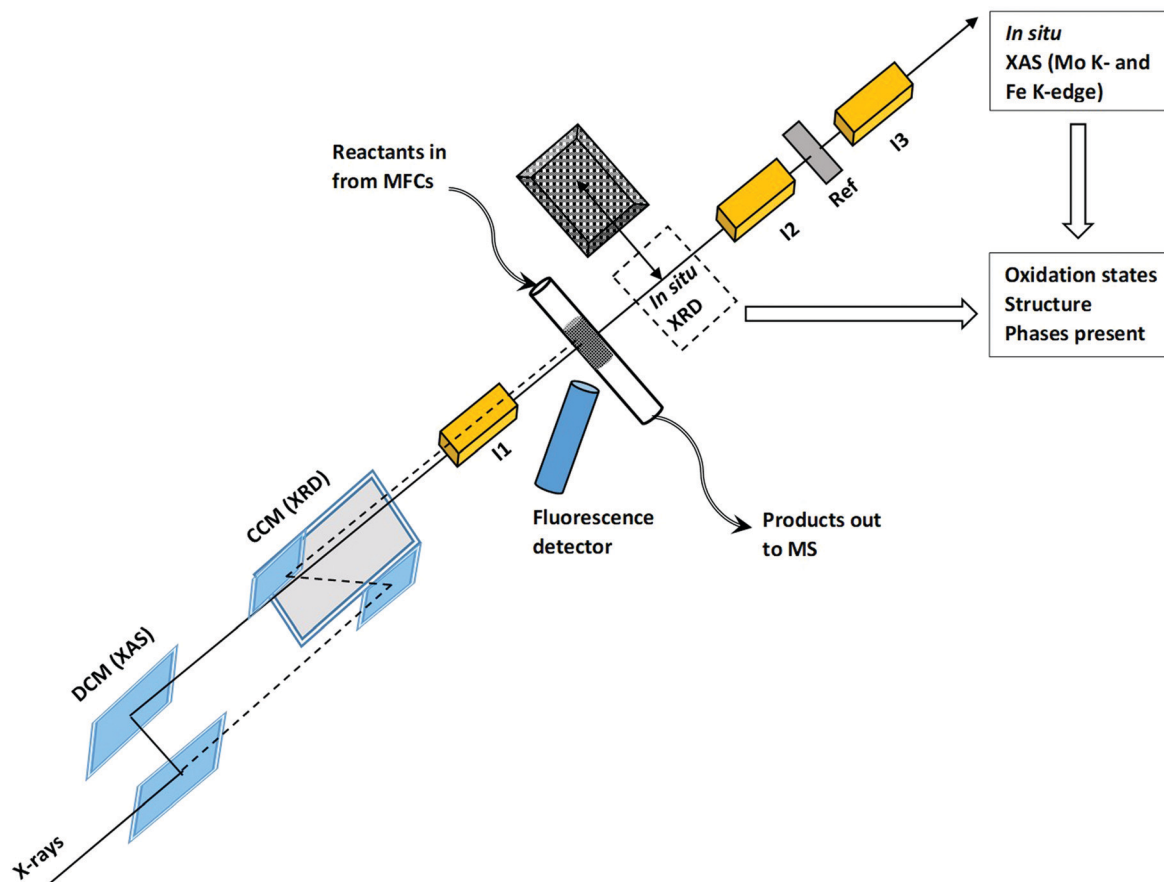


Fig. 1 Schematic setup for *in situ* XAS/XRD experiments performed at the unique setup at SNBL beamline BM31, ESRF. Double crystal monochromator (DCM), channel-cut monochromator (CCM), ionization chambers (I1, I2 and I3), reference foil (Ref), mass flow controllers (MFCs) and mass spectrometer (MS).

The overview of the experimental procedure followed during *in situ* experiments is given in the ESI,† Fig. S1. The catalyst was first heated to 100 °C in a flow of 10% O<sub>2</sub>/He. Then the flow was changed to 5% MeOH/He and the sample was heated to 400 °C at 1 °C min<sup>−1</sup> for temperature programmed reduction (TPR) and was kept at 400 °C for 1 hour. The sample was cooled to 200 °C after which the flow was changed back to 10% O<sub>2</sub>/He. In the next step, the sample was heated to 500 °C at 1 °C min<sup>−1</sup> for temperature programmed oxidation (TPO) and after a stabilization period of 1 hour finally cooled to 100 °C.

XAS data were treated with the software package IFEFFIT.<sup>25</sup> The spectra were energy calibrated, normalized, and background subtracted in Athena. Linear combination fitting (LCF) of XANES spectra was performed in a range of −20 to +50 eV at the Mo K- and Fe K-edges, respectively. EXAFS fitting was performed in Artemis by using model structures obtained from references of Fe<sub>2</sub>(MoO<sub>4</sub>)<sub>3</sub> and MoO<sub>3</sub>, in the case of Mo. At the Fe K-edge, spectra were only measured in the XANES region due to high noise in the EXAFS region. Fittings at the Mo K-edge were performed in *R* – as well as *k*-space for determining the structural parameters which included energy shift of the path ( $\Delta E_0$ ), change in the half path length ( $\Delta R$ ), amplitude reduction factor ( $S_0^2$ ), number of identical paths (*N*) and relative mean-square displacement of the atoms included in the path (Debye–Waller factor,  $\sigma^2$ ).

Principal component analysis (PCA) at the Fe K-edge was performed using XANES spectra in the range of −20 to +80 eV using the PCA subroutine of Demeter.<sup>25</sup>

XRD patterns were recorded with a CMOS-Dexela 2D detector at a short X-ray wavelength of 0.4934 Å (25.13 keV) so that the penetration of X-rays is optimal. For each diffractogram, five bright and additionally five dark images for background subtraction were recorded in the  $2\theta$  range of 2–35° with a data point spacing of approximately 0.01° and subsequently averaged. The sample to detector distance was calibrated with a LaB<sub>6</sub> reference. For XRD data treatment, the diffractograms were normalized to the most intense reflection of the SiO<sub>2</sub> dilutant, and a normalized diffractogram of only the dilutant measured at the same temperature was subtracted using an in-house Matlab script.

## 3. Results and discussion

### 3.1 Initial state and temperature programmed reduction (TPR)

**3.1.1 *In situ* observations at the Mo K-edge.** Fig. 2 shows the *in situ* Mo K-XANES spectra of the FeMo catalyst during heating under reducing conditions of 5% MeOH/He (TPR).



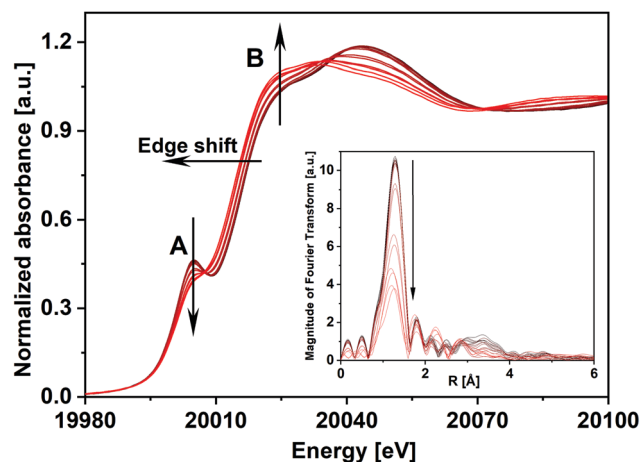


Fig. 2 *In situ* Mo K-edge XANES spectra and corresponding  $k^3$ -weighted Fourier transformed EXAFS spectra (inset) of the FeMo catalyst during TPR under 5% MeOH/He. Arrows point towards increasing temperature.

For comparison, *ex situ* XANES spectra of the FeMo catalyst at 100 °C along with spectra of the reference samples  $\text{Fe}_2(\text{MoO}_4)_3$  and  $\alpha\text{-MoO}_3$  are shown in ESI,† Fig. S2. The Mo K-XANES spectra exhibit a pre-edge peak A at  $\sim 20007$  eV and another feature B at 20027 eV. The pre-edge peak A is attributed to the dipole-forbidden/quadrupole-allowed 1s–4d transition, which is associated primarily with tetrahedral geometry. The feature is also present but to weaker extent in distorted octahedral geometry. Feature B at 20027 eV is assigned to the dipole-allowed 1s–5p transition, a characteristic feature of Mo species with octahedral/distorted octahedral geometry.<sup>26</sup> A comparison of the XANES spectrum of the FeMo catalyst with  $\text{MoO}_3$  showed that the catalyst had a slightly stronger feature A than for  $\text{MoO}_3$  and weaker intensity for peak B. Also, the intensity of peak A for the catalyst was considerably lower compared to the reference spectrum of  $\text{Fe}_2(\text{MoO}_4)_3$ .<sup>27</sup> This showed that in the FeMo catalyst, Mo had both tetrahedral ( $\text{Fe}_2(\text{MoO}_4)_3$  nature) as well as octahedral geometry ( $\text{MoO}_x$  nature, *i.e.*, molybdenum oxide with an average oxidation state between Mo(IV) to Mo(VI)).

As the initial state of the FeMo catalyst at 100 °C indicated mixed Mo(VI) phases, linear combination fitting (LCF) of the XANES spectrum was performed with standards  $\text{Fe}_2(\text{MoO}_4)_3$  and  $\alpha\text{-MoO}_3$ . This showed 76% contribution from ferric molybdate and 24% contribution from molybdenum trioxide. This is in agreement with the composition of the fresh catalyst as determined by XRD containing 83 wt%  $\text{Fe}_2(\text{MoO}_4)_3$  and 17 wt%  $\alpha\text{-MoO}_3$ , which corresponds to 22 mol% of the Mo bound as  $\text{MoO}_3$  and 78 mol% as  $\text{Fe}_2(\text{MoO}_4)_3$ .<sup>7</sup>

The XANES features of the spectra during TPR (Fig. 2) showed that the edge position shifted to lower energies, from 20012.3 to 20010.2 eV, revealing partial reduction of Mo. The decrease in intensity of peak A and simultaneous increase in the intensity of feature B showed a rise in the octahedral nature of Mo centers, corresponding to  $\text{Fe}_2(\text{MoO}_4)_3$  being decomposed to  $\text{FeMoO}_4$  and thereby forming  $\text{MoO}_x$  species *e.g.*  $\text{MoO}_2$  and intermediate  $\text{MoO}_x$  ( $x = 2\text{--}3$ ) species. However, it can be seen that after TPR, peak A was still present showing that Mo(VI) was

not completely reduced to Mo(IV), which does not have this feature. Since  $\text{MoO}_2$  has a strictly octahedral field, the 1s–4d transition is forbidden and no pre-edge peak would be observed.<sup>28</sup> The corresponding Fourier transformed EXAFS spectra during TPR are shown in the inset of Fig. 2 where in the initial state of the catalyst a high amplitude of the first peak (Mo–O<sub>1</sub>) indicated strong tetrahedral nature due to a high content of  $\text{Fe}_2(\text{MoO}_4)_3$ . Metal backscattering at  $\sim 3$  Å (Fe) additionally indicated the presence of  $\text{Fe}_2(\text{MoO}_4)_3$ .<sup>26</sup> The fitting results for the initial state of the catalyst at 100 °C in 10% O<sub>2</sub>/He (see ESI,† Table S1) also supported the presence of multiple Mo phases (see also ESI,† Fig. S3(a)). At about 290 °C, this metal backscattering disappeared due to the reduction of  $\text{Fe}_2(\text{MoO}_4)_3$  (Fig. S3(b), ESI†). In the final state during TPR at 350 °C, the amplitude of the Mo–O<sub>1</sub> contribution was highly reduced showing the presence of disordered Mo oxide phases (Fig. S3(c), ESI†).

In order to determine the contribution of different phases during the TPR, LCF was performed using standard spectra of  $\text{Fe}_2(\text{MoO}_4)_3$ ,  $\text{MoO}_3$  and  $\text{MoO}_2$ .  $\text{FeMoO}_4$  was not available as reference due to unavailability of this compound commercially and its instability during synthesis. Also, at the Mo K-edge weak XANES features present in the  $\text{FeMoO}_4$  phase were found to be overlapping with strong features of the Mo(VI) phases  $\text{Fe}_2(\text{MoO}_4)_3$  and  $\text{MoO}_3$  (for details see ESI,† Fig. S4). Thus, the formation of  $\text{FeMoO}_4$ , and associated reduction of iron, is measured and discussed more clearly using Fe K-edge XANES in Section 3.1.2. It is worth mentioning here that PCA cannot be applied at Mo K-edge due to the unavailability of set of proper standard spectra. The contributions of the different phases obtained from LCF analysis of the XANES spectra at the Mo K-edge during TPR are shown in Fig. 3(a). Reduction of Mo started at approximately 270 °C and the contribution from  $\text{MoO}_2$  was observed at the expense of an initial decrease in  $\text{MoO}_3$  content. In the final state at 400 °C, contributions from  $\text{Fe}_2(\text{MoO}_4)_3$ ,  $\text{MoO}_3$  and  $\text{MoO}_2$  were observed to be 43%, 20% and 37%, respectively. Note that  $\text{MoO}_3$  and  $\text{Fe}_2(\text{MoO}_4)_3$  together represent the Mo(VI) present as  $\text{FeMoO}_4$  after reduction, see also Fe K-edge XANES and XRD results in Section 3.1.2. Thus, at 400 °C, about 40% of the Mo(VI) species were reduced to Mo(IV) species, represented by  $\text{MoO}_2$ . Some  $\text{MoO}_x$  ( $x = 2\text{--}3$ ) may also be presented by a combined contribution from  $\text{MoO}_3$  and  $\text{MoO}_2$  as it is hard to detect these species by their XANES fingerprint. More than 50% of the Mo was still present in oxidation state VI, which is in agreement with a previous *in situ* catalytic study<sup>9</sup> at 250 °C, which showed that during formaldehyde production only Mo(VI) was present throughout the experiment, indicating that gas phase oxygen transfer to molybdenum is rapid under reaction conditions at temperatures near the lower range of industrial conditions (typically 220 to 350 °C).

Corresponding EXAFS spectra were analyzed by performing fitting in *R*-space and variation of selected obtained parameters are given in Fig. 3(b). Initially at 100 °C, the CN of Mo–O<sub>1</sub> was found to be  $5.0 (\pm 0.4)$  at 1.76 Å, which indicates strong contribution from the tetrahedral Mo centers present in  $\text{Fe}_2(\text{MoO}_4)_3$ . A considerable decrease in the CN of Mo–O<sub>1</sub> was observed at approximately 290 °C (see also Fig. S3(b, c) and Table S2, ESI†).





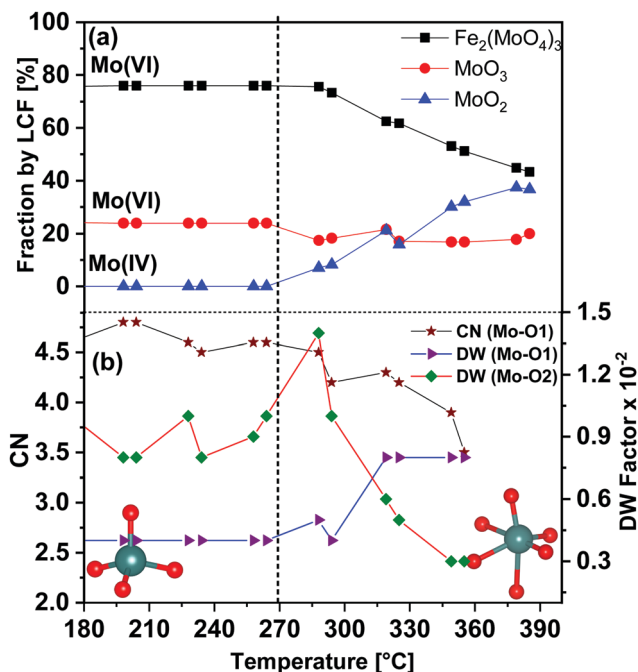


Fig. 3 (a) Mo K-edge XANES linear combination fitting (LCF) results for the FeMo catalyst during heating under 5% MeOH/He employing XANES spectra of the reference compounds  $\text{Fe}_2(\text{MoO}_4)_3$ ,  $\text{MoO}_3$  and  $\text{MoO}_2$ . (b) Variation in the selected parameters during TPR as determined from EXAFS fitting analysis in  $R$  space. Dashed line indicates the temperature at which Mo reduction started. The insets show the transition from tetrahedral to octahedral coordination geometry of Mo during TPR, Mo (blue spheres) and O (red spheres). CN of Mo–O2 bonds were kept fixed to 1.

Also, the DW factor for Mo–O1 started increasing in parallel with the reduction process and was almost doubled at the end of TPR, showing higher disorder in these bonds. Conversely, the DW factor for the longer Mo–O2 bonds (axial position), which corresponds to octahedral Mo centers, decreased at higher temperature showing stronger axial bonds due to an increase in octahedral nature of Mo centers present in  $\text{MoO}_x$  species (the CN of each of Mo–O2 bond were fixed to 1). Hence, with increasing temperature, a decrease in Mo oxidation state and an increase in octahedral nature of Mo centers were observed. These structural changes point towards formation of  $\text{MoO}_2$  and as well as  $\text{MoO}_x$  species during TPR (see also Fe K-edge XANES and XRD results in Section 3.1.2).

### 3.1.2 In situ observations by Fe K-edge XAS and XRD.

Fig. 4(a) shows the *in situ* Fe K-XANES spectra during TPR, measured in the loop with Mo K-edge spectra to simultaneously probe the oxidation state and phase transitions of Fe. The features observed for the initial state of Fe at 100 °C show similarity with the  $\text{Fe}_2(\text{MoO}_4)_3$  spectral features reported earlier.<sup>29</sup> Initially, the Fe K-edge absorption peak maxima was observed at 7131.8 eV, which is similar to the value reported in literature.<sup>29</sup> At higher temperature, the spectra exhibited a gradual shift towards lower energy indicating a change in the average oxidation state of Fe from III to II, mainly corresponding to formation of  $\text{FeMoO}_4$ , however the presence of other Fe(II) species cannot be ruled out. In the temperature

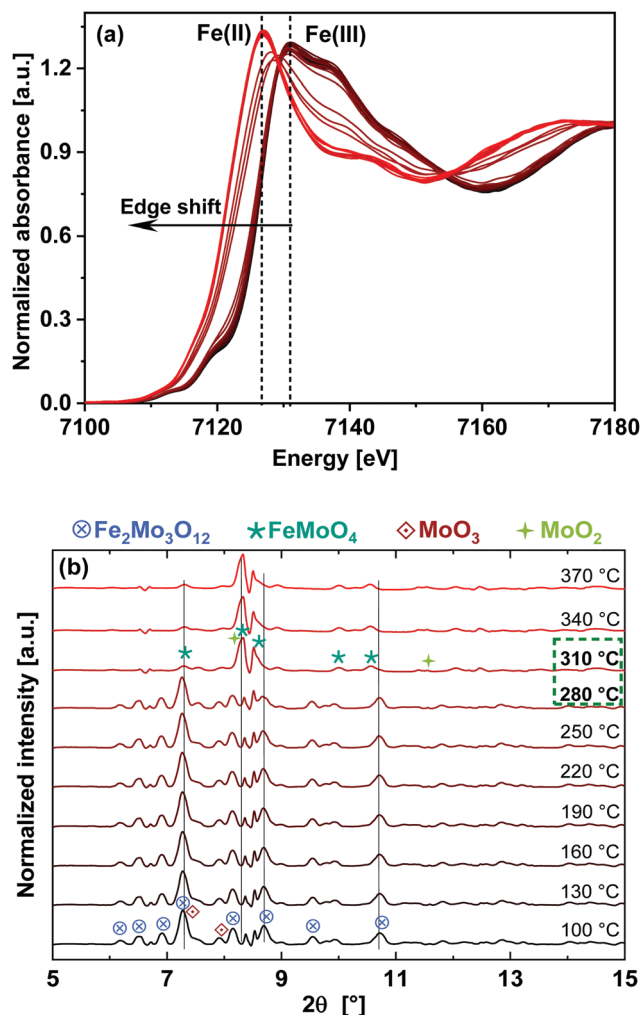


Fig. 4 (a) *In situ* Fe K-edge XANES spectra for the catalyst during heating in 5% MeOH/He (TPR). Dashed lines indicate the energy positions of the white line peak for oxidized and reduced Fe phases. The arrow points towards increasing temperature. (b) *In situ* XRD at 0.4934 Å (25.13 keV) during TPR in 5% MeOH/He. Phase transition to  $\text{FeMoO}_4$  can be observed by the increasing intensity of the reflections at 8.32 and 8.50°.

range 200–300 °C, the presence of iron oxides such as  $\text{Fe}_2\text{O}_3$  and FeO were also expected as intermediate phases. Hence, PCA was performed over Fe K-edge XANES spectra measured in this temperature range to confirm the number components required to fit the spectra by LCF analysis.

PCA uses matrix operations on an experimental spectra data matrix, to give two main outputs in the form of eigenvectors and eigenvalues, thereby providing the minimum number of significant components required to satisfactorily regenerate the data matrix.<sup>30</sup> Thus, absorbance in a set of spectra can be mathematically modelled as a linear sum of individual components, called factors, which correspond to each one of the species present in the sample, plus noise. The PCA result at 275 °C is given as an example in the ESI,<sup>†</sup> Fig. S5 and Table S3. The figure shows the different components weighted by their eigenvalues where the first two components clearly dominate the spectra and the contribution from the third component is

also significant. The fourth component appears to be noise, indicating that 3 components were required to fit the data. Thus, both  $\text{Fe}_2\text{O}_3$  and  $\text{FeO}$  were evaluated as standards during LCF, but  $\text{Fe}_2\text{O}_3$  gave better fit results, confirming its presence as intermediate phase during TPR (200–300 °C).

Results obtained from LCF analysis performed using these XANES spectra are given in the ESI,<sup>†</sup> Fig. S6. It was observed that the reduction of Fe starts at approximately 270 °C, parallel to reduction of Mo.  $\text{Fe(ox)}$  is the initial state which mainly corresponds to  $\text{Fe}_2(\text{MoO}_4)_3$  and  $\text{Fe(red)}$  is the final state after TPR, which have been taken as the standards for LCF analysis. The final reduced state of Fe, having features similar to  $\text{FeMoO}_4$ ,<sup>29</sup> was observed at 330 °C, showing faster reduction of Fe compared to Mo. These results, combined with the Mo K-edge results, indicate that at the end of TPR at 400 °C, the phases present were  $\text{Fe}_2(\text{MoO}_4)_3$ ,  $\text{FeMoO}_4$ , and  $\text{MoO}_2$  (possibly including intermediate phases  $\text{MoO}_x$ ), which confirm the presence of both  $\text{Mo(IV)}$  and  $\text{Mo(VI)}$  species.

Due to the fast interchange between the monochromators, DCM for XAS and CCM for XRD, the XAS measurements at the Mo K- and Fe K-edges were further supported by quasi-simultaneously recorded XRD patterns of the FeMo catalyst during TPR under 5%  $\text{MeOH/He}$ , as shown in Fig. 4(b). An XRD pattern characteristic for  $\text{Fe}_2(\text{MoO}_4)_3$  (most intense reflection at  $2\theta = 7.32^\circ$  and other significant reflections at  $6.93, 7.55\text{--}7.63, 7.92, 8.16, 8.73\text{--}8.82^\circ$ ) at the beginning of the TPR was observed, which disappeared almost completely at 310 °C and reflections corresponding to  $\text{FeMoO}_4$  appeared (reflections at  $8.32$  and  $8.50^\circ$ ). Furthermore, at the end of TPR a weak reflection from  $\text{MoO}_2$  was also observed at  $11.6^\circ$ . In the initial state, reflections from  $\alpha\text{-MoO}_3$  were observed as well (reflections at  $7.43$  and  $8.16\text{--}8.23^\circ$ ), partly overlapping with the reflections of  $\text{Fe}_2(\text{MoO}_4)_3$ , which disappeared as well. Thus, *in situ* XRD confirms that the phase transformation of  $\text{Fe}_2(\text{MoO}_4)_3$  to  $\text{FeMoO}_4$  started at around 280 °C, in agreement with XAS results at the Mo K- and Fe K-edges.

**3.1.3 Gas phase composition during TPR.** The corresponding evolution of the gas phase composition during TPR, overlaid with the fraction of  $\text{Mo(IV)}$  (from Mo K-edge LCF),  $\text{Fe(III)}$  and  $\text{Fe(II)}$  (from Fe K-edge LCF) is shown in Fig. 5 (feed gas composition of 5%  $\text{MeOH/He}$ ). It can be observed from the figure that at  $\sim 270$  °C reduction of Fe started almost simultaneously with Mo. Here, the  $\text{Fe(II)}$  fraction obtained from Fe K-edge LCF represent  $\text{FeMoO}_4$  as no other  $\text{Fe(II)}$  species were confirmed by LCF, which indicate formation of  $\text{FeMoO}_4$  from  $\text{Fe}_2(\text{MoO}_4)_3$ . Looking at the MS trace, the concentration of  $\text{MeOH}$  was observed to decrease in this temperature range, confirming the reaction of methanol with the catalyst in the temperature range from 260 to 400 °C. When staying at this temperature for 1 h, the methanol concentration increased to almost its initial value as shown in the extended X-axis of Fig. 5 for the isothermal period (red line) starting at 400 °C.

Though the MS-traces  $m/z = 29$  and  $18$  corresponding to the formation of  $\text{HCHO}$  and  $\text{H}_2\text{O}$ , respectively, have also been observed as shown in the ESI,<sup>†</sup> Fig. S7, the MS was slow at pumping down from the previous experiment performed.

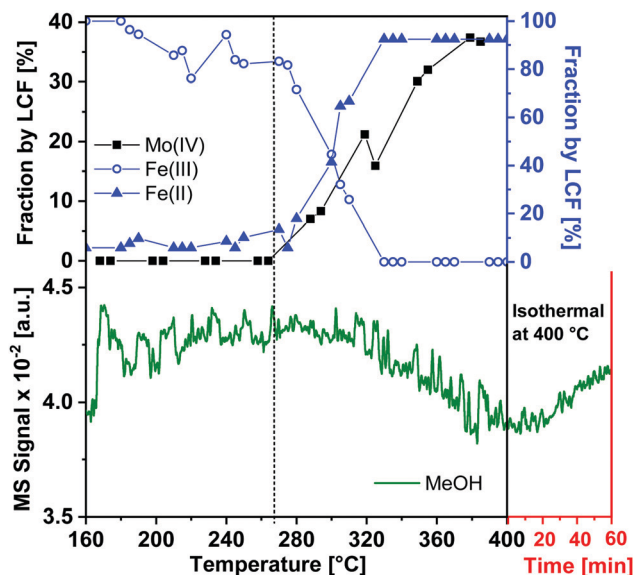
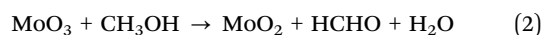
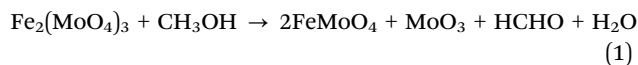


Fig. 5 Fraction of  $\text{Mo(IV)}$  (as determined from Mo K-edge LCF),  $\text{Fe(III)}$  and  $\text{Fe(II)}$  (from Fe K-edge LCF) as function of temperature with MS trace of  $\text{MeOH}$  overlaid during heating (100 to 400 °C at  $1\text{ }^\circ\text{C min}^{-1}$ ) under reducing conditions of 5%  $\text{MeOH/He}$ . The extended X-axis (in red) shows the isothermal period at 400 °C for the next 60 minutes. The dashed line indicates the start of reduction of Mo as well as Fe species.

Hence there was remarkably high background signal of  $\text{HCHO}$  and  $\text{H}_2\text{O}$  during the TPR up to about 300 °C. Nevertheless, it can be observed from Fig. S7 (ESI<sup>†</sup>) that during heat up the concentration of  $\text{MeOH}$  decreased and those of  $\text{HCHO}$  and  $\text{H}_2\text{O}$  increased, as expected, and during the isothermal period, when the reduction was complete, the concentration of  $\text{MeOH}$  increased and those of  $\text{HCHO}$  and  $\text{H}_2\text{O}$  decreased, both with some delay due to the MS pumping down.

Hence, using *in situ* XAS/XRD with MS measurements of the gas phase, it can be concluded that during heating under reducing conditions,  $\text{Fe}_2(\text{MoO}_4)_3$  and  $\text{MoO}_3$  were observed to reduce thereby forming  $\text{FeMoO}_4$  and  $\text{MoO}_2$  which is in agreement with the proposed reactions (1) and (2).



As described earlier, after TPR the catalyst was kept isothermal at 400 °C in 5%  $\text{MeOH/He}$  for 1 hour. Then the temperature was lowered to 200 °C and after stabilization at 200 °C the gas conditions were changed to 10%  $\text{O}_2/\text{He}$  (ESI,<sup>†</sup> Fig. S1). The phases of the catalyst remained mainly intact during the cool down, as well as during switching to oxidizing conditions as indicated by the XAS/XRD results given in the ESI,<sup>†</sup> Fig. S8. At the Mo K-edge a slight shift of the edge was observed towards lower energies owing to a small increase in reduced phases of Mo. At the Fe K-edge in the post edge region some features were enhanced, indicating less distortion in the structure of Fe species after cooling down.

### 3.2 Heating under oxidizing conditions (TPO)

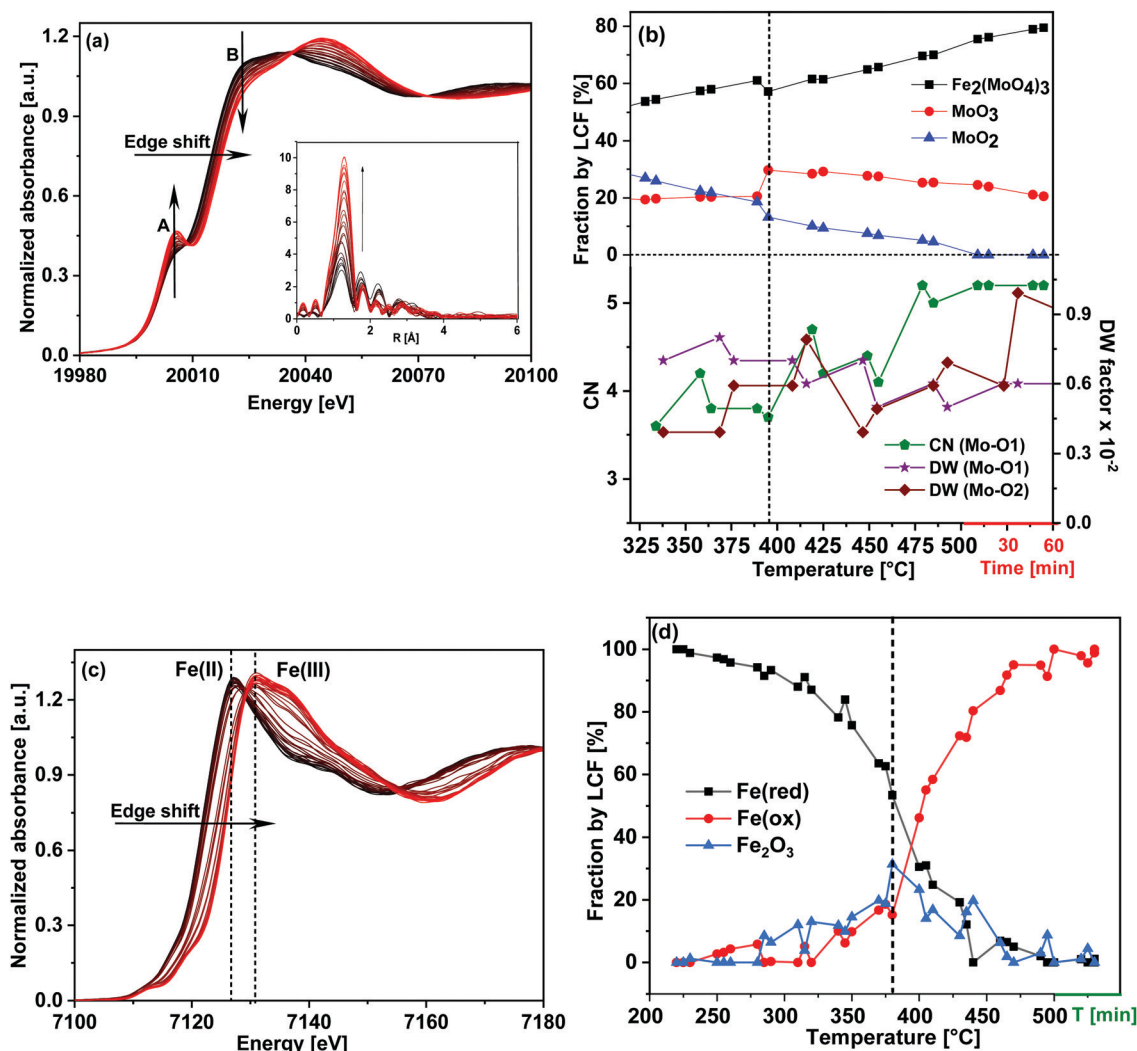
**3.2.1 *In situ* XAS at Mo and Fe K-edges and XRD.** Fig. 6(a) shows the *in situ* Mo K-edge XANES spectra of the catalyst during heating under oxidizing conditions (TPO) after the TPR and cool down to 200 °C in 5% MeOH/He. The edge position shifted towards higher energies showing an increase in the oxidation state of Mo, as expected. The pre-edge peak A regained intensity lost during TPR, corresponding to an increase in tetrahedral nature of the Mo centers, and the simultaneous weakening of feature B indicated lowering of octahedral nature.

LCF analysis of the XANES spectra using the same standards as during TPR and variations of the parameters obtained from EXAFS analysis are shown in Fig. 6(b) in the temperature range 320–500 °C (for the complete LCF results from 200–500 °C please see ESI,† Fig. S9). The initial composition before TPO at 200 °C was 39%  $\text{Fe}_2(\text{MoO}_4)_3$ , 20%  $\text{MoO}_3$  and 41%  $\text{MoO}_2$  which is similar to the composition of the final state after TPR

(43%  $\text{Fe}_2(\text{MoO}_4)_3$ , 20%  $\text{MoO}_3$  and 37%  $\text{MoO}_2$ ) within the margin of error of LCF and considering the temperature difference.

Oxidation of Mo began as soon as the temperature started to rise which is evident from the decreasing  $\text{MoO}_2$  and increasing  $\text{Fe}_2(\text{MoO}_4)_3$  contributions, showing that overall  $\text{MoO}_2$  combined with  $\text{FeMoO}_4$  to give  $\text{Fe}_2(\text{MoO}_4)_3$ . The content of  $\text{MoO}_3$  was quite stable until 380 °C, but in the range of 380–400 °C a sharp increase was observed (Fig. 6(b)), indicating formation of  $\text{MoO}_3$  as well.

At the same temperature, the EXAFS parameters (Fig. 6(b)) showed a decrease in the DW factor of Mo–O1 and an increase in that of Mo–O2 supporting formation of octahedral Mo centers present in  $\text{MoO}_3$ . The contribution from  $\text{MoO}_2$  continuously decreased and became negligible at around 500 °C, showing that almost all of Mo was in the Mo(vi) state. After the sudden increase at approximately 390 °C, the contribution from  $\text{MoO}_3$  fell back to 21%, slightly lower than what was present initially before TPO (24%). The remaining 80% of Mo(vi) was found to be



**Fig. 6** For the FeMo catalyst during heating under oxidizing conditions 10%  $\text{O}_2/\text{He}$  (TPO): (a) Mo K-edge XANES spectra, (b) corresponding XANES LCF results and variation in the selected parameters as determined from EXAFS analysis. At the X-axis, the red line after 500 °C represents the isothermal period, (c) Fe K-edge XANES spectra and (d) corresponding XANES LCF results. In (a) and (c) arrows point towards increasing temperature. The vertical dashed lines in (b) and (d) indicate the temperature  $\sim 390$  °C at which a sudden increase in  $\text{MoO}_3$  and  $\text{Fe}_2\text{O}_3$  content was observed, respectively.





$\text{Fe}_2(\text{MoO}_4)_3$ , which is slightly higher compared to the initial composition of the catalyst at 100 °C before TPR (76%). It is possible that some  $\text{MoO}_3$  formed volatile species and evaporated during the TPR, similar to the evaporation clearly observed under reaction conditions for methanol oxidation.<sup>20</sup> EXAFS analysis also confirmed this conclusion since the CN of Mo–O1 at 1.77 Å increased and the corresponding DW factor decreased, showing a rise in tetrahedral nature of Mo. Similarly, the DW factor of the Mo–O2 (axial) concurrently increased, showing a decrease in octahedral nature of Mo centers (the CN of both Mo–O2 bonds were kept fixed to 1).

Fig. 6(c) shows the Fe K-edge XANES spectra during heating under oxidizing conditions. PCA was performed to confirm the number components required to fit the spectra and the results obtained from PCA analysis for Fe K-edge XANES spectrum at 380 °C is given as an example in ESI,† Fig. S10 and Table S4. In this case also, the first two components clearly dominate the spectra, with a significant contribution from the third component and the fourth component appears to be noise, indicating that 3 components were required to fit the data. Thus,  $\text{Fe}_2\text{O}_3$  was also employed as standard during LCF, which gives better fit results than FeO, confirming its presence as intermediate phase during TPO (300–450 °C). LCF analysis shown in Fig. 6(d) showed that oxidation of Fe effectively started at 280 °C. The contribution from  $\text{Fe}_2\text{O}_3$  was found to be high (around 30%) between 380–410 °C. In this temperature range LCF at the Mo K-edge also showed a high presence of  $\text{MoO}_3$  pointing towards  $\text{FeMoO}_4$  reacting with  $\text{O}_2$  to give  $\text{Fe}_2\text{O}_3$  and  $\text{MoO}_3$  (eqn (3)) in competition with the reactions regenerating the catalyst (eqn (4) and (5)).<sup>31</sup> However, like the excess  $\text{MoO}_3$ , the  $\text{Fe}_2\text{O}_3$  phase was an intermediate and the contribution was found to decrease at higher temperature. At 500 °C Fe was found mostly as  $\text{Fe}_2(\text{MoO}_4)_3$ , due to recombination of  $\text{Fe}_2\text{O}_3$  and  $\text{MoO}_3$  by solid state reaction (eqn (6)). These redox dynamics explain the formation of  $\text{Fe}_2\text{O}_3$  observed under reaction conditions for selective oxidation of methanol at 380 to 400 °C, where  $\text{MoO}_3$  evaporation competes with the recombination and with extended time on stream  $\text{MoO}_3$  eventually evaporates completely.<sup>7</sup>

Corresponding *in situ* XRD results are given in Fig. 7. They also show the presence of  $\text{FeMoO}_4$  and  $\text{MoO}_2$  initially, as observed at the end of TPR. During TPO, a phase transition was observed between 380–410 °C where higher presence of  $\text{MoO}_3$  (by LCF at Mo K-edge data) and  $\text{Fe}_2\text{O}_3$  (by LCF at Fe K-edge data) were observed by XANES. Only  $\text{Fe}_2(\text{MoO}_4)_3$  and  $\text{MoO}_3$  were observed as oxidation products by XRD, which agrees with the overall re-oxidation and recombination reactions given in eqn (3), (4) and (6). The  $\text{Fe}_2\text{O}_3$  detected by LCF of the XANES data was an intermediate, possibly an amorphous, phase and thus was not detected by XRD.

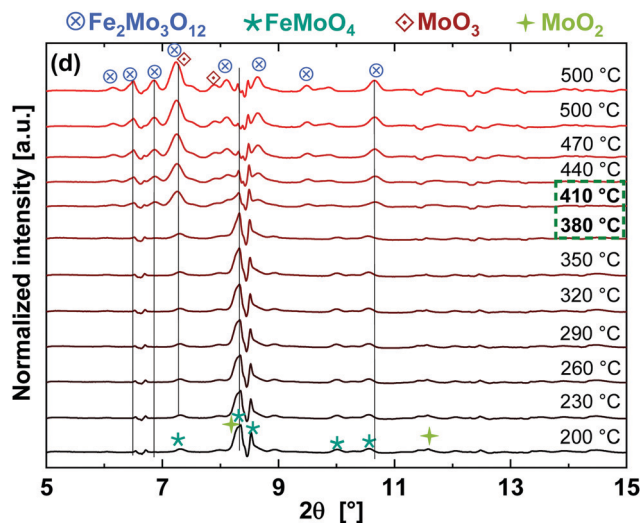
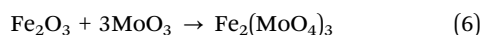
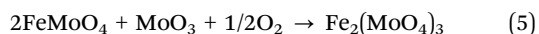
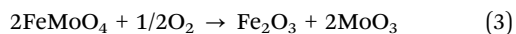


Fig. 7 XRD patterns at different temperatures during TPO (at 25.23 eV corresponding to  $\lambda = 0.4934$  Å).

Overall, the results obtained from XRD correspond to those obtained from LCF of XANES showing that  $\text{FeMoO}_4$  and  $\text{MoO}_2$  were oxidized and recombined to form  $\text{Fe}_2(\text{MoO}_4)_3$ .

**3.2.2 Reversibility of the reduction and re-oxidation.** To check whether the changes that occurred during the complete cycle of TPR and TPO were reversible, the initial state of the catalyst at 100 °C before reduction, the state of the catalyst at 500 °C after reoxidation and the final state of the catalyst after cooling down to 100 °C were compared. Fig. 8(a–d) show the Mo K-edge XANES spectra, corresponding FT spectra, Fe K-edge XANES spectra and XRD reflections of the catalyst at these different stages of the redox cycle. The identical features observed in the Mo and Fe XANES showed that Mo and Fe have attained their initial state with respect to oxidation state and coordination geometry. Similar conclusions can be drawn from the corresponding XRD patterns showing the presence of identical phases at these stages, though at 500 °C there is a temperature shift.

In Fig. 8(b), the FT EXAFS spectra are given and Table 1 shows the fitting parameters (see also ESI,† Fig. S3(a, e and f)). The major difference was observed in the FT spectra in the region of 2.5–4.0 Å where there are backscattering peaks due to Mo–Fe and Mo–Mo contributions. The Mo–Mo peak at 3.7 Å completely disappeared at 500 °C after the redox cycling, which indicates high distortions in the Mo–Mo metal shells related to the  $\text{MoO}_3$  phase. By cooling down to 100 °C, this backscattering peak was observed to be regenerated indicating that the catalyst returns to its crystalline phase. At 500 °C, the CN of the first peak (Mo–O1) was comparatively higher than its value at 100 °C in the initial as well as the final state, which indicate stronger tetrahedral nature of Mo coordination at higher temperature. In case of the longer axial Mo–O2 bonds, a considerable increase in the DW factor was observed between the initial and final states at 100 °C. This indicates loss of octahedral character of the Mo centers during the redox cycle, corresponding to evaporation of some of the excess  $\text{MoO}_3$ . However, changes in the  $\text{MoO}_3$  crystallite particle size or shape during the redox cycling may also occur. As the CN of Mo–O2





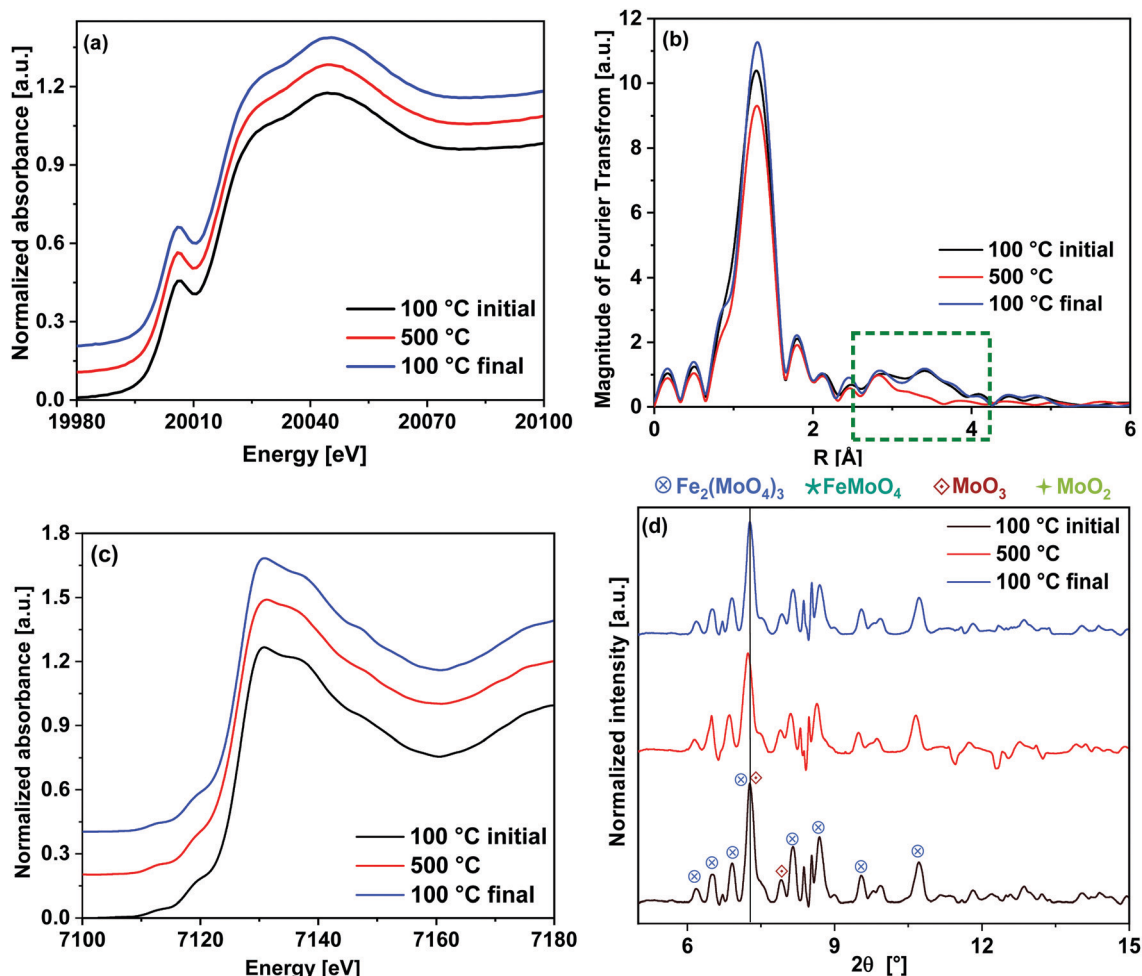


Fig. 8 Comparison of the initial state of the catalyst measured at 100 °C, the state at 500 °C after TPO and the final state after cool-down to 100 °C under 10% O<sub>2</sub>/He: (a) Mo K-edge XANES (b) corresponding  $k^3$ -weighted FT EXAFS spectra with marked region 2.5–4.0 Å. (c) Fe K-edge XANES and (d) XRD patterns with the assignment of phases.

Table 1 Structural parameters obtained from EXAFS fitting at Mo K-edge for the state of the catalyst at 500 °C after reduction and re-oxidation, compared to the initial state of the catalyst at 100 °C and final state after cooling to 100 °C.  $\Delta E_0 = 1.0$  and  $\chi^2_\nu = 65$  for 500 °C,  $\Delta E_0 = 2.1$  and  $\chi^2_\nu = 17$  for 100 °C (final)

Path	100 °C (initial)			500 °C			100 °C (final)		
	$R$ [Å]	CN	$\sigma^2$ [Å <sup>-2</sup> ]	$R$ [Å]	CN	$\sigma^2$ [Å <sup>-2</sup> ]	$R$ [Å]	CN	$\sigma^2$ [Å <sup>-2</sup> ]
Mo–O <sub>1</sub>	1.76 ± 0.01	5.0 ± 0.5	0.0046 ± 0.0006	1.77 ± 0.02	5.2 ± 0.4	0.0054 ± 0.0007	1.77 ± 0.02	4.3 ± 0.4	0.0030 ± 0.0006
Mo–O <sub>2</sub>	2.04 ± 0.06/ 2.36 ± 0.06	1 <sup>a</sup> /1 <sup>a</sup>	0.0084 ± 0.0051	2.04 ± 0.07/ 2.37 ± 0.07	1 <sup>a</sup> /1 <sup>a</sup>	0.0084 ± 0.0031	1.99 ± 0.06/ 2.31 ± 0.06	1 <sup>a</sup> /1 <sup>a</sup>	0.0122 ± 0.0063
Mo–Fe	3.50 ± 0.02	1 <sup>a</sup>	0.0064 ± 0.0025	3.5 ± 0.06	1 <sup>a</sup>	0.0074 ± 0.0031	3.50 ± 0.02	1 <sup>a</sup>	0.0064 ± 0.0021
Mo–Mo	3.73 ± 0.03	1 <sup>a</sup>	0.0064 ± 0.0025	—	—	—	3.7 ± 0.03	1 <sup>a</sup>	0.0064 ± 0.0021

<sup>a</sup> Fixed during the fitting.

bonds were kept fixed to the value of 1, the DW factor obtained from the fitting corresponds to the strength of these bonds.

## 4. Conclusions

We have shown that the advanced setup at the synchrotron facilities makes it possible to obtain bulk structural as well as

chemical information on the catalyst system under investigation. The combination of transmission XAS at the Mo K-edge, fluorescence XAS at the Fe K-edge and XRD at high energy proved to be a successful tool for detection of individual crystalline and amorphous phases *in situ* during the different stages of redox cycling of an iron molybdate catalyst with excess molybdenum trioxide, thereby providing important information about the phase transformations. The simplified



reducing/oxidizing conditions employed in this study are needed to gain deeper insight into the catalytic system and catalytic cycle. Subsequent XAS measurements at the Mo K- and Fe K-edges showed that reduction of Mo as well as Fe started at  $\sim 270^\circ\text{C}$  and at  $400^\circ\text{C}$   $\text{MoO}_2$  and  $\text{FeMoO}_4$  were observed as reduced phases. Complete reduction of  $\text{Fe(III)}$  to  $\text{Fe(II)}$ , forming  $\text{FeMoO}_4$ , was observed at  $330^\circ\text{C}$  whereas 60% Mo was still present as  $\text{Mo(VI)}$  after reduction at  $400^\circ\text{C}$ . Quasi-simultaneously recorded *in situ* XRD at high energy confirmed that the phase transition from  $\text{Fe}_2(\text{MoO}_4)_3$  and  $\text{MoO}_3$  to  $\text{FeMoO}_4$  and  $\text{MoO}_2$  occurred at about  $280\text{--}300^\circ\text{C}$ . Online gas phase analysis recorded simultaneously showed that the concentration of methanol decreased parallel to reduction of Fe and Mo observed by XAS and XRD, confirming the reaction between gas and solid.

Subsequent heating in an oxidative atmosphere from  $200$  to  $500^\circ\text{C}$  caused Mo and Fe to reoxidize thereby forming  $\text{Fe}_2(\text{MoO}_4)_3$ . Intermediates phases  $\text{Fe}_2\text{O}_3$  and excess  $\text{MoO}_3$  were also formed in parallel reactions, but they recombined above  $400^\circ\text{C}$  to give  $\text{Fe}_2(\text{MoO}_4)_3$ , which was observed to be the most dominant phase at  $500^\circ\text{C}$ . XRD confirmed that the main transition occurred at about  $380\text{--}400^\circ\text{C}$ , at which rise in the  $\text{MoO}_3$  and  $\text{Fe}_2\text{O}_3$  contributions were observed by XANES LCF.

The reduction/re-oxidation dynamics observed in this study, particularly the formation of  $\text{FeMoO}_4$  and  $\text{MoO}_2$  during reduction and  $\text{Fe}_2\text{O}_3$  during re-oxidation, shows the segregation of iron and molybdenum (forming  $\text{FeMoO}_4$  and  $\text{Fe}_2\text{O}_3$ ) observed under industrial reaction conditions, where the concurrent  $\text{MoO}_3$  evaporation hinders recombination to form  $\text{Fe}_2(\text{MoO}_4)_3$ , which results in catalyst deactivation and selectivity loss by iron enrichment. In the future, it would be helpful to receive complementary insight by more surface sensitive methods, typically only possible under quasi *in situ* conditions, or using samples with higher surface area that would enhance the sensitivity of XAS for the surface.

## Conflicts of interest

The authors declare no competing financial interest.

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