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Iron molybdate catalysts synthesised *via* dicarboxylate decomposition for the partial oxidation of methanol to formaldehyde†

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A series of iron molybdate catalysts were synthesised *via* a sol gel route using either oxalic acid or malonic acid. Catalysts synthesised using malonic acid were found to give improved formaldehyde yields over those prepared using oxalic acid or a standard coprecipitation method. This was attributed to the iron and molybdenum malonate precursors forming discrete ions that when precipitated gave a homogeneous distribution of iron and molybdenum in the final catalyst. Metal oxalate precursors and materials synthesised using coprecipitation gave less homogeneous structures containing iron rich centres that led to combustion of methanol to carbon oxides.

Introduction

Formaldehyde is produced from methanol *via* two industrial processes; the dehydrogenation of methanol over a silver catalyst,^{1–5} or the selective oxidation of methanol over mixed metal oxide catalysts.^{6–9} The partial oxidation route is increasingly preferred as it uses a lower methanol feed (~10%) and a lower temperature (300 °C) with a more robust catalyst, reducing the cost of formaldehyde production¹⁰ compared with the dehydrogenation route,^{10,11} which relies on high methanol feeds (90%) and high temperatures (600 °C).

The industrial production of formaldehyde using mixed metal oxides is currently conducted using the Formox process, which uses an iron–vanadium, iron–vanadium–molybdenum or iron–molybdenum mixed oxide spinel catalyst. Iron molybdate was first reported as a catalyst for the oxidation of methanol to formaldehyde in 1931¹² and has been used as a commercial catalyst since the 1950s. Industrial catalysts comprise of the mixed metal oxide spinel phase, Fe₂(MoO₄)₃ together with excess MoO₃. The excess MoO₃ is thought to have a dual role in the active catalyst; to improve selectivity by modifying the surface of the iron molybdate^{13–18} and to reduce deactivation due to loss of molybdenum leading to the formation of iron-rich phases,^{19–21} which lead to combustion products.^{6,22,23}

Traditionally, iron molybdate catalysts have been synthesised by co-precipitating metal nitrate solutions using a base to yield precursors that are then calcined to form oxide catalysts with a low surface area (typically <10 m² g^{−1}).^{8,13,14} This preparation route is difficult to control leading to an inhomogeneous distribution of the metals and materials that contain mixtures of mixed oxide and single oxide phases. For iron molybdate materials this can lead to the formation of iron rich phases such as FeO_x and FeMoO₄, which reduce the selectivity of the catalysts.^{6,22,23}

A number of different synthesis methods have been investigated to try and improve the homogeneity of the materials and increase the selectivity of the catalysts to formaldehyde. Mechanochemistry has been investigated as a synthesis methodology by Huang *et al.*²⁴ and Dong *et al.*²⁵ The mixtures of iron oxide and molybdenum oxide were ground at 600–700 °C, but the limitations of the solid-state reactions meant that the resultant materials were low surface area with molybdenum rich and iron rich phases present in the final catalysts. Hydrothermal synthesis from iron nitrate and ammonium heptamolybdate has been investigated, although these offer no improvement in homogeneity resulting in a mixture of phases similar to the coprecipitated materials.^{26,27} Alternative synthetic strategies have been used to obtain phase pure materials by designing nano-structured catalysts such as Fe₂(MoO₄)₃ supported on MoO₃ nanorods²⁸ or MoO_x/FeO_x core shell structures,^{29,30} which showed high selectivity to formaldehyde, demonstrating the importance that the control of the catalyst phases has on the performance.

Sol–gel chemistry is a widely used technique for obtaining homogeneous mixed oxide materials by allowing the components to be atomically mixed in the precursor³¹ phase.

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The gel precursor contains a homogeneous distribution of ions at room temperature, although phase segregation can still occur during calcination at elevated temperatures.³² Traditional sol-gel methodologies utilise the hydrolysis and condensation of alkoxides, but the difference in hydrolysis rates of different metal alkoxides means the homogeneity is difficult to control for mixed oxides.

An alternative sol-gel strategy is to use small chelating molecules (traditionally citric acid) in aqueous solutions of metals to form the gels. This methodology gives more homogeneous solutions than traditional sol-gel chemistry and can be used for binary, ternary and quaternary metal oxides.³² Often a base is added as pH is a key factor in obtaining a homogeneous distribution of the different metals in the gel. Soares *et al.* investigated propanoic acid using this methodology to synthesise iron molybdate catalysts and found that the catalysts formed had a high surface area compared to coprecipitated materials. However, the formaldehyde selectivity was not maintained at high temperatures,^{8,33} which was thought to be due to loss of Mo during calcination, leading to Fe rich sites at the surface. Oudghiri-Hassani³⁴ investigated dissolving iron and molybdenum nitrates in oxalic acid which acts as the chelating agent and solvent. The oxalate precursors were calcined to form the iron molybdate materials. Oudghiri-Hassani did not investigate the materials for methanol oxidation, but subsequently Yeo *et al.*³⁵ found that they showed high selectivity to formaldehyde.

In this study we have extended our previous work to include malonic acid as a chelating agent/solvent to investigate the effect of the diacid chain length on the catalyst properties. Previous studies have suggested that the increased chain length allows the malonic acid to act as a bidentate ligand, leading to more isolated iron centres rather than the polymeric chains found with oxalic acid. This may improve the homogeneity of the materials, limiting the formation of iron rich species in the final catalyst, leading to improved performance for methanol oxidation to formaldehyde.

Experimental

Catalyst preparation

The preparation of the iron molybdate catalysts with Fe:Mo ratios of 1:1.5, 1:2.2 and 1:3.0 were carried out using a procedure described below with both oxalic acid and malonic acid.

Iron nitrate, $\text{Fe}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$, ammonium molybdate, $(\text{NH}_4)_6\text{Mo}_7\text{O}_{24} \cdot 4\text{H}_2\text{O}$, and the organic acid (molar ratios = 1/0.21/10, 1/0.31/10 and 1/0.42/10) were physically ground using a pestle and mortar for 10 min. The solid precursors were then heated on a hotplate at 160 °C (oxalic acid) or 130 °C (malonic acid) for 3 h. On cooling the resultant solids were calcined under flowing air (20 ml min⁻¹) in a tubular furnace (500 °C, 2 h, 10 °C min⁻¹) resulting in the final iron molybdate catalysts.

The oxalic acid derived materials were designated 1:1.5-O, 1:2.2-O and 1:3.0-O depending on the Fe:Mo atomic ratio, with the malonic acid materials designated 1:1.5-M, 1:2.2-M and 1:3.0-M.

A catalyst was also prepared using a conventional coprecipitation method. A solution of ammonium heptamolybdate tetrahydrate, $(\text{NH}_4)_6\text{Mo}_7\text{O}_{24} \cdot 4\text{H}_2\text{O}$, (9.71 g in 150 ml distilled water) was acidified to pH 2 using conc. HNO_3 (70%). A solution of iron nitrate nonahydrate, $\text{Fe}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$ (10.1 g in 150 ml distilled water) was added dropwise with vigorous stirring. The suspension was aged at 80 °C for 3 h, and the solution was cooled to room temperature and the precipitate was recovered by filtration, dried at 120 °C for 16 h and calcined in flowing air at (500 °C, 2 h, 10 °C min⁻¹, 20 ml min⁻¹) resulting in the final iron molybdate catalyst with a Fe:Mo atomic ratio of 1:2.2. This material was designated 1:2.2-C.

Catalyst characterization

XRD measurements were obtained using a Panalytical X'pert Pro diffractometer using Cu K α X-ray source operating with an accelerator voltage 40 kV and 40 mA current. X-ray patterns were recorded in the range 10–80° 2 θ . The patterns produced were compared to reference patterns supplied by the International Centre for Diffraction Data (ICDD).

Raman spectra were obtained using a Renishaw Ramascope, using an Ar⁺ laser of wavelength 514 nm as the light source. An Olympus BH2-UMA fitted with a 20 \times Leica optical zoom lens was used to focus the laser light. Samples were placed onto aluminium backing plates which are mounted onto an automated stage to allow for mapping operations. A series of spectra were obtained from a large area of the sample to confirm how homogeneous the samples were.

Elemental composition and oxidation state analysis of the calcined and uncalcined iron molybdate samples surfaces were conducted using a Thermo Scientific K α X-ray photoelectron spectrometer (XPS), with monochromatic Al radiation operating at 72 W power with a spot size of 400 μm . Dual low energy electron and Ar⁺ neutralisation was used and all results calibrated against C(1s) results where applicable. The data was analysed using CasaXPS software using Scofield sensitivity factors corrected with an energy dependence of 0.6, after application of a Shirley background.

BET surface areas were determined by N₂ absorption at -196 °C using a Quantachrome Quadrasorb-evo instrument. The samples (0.5–3.0 g) were prepared for analysis by removing physisorbed water at 120 °C for 2 h under a vacuum. Both the reference and sample were cooled to -196 °C and tested to attain an equilibrium measurement. Analysis was conducted at 5 different pressures along the isotherm before results were converted into surface area measurements (m² g⁻¹).

TGA experiments were undertaken using a Perking Elmer TGA 4000 equipped with an autosampler. 10–20 mg of



sample were loaded into pre-weighed calcination boats and were heated 30–800 °C at a rate of 5 °C min⁻¹. Both flowing air and nitrogen atmospheres were used where applicable using a flow rate of 30 ml min⁻¹.

Catalyst testing

The catalytic performance of the iron molybdate catalysts was evaluated for the partial oxidation of methanol to formaldehyde in a laboratory plug flow microreactor. The catalyst was pressed and sieved between 400 and 600 µm. Typically, 0.3 g of the catalyst was placed in a quartz reactor tube (8 mm internal diameter) held between plugs of quartz wool. The reactor was placed in a tubular furnace (Carbolite) and temperature was monitored using a k-type thermocouple at the centre of the catalyst bed. Helium was delivered to a saturator containing methanol (99.5% Sigma-Aldrich) which was maintained at 5.2 °C in a thermostatically controlled water bath. The methanol/helium and oxygen were introduced using mass flow controllers (Bronkhorst) to give a total flow rate of 60 ml min⁻¹ (MeOH:O₂:He = 5:10:85). Both inlet and outlet lines were heated to 130 °C to prevent condensation. The data was collected and analyzed using an on-line gas chromatograph (Agilent 7820A) equipped with a Porapak Q (1 m) column and a Molsieve 13 Å (80–100 mesh) column for separation of the products. The products passed through a methaniser to convert them to methane, in order to overcome detection limitations, before analysis with an FID.

Results and discussion

The XRD patterns of the materials synthesised using malonic acid and oxalic acid are shown in Fig. 1, together with the standard coprecipitated material (1:2.2-C).

For all the materials, the mixed oxide phase, Fe₂(MoO₄)₃, MoO₃ and small amounts of Fe₂O₃ were identified using powder diffraction patterns published by the International Centre for Diffraction Data (α-Fe₂(MoO₄)₃ – ICDD: 01-083-1701, α-MoO₃ – ICDD: 00-005-0508, α-Fe₂O₃ ICDD: 01-071-

5088). Using the reference intensity ratios (RIR) from these reference patterns, based on the intensity of the reflections in the powder pattern compared to corundum (I/I_c), the relative amounts of each phase could be estimated, and these are shown in Table 1.

Although the use of RIR can only be considered semi-quantitative, there are some general observations that can be made from the analysis. The first thing to note is that all samples contain iron oxide. This is perhaps not surprising for the materials that contain a stoichiometric amount molybdenum (Fe:Mo = 1:1.5), but is unexpected in the materials containing excess molybdenum as this is thought to limit the formation of iron rich phases. The relative amount of iron rich phases is seen to increase with increasing molybdenum content which is counter intuitive and suggests that X-ray amorphous MoO₃ is formed as the molybdenum content increases.

A further observation to note from the XRD patterns is that the different synthetic methods do not lead to a large difference in composition, with the relative amounts of each phase consistent for the coprecipitated material and sol-gel samples made using oxalic and malonic acid (Table 1). This is surprising, as one of the considerations for using sol-gel chemistry is to improve the homogeneity of the final material, minimizing the amount of iron rich phases present in the final catalysts.

Katelnikovas *et al.* have evaluated different chelating agents using the expression:³⁶

$$k = n + \log M$$

where n is the number of OH groups and M is the molar mass of the molecule. When the synthesis of yttrium aluminium garnet was investigated a k value of >3 was required to synthesise the phase pure mixed oxide and chelating agents with a k value of <3 resulted in single phase formation. Using this expression, oxalic and malonic acid would have k values of 1.95 and 2.02 respectively which could explain the presence of Fe₂O₃ in the catalysts, even with excess molybdenum. However, Katelnikovas *et al.* also showed that optimum pH and chelating agent concentrations were important factors. These variables were not investigated for the synthetic methodology used in the

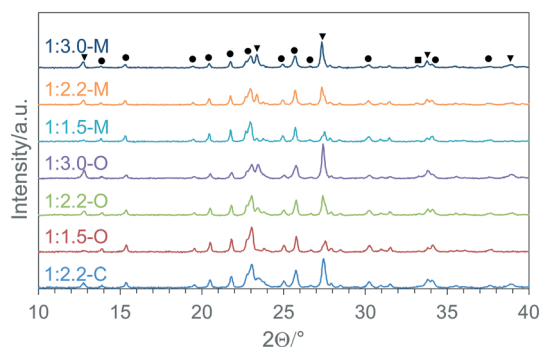


Fig. 1 X-ray diffraction patterns of iron molybdate materials synthesised using malonic acid, oxalic acid and coprecipitation (key: ▼ MoO₃; ● Fe₂(MoO₄)₃; ■ Fe₂O₃).

Table 1 Relative phase composition of iron molybdate samples determined from the reference intensity ratios from published powder patterns in the ICDD database

Sample	Composition from powder XRD patterns (%)			Theoretical composition (%)	
	Fe ₂ (MoO ₄) ₃	MoO ₃	Fe ₂ O ₃	Fe ₂ (MoO ₄) ₃	MoO ₃
1:2.2-C	65.3	33.2	1.5	68.2	31.8
1:1.5-O	85.7	13.5	0.8	100.0	0.0
1:2.2-O	71.0	27.9	1.1	68.2	31.8
1:3.0-O	48.4	49.5	2.1	50.0	50.0
1:1.5-M	88.8	10.4	0.8	100.0	0.0
1:2.2-M	68.8	28.7	2.5	68.2	31.8
1:3.0-M	49.0	47.1	3.9	50.0	50.0



current study that used oxalic acid ($pK_{a1} = 1.46$, $pK_{a2} = 4.40$) and malonic acid ($pK_{a1} = 2.5$, $pK_{a2} = 8.7$) as both the solvent and chelating agent.

The composition of the materials was further studied using Raman spectroscopy and X-ray photoelectron spectroscopy (XPS). Representative Raman spectra taken from different areas of the materials (Fig. 2 and S1†) show that the materials display some compositional inhomogeneity with the $\text{MoO}_3:\text{Fe}_2(\text{MoO}_4)_3$ ratio varying across the samples, although this is less pronounced as the molybdenum content increases. However, it is clear that when a large number of spectra are compared from different areas of the materials there is the expected trend, with MoO_3 peaks become more intense with increasing Mo:Fe ratio in the samples (Fig. S1†). One thing to note is that Fe_2O_3 was not observed by Raman spectroscopy, although this is unsurprising due to hematite being a good absorber of the incident radiation, leading to low scattering intensities compared to the molybdenum containing phases.

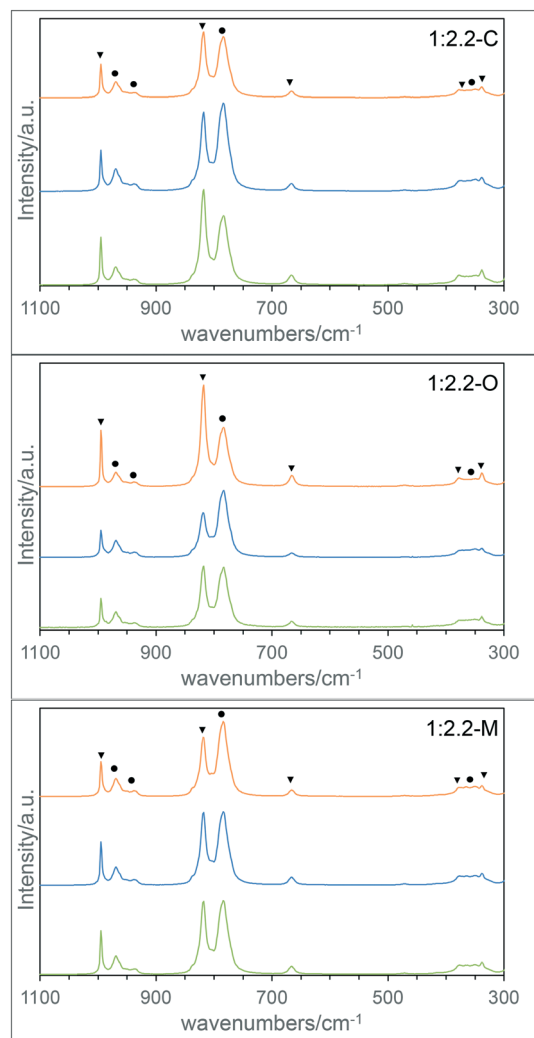


Fig. 2 Representative Raman spectra from the 1:2.2-C, 1:2.2-O and 1:2.2-M materials (key: ▼ MoO_3 ; ● $\text{Fe}_2(\text{MoO}_4)_3$).

XPS spectroscopy showed that all samples had a molybdenum rich surface compared to the bulk, and that this was higher for the oxalic acid and malonic acid derived materials than the coprecipitated sample (Table 2). This might be expected as an MoO_3 monolayer on the surface of $\text{Fe}_2(\text{MoO}_4)_3$ has been postulated to be the active site for methanol oxidation¹⁴ and surface enrichment of Mo has been observed previously by electron microscopy,³⁷ XPS³⁸ and low energy ion scattering on the outermost surface layer.¹⁶ In this study, the use of the dicarboxylate chelating agents seems to promote the formation of a Mo rich surface layer compared to the standard coprecipitated material (1:2.2-C).

BET surface areas were also obtained for the samples (Table 2). The oxalate route gave the highest surface areas which were found to increase with Mo content for both the sol-gel preparation methods. Previous studies using standard coprecipitation found that the surface area decreases with molybdenum content due to the increase in low surface area MoO_3 formed.^{13,16,39} However, this rise in surface area with increased molybdenum content was also observed by Soares *et al.* using a sol-gel method and similar Fe:Mo ratios,⁸ which was attributed to the sponge-like MoO_3 morphologies observed.

The iron molybdate materials were then tested as catalysts for the partial oxidation of methanol to formaldehyde (Fig. 3). The conversion of methanol over the catalysts can be linked to the method of preparation, with catalysts prepared using the sol-gel chemistry outperforming the coprecipitated catalysts, and the oxalate route showing better performance than those prepared using the malonate route. The activity is similar for all the sol-gel synthesised materials apart from the 1:1.5-M, which shows comparable activity to the coprecipitated catalyst (1:2.2-C) and does not reach full conversion until $\sim 380^\circ\text{C}$. This is perhaps not surprising as the 1:1.5-M material has the lowest surface area and the lowest Fe:Mo surface ratio (Table 2), both of which are key indicators of performance.

For the catalysts synthesised using malonic acid the selectivity to formaldehyde was high, with $>95\%$ observed across a broad temperature and conversion range. As the temperature was increased the selectivity to formaldehyde decreased as a greater amount of CO_x was produced. The catalysts prepared with oxalic acid showed a lower selectivity

Table 2 BET surface area and XPS surface composition of the iron molybdate materials after calcination

Sample	Surface area/ $\text{m}^2 \text{g}^{-1}$	Fe:Mo surface ratio
1:2.2-C	3.9	1:3.0
1:1.5-O	4.4	1:2.2
1:2.2-O	4.6	1:3.6
1:3.0-O	7.0	1:3.6
1:1.5-M	2.2	1:2.0
1:2.2-M	3.1	1:3.3
1:3.0-M	3.4	1:3.3



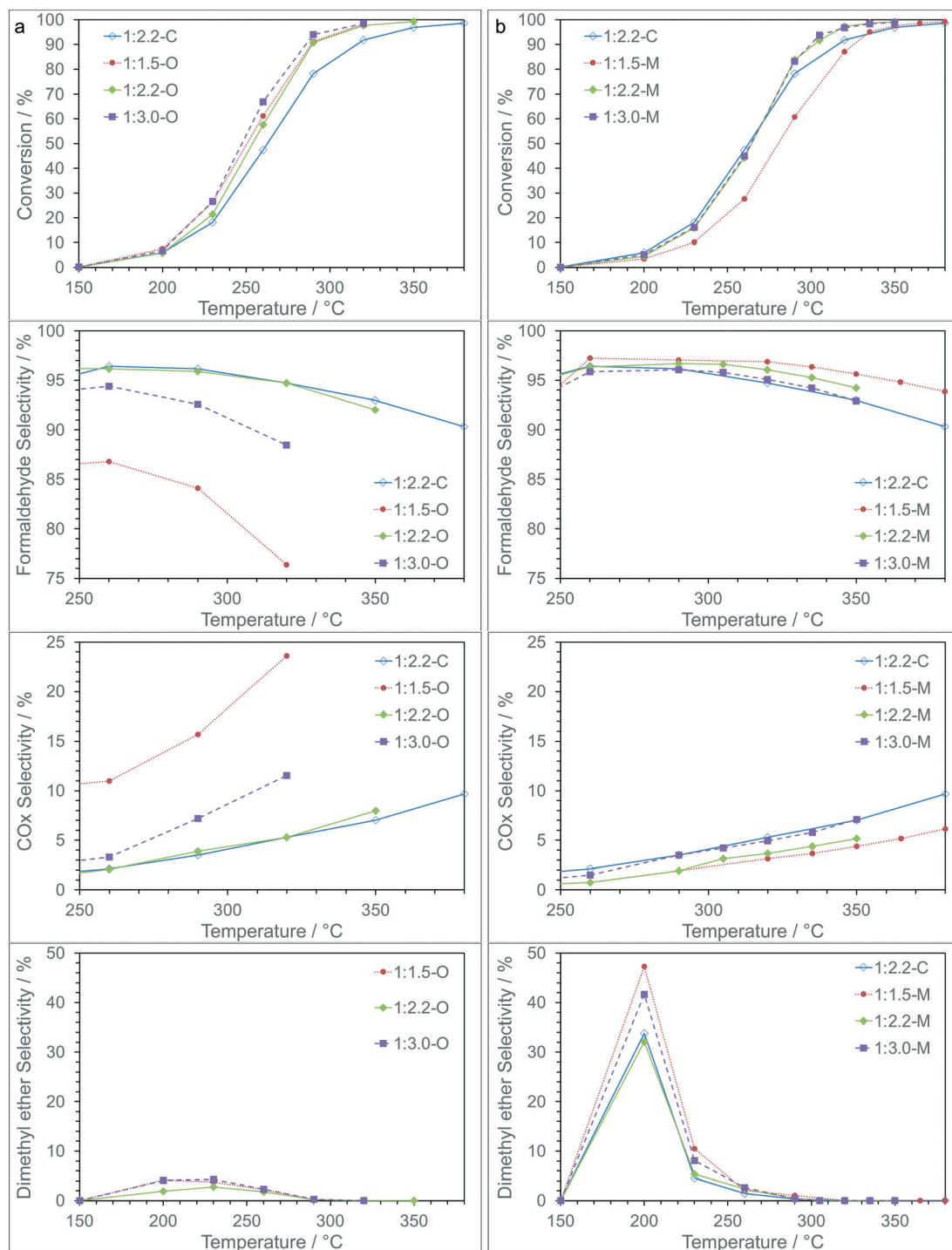


Fig. 3 Catalyst testing for methanol partial oxidation for materials prepared using (a) oxalic acid and (b) malonic acid compared with a standard coprecipitated catalyst. Reaction conditions: 0.3 g of catalyst, flow rate = 60 ml min⁻¹ (MeOH : O₂ : He = 5 : 10 : 85).

to formaldehyde, with the 1:1.5-O and 1:3.0-O in particular showing lower selectivity at fairly modest temperatures due to increased CO_x formation. The 1:2.2-O material showed very similar behavior to the coprecipitated iron molybdate (1:2.2-C). At low conversion dimethyl ether (DME) was observed for all catalysts (Table S1†), with higher selectivity observed over 1:2.2-C and the materials synthesized using

malonic acid. As conversion increased, DME selectivity reduced to >1% above 300 °C.

Based on these results the malonate route shows clear advantages over the oxalate and coprecipitation routes with higher yields of formaldehyde over an extended temperature range and reaction rates (Fig. S2†) suggesting an increase in stability of these materials.



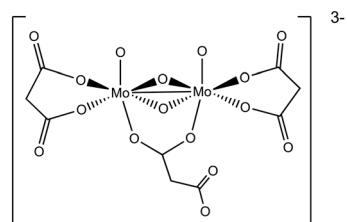
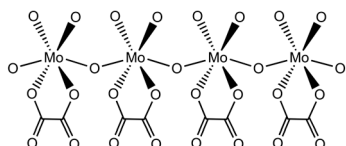
a The structure of $[\text{Mo}^{\text{VI}}_2\text{O}_4(\text{CH}_2\text{C}_2\text{O}_4)_3]^{3-}$ b The polymeric structure of $[\text{Mo}^{\text{VI}}\text{O}_3(\text{C}_2\text{O}_4)]^{2-}$

Fig. 4 (a) The structure of $[\text{Mo}^{\text{VI}}_2\text{O}_4(\text{CH}_2\text{C}_2\text{O}_4)_3]^{3-}$. (b) The polymeric structure of $[\text{Mo}^{\text{VI}}\text{O}_3(\text{C}_2\text{O}_4)]^{2-}$.

The precursors synthesised using the sol-gel routes were further characterised using XPS and thermogravimetric analysis (TGA) to understand the relationship between the catalyst performance and the species present in the precursors. The oxalate and malonate precursors can exist in different forms, depending on the chelating agent and the oxidation state of the iron and molybdenum.

The conjugate bases of both oxalic and malonic acid are well-established chelating ligands for metal cations that have been applied for the dissolution of metals in applications such as removing unwanted metal contaminants^{40,41} and selectively leaching of metals from raw materials.⁴² Both the organic acids chelate to molybdenum in a similar way, forming polymeric structures depending on the Mo oxidation state.^{43,44} Mo(v) dimers form which complex to give $[\text{Mo}_2\text{O}_4(\text{C}_2\text{O}_4)_3]^{3-}$ or $[\text{Mo}_2\text{O}_4(\text{CH}_2\text{C}_2\text{O}_4)_3]^{3-}$ which are linked together by bridging oxalate or malonate ions (Fig. 4),⁴⁴ whereas, Mo(vi) forms $[\text{MoO}_3(\text{C}_2\text{O}_4)_2]^{2-}$ or $[\text{MoO}_3(\text{CH}_2\text{C}_2\text{O}_4)_2]^{2-}$ anions, which are bound together in polymeric species *via* Mo–O–Mo linear chains⁴³ (Fig. 4b).

However, the acids show different behaviour in the presence of iron. It has been proposed that the longer chain length allows the malonic acid to act as a bidentate ligand,^{45–47} leading to isolated iron centres rather than the polymeric chains found with oxalic acid. This suggests that the different chain length of the diacid chelating agent could play a role in determining the homogeneity of the precursors, limiting the formation of iron rich regions within the catalysts. For both acids the iron complexes polymerise to form infinite chain structures as shown in Fig. 5a. For the Fe(III) chains the additional ligand can lead to crosslinking of these chains (Fig. 5b).

However, for malonic acid the longer carbon chain allows the formation of $[\text{Fe}^{\text{III}}(\text{CH}_2\text{C}_2\text{O}_4)_3]^{3-}$ with the three bidentate

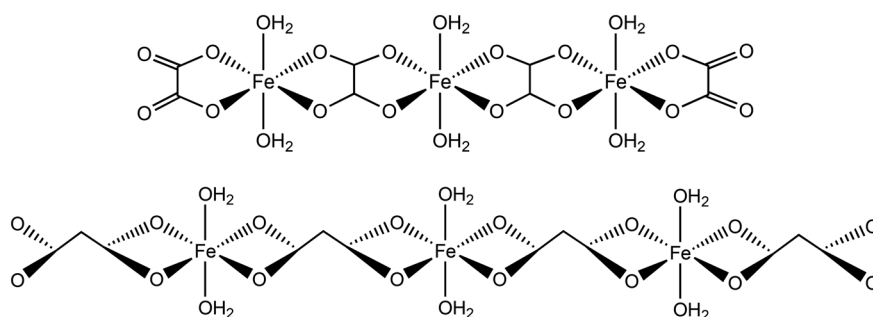
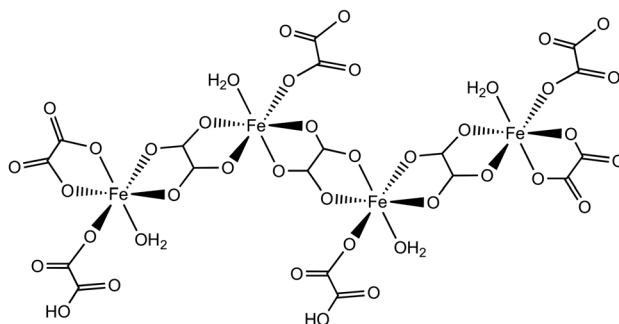
a The polymeric structure of $\text{Fe}^{\text{III}}(\text{C}_2\text{O}_4) \cdot 2\text{H}_2\text{O}$ and $\text{Fe}^{\text{III}}(\text{CH}_2\text{C}_2\text{O}_4) \cdot 2\text{H}_2\text{O}$.b The crosslinked polymeric structure of $\text{Fe}^{\text{III}}(\text{C}_2\text{O}_4)_3 \cdot \text{H}_2\text{O}$.

Fig. 5 (a) The polymeric structure of $\text{Fe}^{\text{III}}(\text{C}_2\text{O}_4) \cdot 2\text{H}_2\text{O}$ and $\text{Fe}^{\text{III}}(\text{CH}_2\text{C}_2\text{O}_4) \cdot 2\text{H}_2\text{O}$. (b) The crosslinked polymeric structure of $\text{Fe}^{\text{III}}(\text{C}_2\text{O}_4)_3 \cdot \text{H}_2\text{O}$.



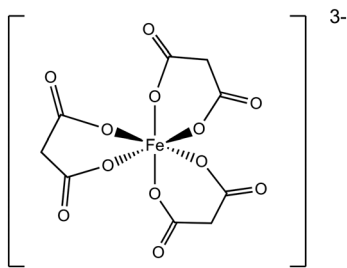


Fig. 6 The structure of $[\text{Fe}^{\text{III}}(\text{CH}_2\text{C}_2\text{O}_4)_3]^{3-}$.

ligands chelating to one Fe^{III} centre to form discrete anions that do not polymerise (Fig. 6).

Therefore, the oxidation state of the cations can have a big influence on the composition and structure of the precursors and the homogeneity of the final oxide materials, depending on whether polymeric chains or discrete anions are present in the samples. XPS analysis of molybdenum could therefore give important information as to the species formed during the melting that occurs during the initial heat treatment step as Mo^{IV} , Mo^{V} and Mo^{VI} have clearly defined binding energies for $\text{Mo } 3d_{5/2}$ – $\text{Mo } 3d_{3/2}$ doublets of 229.9–233.0, 231.2–234.3 and 232.5–235.6 respectively.⁴⁸

From the XPS analysis (Table 3, Fig. S3†) it is clear that the oxalate and malonate samples had a large amount of Mo^{V} present in the precursor, which is likely to lead to a higher proportion of isolated Mo ions rather than polymeric chains. This in turn could lead to a higher degree of mixing between iron and molybdenum in the precursors, limiting the formation of iron oxides which were found to be higher from the XRD analysis (Table 1). Conversely, standard coprecipitation methods (1:2.2-C) resulted in exclusively Mo^{VI} species due to the formation of $\text{Fe}_2(\text{MoO}_4)_3$ and MoO_3 prior to calcination.

TGA of the materials adds further evidence for the homogeneous mixing of the iron and molybdenum shown in the XPS results (Fig. 7).

For the oxalate samples, two large regions of mass loss were observed. The initial loss at 160–200 °C corresponded to iron oxalate decomposition in air as observed previously.^{49,50} The second major feature was between 230–270 °C which has been assigned to the decomposition of oxomolybdate oxalate which is again supported by literature examples.^{51–53} In addition to the major features, mass loss between 325–350 °C was assigned

to the decomposition of ammonium molybdate species to form crystalline MoO_3 ,⁵⁴ with the minor change observed at 350–400 °C was attributed to the formation of $\text{Fe}_2(\text{MoO}_4)_3$ during a solid phase reaction between Fe_2O_3 and MoO_3 species.⁵⁵

In all materials there was an overlap between the iron and molybdenum oxalate decompositions between 200–230 °C, which could be due to be secondary catalytic decarboxylation of oxomolybdate oxalate.⁵⁶ Majumdar *et al.* studied the thermal decomposition of $\text{FeC}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$ and $\text{ZnC}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$ mixtures and observed a significant decrease in zinc oxalate decomposition temperatures. This was ascribed to the exothermic catalytic oxidation of CO to CO_2 over Fe_3O_4 and the oxidation of Fe_3O_4 to Fe_2O_3 .⁴⁹ This caused localised high temperatures around these iron centres causing the activation of neighbouring carboxylate species. Increases in this overlapping region can be an indication of mixing of the oxalates and possibly the overall homogeneity of the material produced post calcination.

For the malonate samples, TGA shows similar behaviour to the oxalate samples, with decompositions assigned to molybdenum malonate at 176–214 °C (ref. 57) and iron malonate at 220–358 °C (ref. 46) respectively. As for the oxalate samples, there is an overlapping region in between these features, which becomes more prevalent with increasing molybdenum content and can be seen as a defined peak in the 1:2.2-M and 1:3.0-M precursors. If this is attributed to the exothermic decomposition of the molybdenum malonate caused the activation of neighbouring malonates, the increase in the mass loss in this region is a clear indicator of an increase in the intimate mixing between the two phases. This correlates with the XPS analysis showing a higher amount of Mo^{V} present in the sample, which is less likely to form long polymeric chains and can lead to a better dispersion of Fe and Mo in the precursors.

Unlike isolated iron oxalate, which was linked to Fe_2O_3 production, decomposition of iron malonate has been shown to produce $\gamma\text{-Fe}_2\text{O}_3$ nanoparticles by Stefanescu and Stefanescu.⁴⁵ This might have aided the formation of more homogeneous distribution by preventing the formation of large isolated areas of iron enrichment observed on the surface, although larger volumes of isolated iron malonate could be detrimental if not dispersed leading to localised high iron concentrations.

Conclusions

The synthesis of mixed metal oxide catalysts has been widely studied with a key focus on improving the homogeneity of the materials. In this study we have investigated the use of malonic and oxalic acids as chelating agents in the sol gel synthesis of iron molybdate. The metal malonate and oxalate precursors were decomposed and the resultant oxide catalysts were found to give improved activity over coprecipitated catalysts for the selective oxidation of methanol to formaldehyde. The highest yields of formaldehyde were observed for the catalysts synthesised using malonic acid with higher than stoichiometric molybdenum content (1:2.2-M and 1:3.0-M). This is proposed to

Table 3 XPS surface composition of the iron molybdate precursors before calcination

Sample	$\text{Mo}^{\text{V}} : \text{Mo}^{\text{VI}}$ surface ratio
1:2.2-C	0 : 1
1:1.5-O	1 : 1.2
1:2.2-O	1 : 1.1
1:3.0-O	1 : 2.1
1:1.5-M	1 : 2.7
1:2.2-M	1 : 1.5
1:3.0-M	1 : 2.4



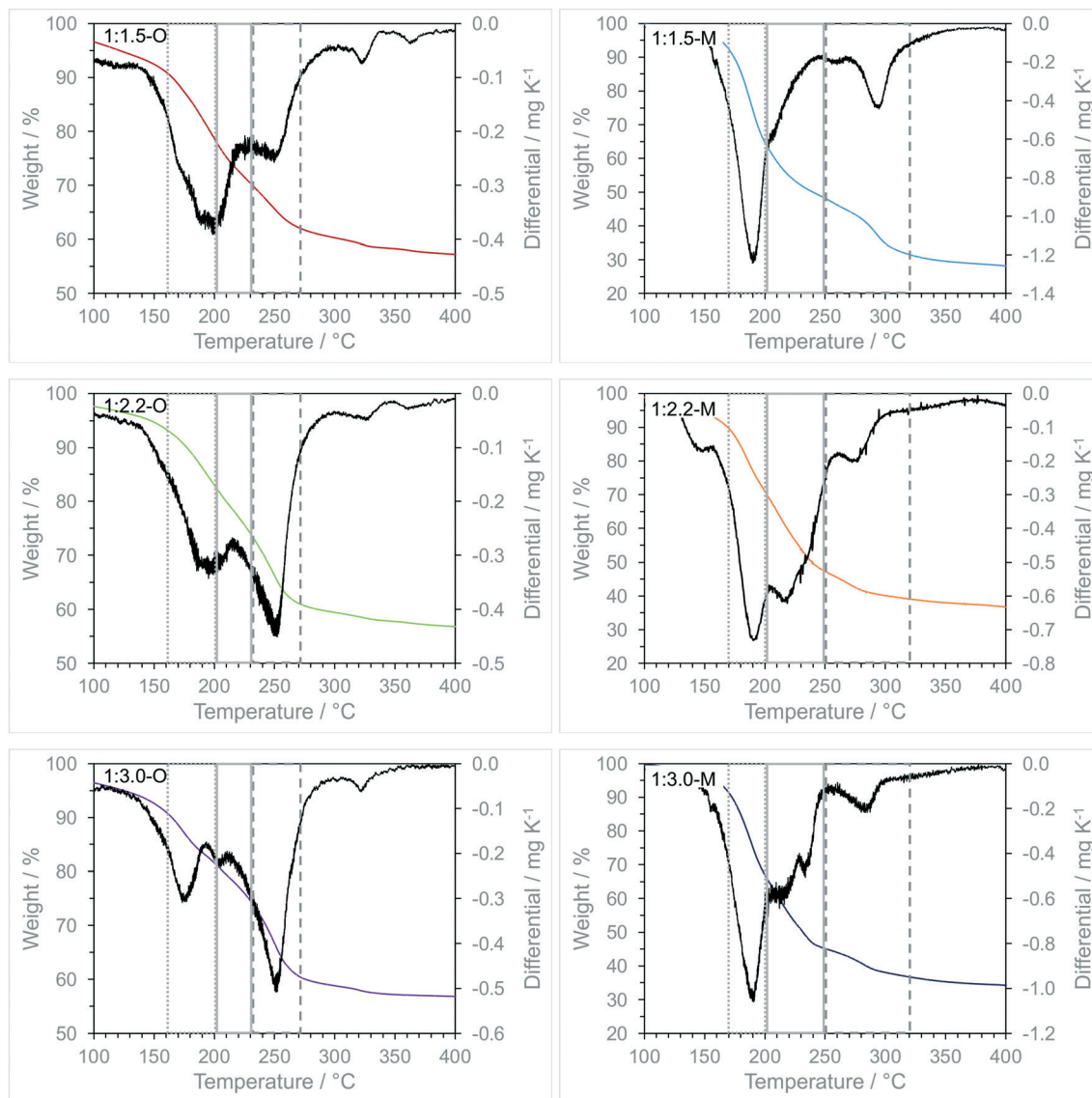


Fig. 7 Thermogravimetric analysis of iron molybdate precursors synthesised using oxalic acid and malonic acid.

be due to the longer chain length of malonic acid allowing it to act as a bidentate ligand, resulting in more isolated iron and molybdenum centres than in the polymeric chains formed with oxalic acid. These isolated metal centres can form a more homogenous material, reducing the amount of iron rich centres that have been attributed to reduced selectivity for iron molybdate catalysts.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 E. Cao and A. Gavrilidis, *Catal. Today*, 2005, **100**, 154–163.
- 2 W.-L. Dai, Q. Liu, Y. Cao and J.-F. Deng, *Appl. Catal.*, 1998, **175**, 83–88.
- 3 L. Lefferts, J. G. Van Ommen and J. R. H. Ross, *Appl. Catal.*, 1986, **23**, 385–402.
- 4 M. Qian, M. A. Liauw and G. Emig, *Appl. Catal.*, A, 2003, **238**, 211–222.
- 5 A. N. Pestryakov, N. E. Bogdanchikova and A. K. Gericke, *Catal. Today*, 2004, **91**, 49–52.
- 6 A. Andersson, M. Hernelind and O. Augustsson, *Catal. Today*, 2006, **112**, 40–44.
- 7 K. I. Ivanov and D. Y. Dimirtov, *Catal. Today*, 2010, **154**, 250–255.
- 8 A. P. V. Soares, M. F. Portela, A. Kiennemann and L. Hilaire, *Chem. Eng. Sci.*, 2003, **58**, 1315–1322.



- 9 V. Diakov, B. Blackwell and A. Varma, *Chem. Eng. Sci.*, 2002, **57**, 1563–1569.
- 10 G. Reuss, W. Disteldorf, O. Grundler and A. Hilt, *Ullmann's Encyclopedia of Industry Chemistry*, VCH, Weinheim, Germany, 5th edn, 1992, vol. A11.
- 11 A. B. Stiles and T. A. Koch, *Catalysis Manufacture*, Marcel Dekker, New York, 2nd edn, 1995.
- 12 H. Adkins and W. R. J. Peterson, *J. Am. Chem. Soc.*, 1931, **53**, 1512–1520.
- 13 A. P. V. Soares, M. F. Portela, A. Kiennemann, L. Hilaire and J. M. M. Millet, *Appl. Catal., A*, 2001, **206**, 221–229.
- 14 E. Söderhjelm, M. P. House, N. Cruise, J. Holmberg, M. Bowker, J.-O. Bovin and A. Andersson, *Top. Catal.*, 2008, **50**, 145–155.
- 15 M. Badlani and I. E. Wachs, *Catal. Lett.*, 2001, **75**, 137–149.
- 16 K. Routray, W. Zhou, C. J. Kiely, W. Grünert and I. E. Wachs, *J. Catal.*, 2010, **275**, 84–98.
- 17 B. Delmon and G. F. Froment, *Catal. Rev.: Sci. Eng.*, 1996, **38**, 69–100.
- 18 I. E. Wachs and K. Routray, *ACS Catal.*, 2012, **2**, 1235–1246.
- 19 V. Diakov, D. Lafarga and A. Varma, *Catal. Today*, 2001, **67**, 159–167.
- 20 A. P. V. Soares, M. F. Portela, A. Kiennemann and J. M. M. Millet, *React. Kinet. Catal. Lett.*, 2002, **75**, 13–20.
- 21 K. V. Raun, J. Johannessen, K. McCormack, C. C. Appel, S. Baier, M. Thorhauge, M. Høj and A. D. Jensen, *Chem. Eng. J.*, 2019, **361**, 1285–1295.
- 22 M. Carbucicchio and F. Trifirò, *J. Catal.*, 1976, **45**, 77–85.
- 23 T. H. Kim, B. Ramachandra, J. S. Choi, M. B. Saidutta, K. Y. Choo, S. D. Song and Y. W. Rhee, *Catal. Lett.*, 2004, **98**, 161–165.
- 24 Y. Huang, L. Cong, J. Yu, P. Eloy and P. Ruiz, *J. Mol. Catal. A: Chem.*, 2009, **302**, 48–53.
- 25 L. Dong, K. Chen and Y. Chen, *J. Solid State Chem.*, 1997, **129**, 30–36.
- 26 A. M. Beale, S. D. M. Jacques, E. Sacaliuc-Parvalescu, M. G. O'Brien, P. Barnes and B. M. Weckhuysen, *Appl. Catal., A*, 2009, **363**, 143–152.
- 27 Y. Ding, S.-H. Yu, C. Liu and Z.-A. Zang, *Chem. – Eur. J.*, 2007, **13**, 746–753.
- 28 G. Jin, W. Weng, Z. Lin, N. F. Dummer, S. H. Taylor, C. J. Kiely, J. K. Bartley and G. J. Hutchings, *J. Catal.*, 2012, **296**, 55–64.
- 29 M. Bowker, M. House, A. Alshehri, C. Brookes, E. K. Gibson and P. P. Wells, *Catal., Struct. React.*, 2015, **1**, 95–100.
- 30 C. Brookes, P. P. Wells, G. Cibir, N. Dimitratos, W. Jones and M. Bowker, *ACS Catal.*, 2014, **4**, 243–250.
- 31 B. L. Cushing, V. L. Kolesnichenko and C. J. O'Connor, *Chem. Rev.*, 2004, **104**, 3893–3946.
- 32 A. E. Danks, S. R. Hall and Z. Schnepf, *Mater. Horiz.*, 2016, **3**, 91–112.
- 33 A. P. V. Soares, M. F. Portela and A. Kiennemann, *Stud. Surf. Sci. Catal.*, 1997, **110**, 807–816.
- 34 H. Oudghiri-Hassani, *Catal. Commun.*, 2015, **60**, 19–22.
- 35 B. R. Yeo, G. J. F. Pudge, K. G. Bugler, A. V. Rushby, S. Kondrat, J. K. Bartley, S. Golunski, S. H. Taylor, E. Gibson, P. P. Wells, C. Brookes, M. Bowker and G. J. Hutchings, *Surf. Sci.*, 2016, **648**, 163–169.
- 36 A. Katelnikovas, J. Barkauskas, F. Ivanauskas, A. Beganskiene and A. Kareiva, *J. Sol-Gel Sci. Technol.*, 2007, **41**, 193–201.
- 37 M. Bowker, R. Holroyd, M. House, R. Bracey, C. Bamroongwongdee, M. Shannon and A. Carley, *Top. Catal.*, 2008, **48**, 158–165.
- 38 B. I. Popov, A. V. Pashis and L. N. Shkuratova, *React. Kinet. Catal. Lett.*, 1986, **30**, 129–135.
- 39 M. P. House, A. F. Carley, R. Echeverria-Valda and M. Bowker, *J. Phys. Chem. C*, 2008, **112**, 4333–4341.
- 40 F. Vegliò, B. Passariello and C. Abbruzzese, *Ind. Eng. Chem. Res.*, 1999, **38**, 4443–4448.
- 41 C. Ash, V. Tejnecký, L. Borůvka and O. Drábek, *J. Contam. Hydrol.*, 2016, **187**, 18–30.
- 42 P. Hu, Y. Zhang, T. Liu, J. Huang, Y. Yuan and Q. Zheng, *J. Ind. Eng. Chem.*, 2017, **45**, 241–247.
- 43 M. Cindrić, N. Strukan, V. Vrdoljak, M. Devčić, Z. Vekšli and B. Kamenar, *Inorg. Chim. Acta*, 2000, **304**, 260–267.
- 44 B. Modéc, J. V. Brenčić and J. Koller, *Eur. J. Inorg. Chem.*, 2004, **2004**, 1611–1620.
- 45 O. Stefanescu and M. Stefanescu, *J. Organomet. Chem.*, 2013, **740**, 50–55.
- 46 M. M. Rahman, V. A. Mukhedkar, A. Venkataraman, A. K. Nikumbh, S. B. Kulkarni and A. J. Mukhedkar, *Thermochim. Acta*, 1988, **125**, 173–190.
- 47 D. Xiao, Y. Guo, X. Lou, C. Fang, Z. Wang and J. Liu, *Chemosphere*, 2014, **103**, 354–358.
- 48 J.-G. Choi and L. T. Thompson, *Appl. Surf. Sci.*, 1996, **93**, 143–149.
- 49 B. Boyanov, D. Khadzhiev and V. Vasilev, *Thermochim. Acta*, 1985, **93**, 89–92.
- 50 E.-H. M. Diefallah, M. A. Mousa, A. A. El-Bellihi, E.-H. El-Mossalamy, G. A. El-Sayed and M. A. Gabal, *J. Anal. Appl. Pyrolysis*, 2002, **62**, 205–214.
- 51 M. I. Diaz-Guemes, A. S. Bhatti and D. Dollimore, *Thermochim. Acta*, 1986, **106**, 125–132.
- 52 J. Gopalakrishnan, B. Viswanathan and V. Srinivasan, *J. Inorg. Nucl. Chem.*, 1970, **32**, 2565–2568.
- 53 S. P. Goel and P. N. Mehrotra, *Thermochim. Acta*, 1983, **70**, 201–209.
- 54 A. O. Kostynyuk, F. Gutenuar, A. N. Kalashnikova, Y. V. Kalashnikov and N. V. Nikolenko, *Kinet. Catal.*, 2014, **55**, 649–655.
- 55 M. Bowker, R. Holroyd, A. Elliott, P. Morrall, A. Alouche, C. Entwistle and A. Toerncrona, *Catal. Lett.*, 2002, **83**, 165–176.
- 56 R. Majumdar, P. Sarkar, U. Ray and M. R. Mukhopadhyay, *Thermochim. Acta*, 1999, **335**, 43–53.
- 57 M. E. Brown, T. T. Bhengu and D. K. Sanyal, *Thermochim. Acta*, 1994, **242**, 141–152.

