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Evolution of Mo species and ZSM-5 microstructure with temperature and its impact on methane dehydroaromatisation activity†

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Mo-ZSM-5 is a well-studied catalyst for its ability to convert methane to aromatics and hydrogen. Since the process is thermodynamically 'uphill', high temperatures are necessary to achieve acceptable product yields. However, these temperatures have been shown to negatively impact the stability of the zeolite, with the formation of aluminum molybdates being proposed as particularly responsible for framework collapse. Here we use X-ray absorption and optical Raman spectroscopy as well as X-ray diffraction to characterize the initial Mo species present after introduction to ZSM-5, thermal activation and after structural collapse. The results demonstrate that the formation of mononuclear Mo⁶⁺ species coordinated to oxygen is retained despite increasingly high temperature treatment and which is able to perform methane dehydroaromatisation although with decreasing efficiency as zeolite porosity is lost.

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Introduction

Methane dehydroaromatisation (MDA) is a promising reaction to upgrade methane directly into aromatics and hydrogen.^{1,2} A variety of catalysts based on metal-ion exchanged zeolites have been explored for the MDA reaction. Among the tested zeolite topologies (for example ZSM-5, MCM-22, and MCM-49) and metal cations, the combination of Mo supported on 10-membered ring zeolites (ZSM-5 and MCM-22) remains one of the most active catalysts to date.³ Mo-containing zeolite catalysts are normally prepared by mixing the solid oxides of Mo compounds with zeolite supports followed by calcination at high temperatures, which leads to the decomposition and distribution of the MoO_x species into the pores of the zeolite. Extensive research has been carried out to resolve the nature and location of this dispersed MoO_x species.^{4–6} Although there is no consensus, a recent study has shown that after calcination, isolated Mo-oxo species anchored at the channel intersection are

the dominant species.⁷ In general, it is accepted that after migration into the zeolite micropores, the MoO_x species anchor at the Brønsted acid sites, releasing water molecules.⁸ This reaction also distorts or even extracts framework Al and, in some cases, gives rise to the formation of Al₂(MoO₄)₃ species.⁹

Despite all the promise, Mo/H-ZSM-5 suffers from rapid deactivation due to both coke formation and the migration of the active Mo species to the outer parts of the zeolite crystals.^{10,11} Overcoming this problem has been the focus of many studies, attempting to: (1) modify and optimise the catalyst zeolite support and the active metal cations^{12,13} and, (2) regenerate the coked catalyst in a reductive or oxidative atmosphere.¹⁴ It has been found that at typical MDA reaction temperatures Mo/H-ZSM-5 is not stable enough under the oxidative conditions needed for regeneration, leading to the sublimation of the MoO_x species and more importantly, to irreversible alteration to the zeolite framework often resulting in the collapse of the zeolite.⁸

In the current study, we focus on the thermal stability of the Mo/H-ZSM-5 catalysts with an emphasis on the evolution of the chemical species as the zeolite undergoes structural collapse. The thermal stability of the catalysts was examined firstly by means of differential scanning calorimetry (DSC). The phase and morphological evolution of the catalysts during the course of the collapse was further investigated by powder X-ray diffraction (PXRD) and scanning electron microscopy (SEM), while changes in Mo speciation of Mo cation were characterised by X-ray absorption fine structure (XAFS). It is thought that modality of zeolite framework collapse is intimately related to the microscopic framework Al distribution.

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Experimental section

Catalyst preparation

NH₄-ZSM-5 was supplied by Zeolyst International (CBV3024E, Si:Al = 15). SSZ-13 (Si:Al = 15) was synthesized in fluoride media following the reported methods.¹⁵ H-form zeolites were obtained by calcination at 550 °C (ZSM-5) and 620 °C (SSZ-13). Mo/H-zeolites were prepared *via* the established solid-state ion-exchange method (SSIE).¹⁶ Mo/H-ZSM-5 samples are designated with the name Mo_x-Z5, where *x* denotes the weight percentage of the Mo loading. Samples were calcined to the point of collapse and beyond in static air using a ramp rate of 10 °C min⁻¹.

Catalyst characterization

PXRD patterns were recorded using a Rigaku MiniFlex 600 X-ray diffractometer fitted with Cu K α radiation. The Kerr-gated Raman spectra were recorded at the ULTRA facility at the Central Laser Facility, using a 400 nm wavelength excitation source. The data were calibrated to the spectra of toluene-impregnated H-ZSM-5. Surface characterization was carried out using N₂ physisorption at 77.3 K on a Quadrasorb EVO QDS-30. Samples were outgassed at 350 °C overnight. The Brunauer–Emmett–Teller (BET) method was used for surface area analysis. The *t*-plot method was used for micropore volume analysis. Solid-state ²⁷Al MAS-NMR were carried out on a standard Bruker 4 mm double-resonance MAS probe at the UCL Department of Chemistry, and samples were fully hydrated prior to each measurement. Differential scanning calorimetry (DSC) was recorded on a NETZSCH STA 449 F3 Jupiter at the UCL Department of Chemistry. Samples were heated up to 1400 °C at 10 °C min⁻¹ under a 30 mL min⁻¹ flow of He. An empty fused alumina sample pan was used as a reference. Scanning electron micrographs (SEM) and energy dispersive X-ray spectroscopy (EDX) analysis were carried out with a JEOL JSM-6610LV at RCaH and JEOL6700F FEG at the UCL Department of Chemistry. XRF elemental distribution maps were recorded at beamline 2-ID-E at the Advanced Photon Source.¹⁷

X-ray absorption fine structure

Mo K-edge XAFS studies were performed at the European Synchrotron Radiation Facility (ESRF), France, on BM26A Beamline and the Diamond Light Source (DLS), UK, on beamline B18. The measurements were performed in transmission mode using ion chambers with a fast-scanning Si(111) double-crystal monochromator. Each scan took 5 min to acquire at B18 and 10 min at BM26A. Three scans were taken for each sample and averaged. XAFS data reduction and analysis were performed using the Demeter package over a *k*-range of 2–10 Å and *R*-range of 0.5–3 Å.¹⁸ Linear combination fitting (LCF) of XANES spectra was performed in an energy range of –20 to +70 eV at Mo K-edge. Morlet wavelet transforms are calculated by the Fortran version of the HAMA software developed by Funke *et al.*¹⁹

MDA catalytic activity tests

The catalytic testing was carried out by placing 0.6 g of catalyst (150–425 μ m sieved fractions) in a tubular quartz reactor with an internal diameter of 0.7 mm. The catalysts were secured in the isothermal zone of the oven by quartz wool. A total gas flow of 30 mL min⁻¹ (GHSV = 1500 h⁻¹) was fed to the reactor by a set of mass flow controllers. The samples were first calcined to 700 °C at 5 °C min⁻¹ under a 20% O₂ in He flow. Then still at 700 °C the reactor was flushed by a stream of pure Ar for 20 min, before switching to MDA gas composition (50% CH₄ in Ar). The product formation was analyzed by an online OmniStar GSD 320O1 mass spectrometer.

Results and discussion

Catalyst characterization

The solid-state Mo-exchanged zeolite materials used in this work are typical for relevant MDA catalyst.

2 and 4 wt% Mo/H-ZSM-5 were synthesized from high-temperature calcination of a ground mixture of H-ZSM-5 and MoO₃. In Fig. 1(b) is shown the PXRD of the calcined Mo/H-ZSM-5 against the physical mixture. Reflections of crystalline MoO₃ in the after high temperature treatment fall below the

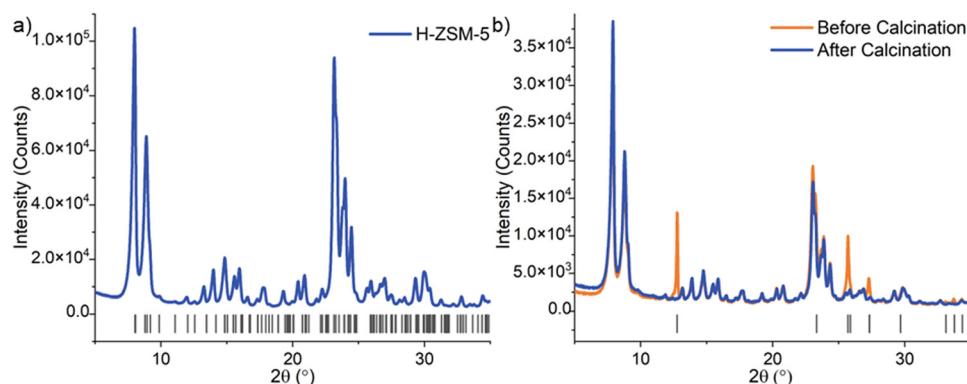


Fig. 1 PXRD patterns of zeolite precursors H-ZSM-5 with indexing lines of the references ZSM-5 structures (a) and Mo4-Z5 before (blue line) and after (orange line) calcination with indexing lines of the crystalline MoO₃ phase are shown below the patterns (b).



detection level in sample Mo4-Z5. It is known that during the thermal treatment, Mo species migrate into the pores of zeolite and ion exchange at the Brønsted acid sites. A change in peak position, width, and intensity pertaining to the MFI framework is not observed in the pattern of Mo4-Z5, indicating the zeolite structure is retained with no observable loss in crystallinity. No other crystalline Mo phase such as $\text{Al}_2(\text{MoO}_4)_3$ was detected. Mo loading obtained from chemical analysis and texture properties of the Mo/H-ZSM-5 samples are summarized in Table S1 (ESI[†]). The introduction of Mo leads to a decrease in BET micropore area and volume (up to 16% and 18% respectively) while the severity in reduction directly relates to the Mo loading. This observed decrease is possibly caused by the Mo introduced partially hindering diffusion throughout the micropores and/or a loss of microporous structure due to dealumination,¹⁴ which is also evident by the increase in amount of octahedral extra-framework Al (EFAL) and $\text{Al}_2(\text{MoO}_4)_3$ species observed by ²⁷Al MAS-NMR (Fig. 2).

The interaction between Mo and the zeolite framework can lead to extraction of tetrahedral framework Al,²⁰ in some cases leading to an irreversible framework degradation and the formation of mesopores – a process accompanied by the formation of aluminium molybdate.⁸ To gain insight into Al speciation, Mo-exchanged zeolite catalysts were characterized by ²⁷Al MAS-NMR. The introduction of the Mo ion modifies substantially the NMR spectra of ZSM-5 zeolite (Fig. 2). The extraction of framework Al is undoubtedly evidenced by a significant decrease in the tetrahedral framework Al signals at 54 ppm. This is accompanied by an increase in the peak width, likely caused by an increased quadrupolar broadening effect. This framework dealumination becomes more noticeable with increasing molybdenum loading, while a shift of 2 ppm towards a higher field for the framework Al peak is also observed. Tessonnier *et al.* reported that peak shift is related to the distortion of framework AlO_4 units and alteration of the T-O-T angle.⁵ Furthermore, an increase in signal intensity

around 0 ppm, corresponding to an increase in the amount of octahedral extra-framework Al (EFAL) was observed, along with a new signal at 14 ppm. Kosinov *et al.* assigned the signals at 14 ppm and –14 ppm to bulk and well-dispersed $\text{Al}_2(\text{MoO}_4)_3$ respectively,⁸ whereas Ma *et al.* argued that signal at –14 ppm represents the $\text{Al}_2(\text{MoO}_4)_3$ species, and it shifts to 14 ppm upon hydration.⁹ The bulk $\text{Al}_2(\text{MoO}_4)_3$ reference measured here shows an intense signal at –13.8 ppm (Fig. S2, ESI[†]). Given that no bulk crystalline $\text{Al}_2(\text{MoO}_4)_3$ phases are detected by PXRD and the Mo samples were hydrated prior to measurement, the signal at 14 ppm likely originates from hydrated $\text{Al}_2(\text{MoO}_4)_3$. Signal intensity from both octahedral EFAL and $\text{Al}_2(\text{MoO}_4)_3$ surge with increasing Mo loading. Therefore, it can be concluded that Mo ion-exchange leads to framework distortion, significant framework dealumination and formation of octahedral EFAL and $\text{Al}_2(\text{MoO}_4)_3$ species. It is interesting to note that these changes have not manifested into long-range structural changes in the zeolite framework as evidenced by the unchanging PXRD patterns (see Fig. 1).

The structure of dispersed Mo oxide species was further investigated by Kerr-gated Raman spectroscopy, as shown in Fig. 3.²¹ The spectrum of the H-form ZSM-5 zeolite is characterized by two broad bands, one centered around 400 cm^{-1} with shoulders on either side at 300 and 470 cm^{-1} , and the second at 812 cm^{-1} , due to the vibrations of the MFI framework.^{22,23} The spectrum of the MoO_3 and H-ZSM-5 physical mixture, in contrast, is dominated by strong Raman signals characteristic of MoO_3 ; *i.e.* 293 cm^{-1} and 364 cm^{-1} (M=O bending), 473 cm^{-1} and 668 cm^{-1} edge-sharing (Mo-O(3) stretching), 819 cm^{-1} (Mo-O-Mo asymmetric stretching), and 995 cm^{-1} (Mo=O asymmetric stretching).²⁴

After SSIE and calcination (see Fig. 3), Mo2-Z5 and Mo4-Z5, the Raman bands from MoO_3 are clearly diminished, whereas new Raman bands emerge at 858 cm^{-1} and 944 cm^{-1} . The 944 cm^{-1} band can be readily attributed to the Mo=O stretching mode from distorted octahedral and is often attributed to isolated monomeric species,⁶ while the 858 cm^{-1} band is due to

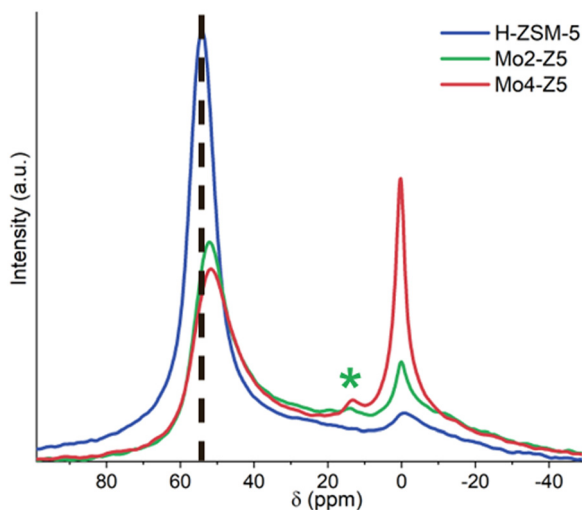


Fig. 2 ²⁷Al MAS NMR spectra of H-ZSM-5, Mo2-Z5 and Mo4-Z5 (the * indicates the location of the chemical shift for $\text{Al}_2(\text{MoO}_4)_3$ at 14 ppm).

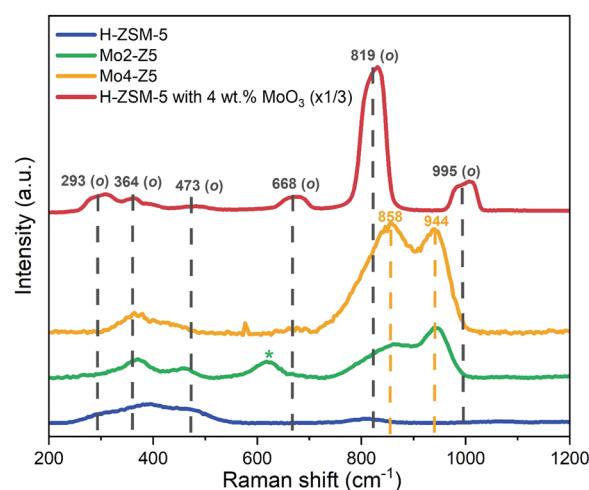


Fig. 3 Kerr-gated Raman spectra for H-ZSM-5 (blue), Mo2-Z5 (green), Mo4-Z5 (orange), and a physical mixture of MoO_3 with H-ZSM-5 (red).



a Mo–O–Mo antisymmetric stretch, suggestive of the presence of dimeric Mo species. Nevertheless, polymolybdate species also possess symmetric and antisymmetric Mo–O–Mo stretches at 630 cm^{-1} and 860 cm^{-1} , and Mo=O terminal stretches at 934 cm^{-1} , so it is difficult to draw definitive conclusions. Further Raman bands at lower frequencies are also observed and their position overlaps with signals from the zeolite framework. The 858 cm^{-1} band however, presents a slight asymmetry at the low-frequency side, which might be due to a contribution from residual MoO_3 . These observations indicate after the SSIE and calcination most of the Mo cations migrate into the zeolite channels and become highly dispersed species, although the presence of polymeric molybdate species cannot be completely ruled out.

Thermal stability and zeolite structural evolution

The thermal stability and behavior of Mo/H-ZSM-5 catalysts, including a 6 wt% sample designated by Mo6-Z5 and synthesized using the same method described above, were evaluated with DSC. All four DSC traces (Fig. 4) exhibit an endotherm below 200 which is due to dehydration. Above $1000\text{ }^\circ\text{C}$, an exothermic peak is seen and followed by a clear endothermic step. The exotherms are assigned to the amorphisation and collapse of the zeolite framework and are indicative of heat release upon the collapse. This dynamic has been reported for the collapse of many types of zeolites such as zeolite Y and zeolite A.^{25,26}

The thermal stability of the zeolite studied here is defined by the collapse temperature of the zeolite (t_c), which is the maximum of the exotherm. For the H-form zeolite, the measured t_c is $1218\text{ }^\circ\text{C}$, which is seen to drop for Mo2-Z5. Interestingly the measured t_c rises and plateaus for both Mo4-Z5 and Mo6-Z5,

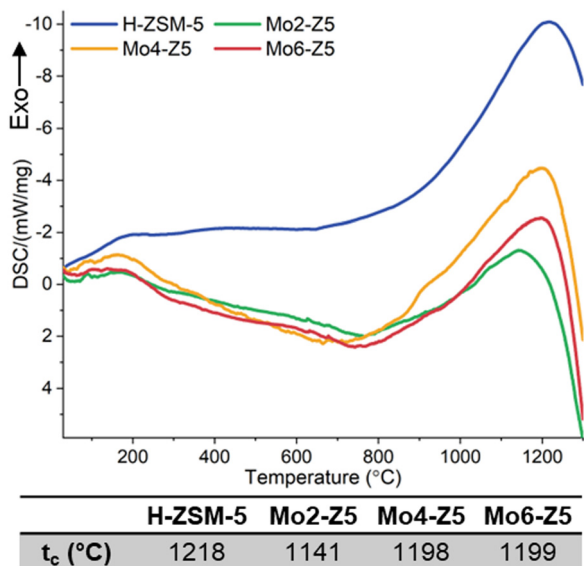


Fig. 4 DSC traces of H-ZSM-5, Mo2-Z5, Mo4-Z5, and Mo6-Z5. t_c s of the Mo exchanged H-ZSM-5 samples and H-form precursor are listed in Table 1. Note the weak shoulders at $900\text{ }^\circ\text{C}$ in the Mo4-Z5 and Mo6-Z5 samples are probably due to Mo volatilisation.

and its magnitude is on par with that of the H-form precursor. This unexpected rise happens against the backdrop of severe framework dealumination and the EFAl formation caused by Mo ions.

To verify the DSC measurement can indeed be correlated to zeolite the structural collapse and to further investigate the phase evolution during framework collapse, Mo2-Z5 and Mo4-Z5 were calcined to temperatures just below ($1150\text{ }^\circ\text{C}$), at ($1240\text{ }^\circ\text{C}$), and above ($1350\text{ }^\circ\text{C}$) the t_c . Samples calcined to these temperatures are given the label Mox-Z5-BC, Mox-Z5-@C and Mox-Z5-AC respectively with x denoting the Mo loading. At $1150\text{ }^\circ\text{C}$, below the collapse temperature (t_c), all but the most intense low-angle reflections of the MFI structure (Fig. 5a) diminished below the background for both Mo loading. Intensities of the reflections at 2θ angles of 7.92 and 8.87° were integrated for both samples and the result shows that the crystallinity determined for Mo2-Z5-BC and Mo4-Z5-BC are at 8.84 and 8.50% of the initial intensity, respectively. A broad diffuse scattering peak centered at 22° is also observed, and it is believed to originate from amorphous silica/alumina phases. Therefore, at temperatures just below t_c , Mo/H-ZSM-5 catalysts are already mostly PXRD amorphous. Interestingly, mullite ($\text{Si}_2\text{Al}_6\text{O}_{13}$), an aluminum-rich dense aluminosilicate phase, is detected alongside ZSM-5, suggesting that the recrystallisation of the thermodynamically more stable, denser, aluminous ceramic phase happens before the complete collapse of the zeolite phase.

Fig. 5b shows the PXRD patterns of Mo/H-ZSM-5 samples calcined to and above the t_c . At the onset of t_c , the remaining reflections from the MFI zeolite phase completely disappear, and an increase in mullite reflection intensity indicates a greater extent of crystallisation of the dense aluminous ceramic phase. BET specific surface area measurements show that Mo4-Z5-@C and Mo4-Z5-AC possess a total surface area of 8.50 and $2.30\text{ m}^2\text{ g}^{-1}$ respectively, confirming the microporous structure of the ZSM-5 is lost completely. Note that recrystallisation of the amorphous silica is not observed at t_c . In turn, at temperatures well above t_c , recrystallisation of the dense silicate phase in form of α -cristobalite (SiO_2) is observed, accompanied by a reduction in intensity of the broad amorphous peak. No molybdenum-containing crystalline phase was detected by the PXRD measurement, suggesting that molybdenum ions remain either highly dispersed in an amorphous phase or trapped in the crystalline phases formed.

The partial collapse of the zeolite framework at temperatures below t_c with co-current recrystallisation of the dense ceramic phase evidenced by PXRD point to a possible inhomogeneity in the framework collapse. One possible source for inhomogeneity in collapse behaviour is Al content as numerous studies with isostructural zeolites have concluded that the thermal stability increases with increasing the Si/Al ratio caused by the increasing covalent character associated with more Si–O bonds.^{27,28} EDX coupled with transmission electron microscopy (Fig. 6) further reveal anisotropy in Al distribution for Mo4-Z5 *i.e.* Al rich domains appear to be present in the sample. However, with its small(er) particle size (hundreds of



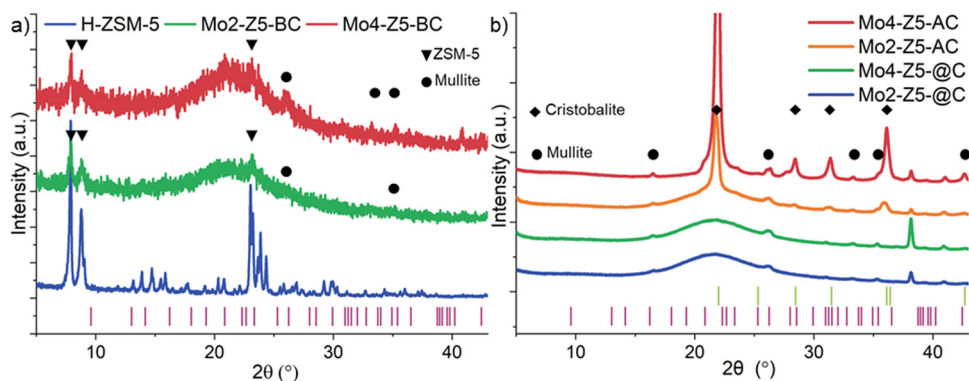


Fig. 5 PXRD patterns of Mo4-Z5-BC, and Mo2-Z5-BC compared to that of H-ZSM-5 (a), and Mo4-Z5-@C, Mo2-Z5-@C, Mo4-Z5-AC and Mo2AC (b). Indexing lines show position reflections from the mullite (purple) and cristobalite (light green).

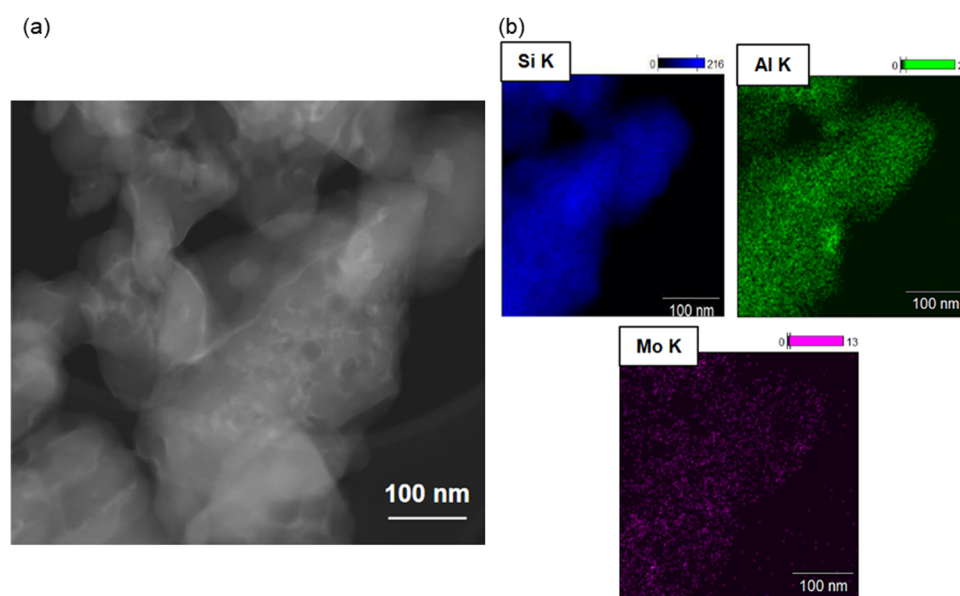


Fig. 6 Dark-field TEM image (a) and Si, Al, and Mo elemental EDX maps (b) for Mo4-Z5.

nanometers) it is inherently difficult to investigate distribution effects using this particular material. To clearly visualize if Al distribution influences thermal stability, a model SSZ-13 catalyst with a large crystallite size (*ca.* $30 \times 30 \times 30 \mu\text{m}$) – synthesized using a method described previously.¹⁷ Since differentiating between Si and Al is challenging using X-rays, Cu was added to the sample *via* conventional wet ion-exchange and XRF mapping on the basis that the Cu will provide framework charge compensation mimicking the Al distribution. Fig. S3 (ESI[†]) contains an XRF map of the Cu exchanged SSZ-13 revealing the Al, Si and Cu distributions from the crystal. Although the signals from the Al and Si appear to overlap, relative to the Si signal most of the Cu intensities are concentrated in a region around $20 \mu\text{m}$ in length and located at the core of the crystal, and the outer $5\text{--}7 \mu\text{m}$ of the crystal shows almost no Cu intensities at all. This type of distribution is indicative of the so-called “zoning effect”.

2 and 4 wt% Mo/H-SSZ-13 samples prepared using the SSIE method were calcined to $1300 \text{ }^\circ\text{C}$, the t_c of 2 wt% Mo/H-SSZ-13 (equivalent to a temperature just below the t_c of 4 wt% Mo/H-SSZ-13), and named Mo-S13-@C and Mo-S13-BC, respectively. The zeolite collapse and amorphisation of Mo exchanged model SSZ-13 catalyst are illustrated by SEM (Fig. 7). For Mo-S13-BC, although the calcination temperature is below t_c , the Al rich core of the crystal displays severe strain in the form of cavitation and delamination as if the core had imploded under stress. This type of morphology is typical of thermally collapsed zeolites transitioning from a rigid to an amorphous state. Contrary to the core of the crystal, the Si rich shell remains almost completely intact with no sign of fracturing or crumbling while PXRD measurement (Fig. S4, ESI[†]) shows some of the CHA structure is maintained. When the calcination temperature reaches the t_c , the $30 \mu\text{m}$ sized crystals start to adopt a globular morphology that is reminiscent of a melting solid



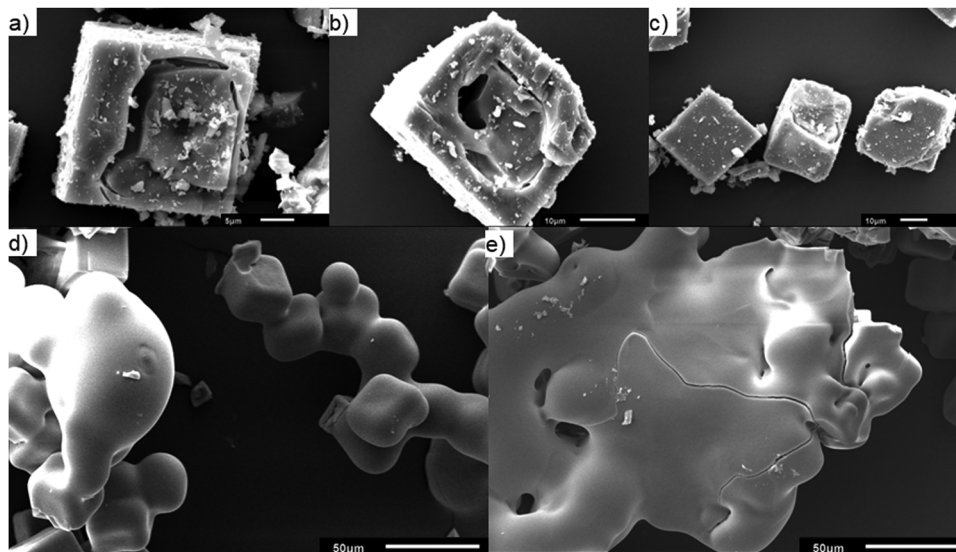


Fig. 7 SEM micrographs of Mo-S13-BC (a)–(c) and Mo-S13-@C (d) and (e).

(Fig. 7d and e). Evidence of flow is clearly demonstrated in SEM micrographs, where the amorphised crystals have merged into a large, single globular unit.

As summarised by the scheme in Fig. 8, it appears that within a single crystallite of zeolite SSZ-13, the Al rich and Si rich regions are not only amorphised at different temperatures but also with different modalities. The Al rich core is amorphised *via* framework collapse at a relatively low temperature below t_c and forms a rigid amorphous solid first before recrystallisation. The Si rich shell is amorphised in a fashion typical to classical solid melting. This difference in collapse temperature is perhaps not surprising since numerous studies on the effects of varying Si/Al ratio of a series of isostructural zeolites have concluded that the thermal stability increases with increasing the Si/Al ratio. However, previous effort mostly focused on the macroscopic aspect while the microscopic aspect such as the

effects of framework Al distribution within a single particle of zeolite has been overlooked.

Mo speciation after collapse, a XAFS study

As the state of Mo ions in the collapsed zeolite catalysts was not accounted for by the PXRD, XAFS measurements were performed to obtain insight into the speciation of molybdenum during the collapse. Four reference materials, two calcined 4 wt% Mo/H-ZSM-5, one measured in ambient conditions at RT (MoZ-RT) and the other measured *in situ* in Ar atmosphere after the 700 °C calcination and cooled to RT (MoZ-700), as well as aluminium molybdate $\text{Al}_2(\text{MoO}_4)_3$, and molybdenum trioxide MoO_3 are also measured. The collapsed Mo/H-ZSM-5 (Fig. 9a) shows a very similar spectral profile to those of $\text{Al}_2(\text{MoO}_4)_3$ and MoZ-700 reference where the pre-edge region

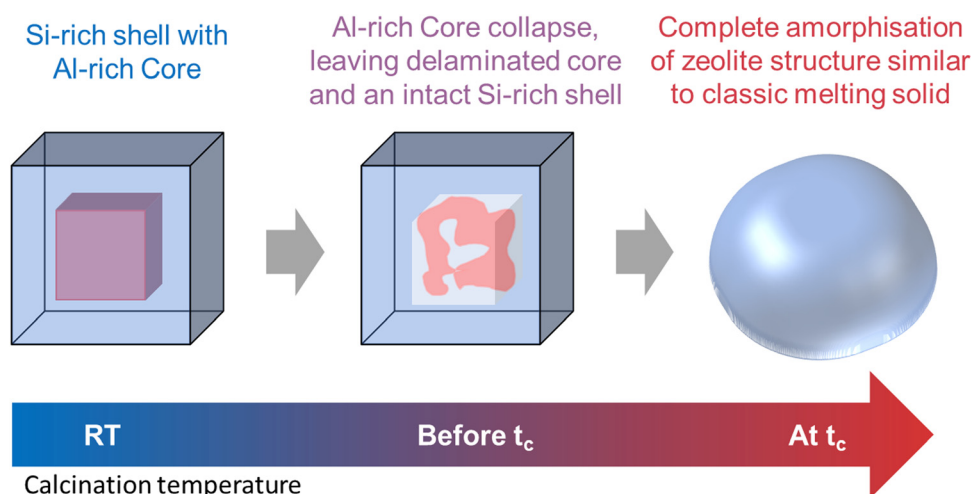


Fig. 8 Schematic representation depicting the temperature dependent multi-modal collapse mechanism of a single crystallite of zeolite SSZ-13.



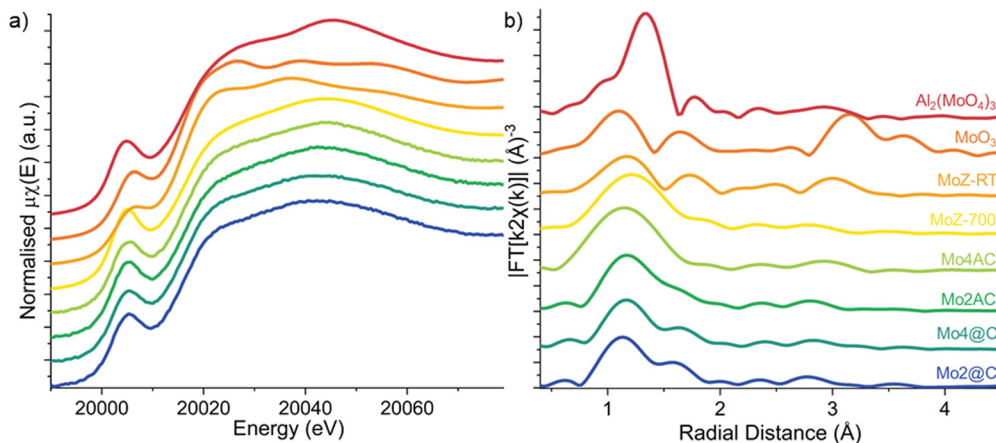


Fig. 9 Mo K-edge XANES spectra (a) and k^2 weighted FT-EXAFS magnitude (b) of Mo2-Z5-@C, Mo4-Z5-@C, Mo2-Z5-AC, and Mo4-Z5-AC sample, and MoZ-700, MoZ-RT, MoO₃, and Al₂(MoO₄)₃ reference (from bottom to top).

is dominated by a pre-edge peak at 20 005 eV and it is assigned to the dipole forbidden (quadrupole allowed) $1s \rightarrow 4d$ transition, while the white line at 20 014 eV is attributed to the $1s \rightarrow 5p$ dipole transition. Note the increased intensity for the pre-edge peak for Mo in Al₂(MoO₄)₃ when compared to MoO₃ since the former contains monomeric tetrahedral [MoO₄]²⁻ species which leads to a relaxing of the dipole selection rule due to the removal of orbital parity in a tetrahedral environment. In the case of Mo/H-ZSM-5, studies have shown evidence for the presence of isolated monomeric Mo-oxo species at low Si/Al ratios,^{2,6} although the presence of dimeric species cannot be ruled out. More recently, utilizing a combination of *in situ* and *operando* XAFS and high-resolution PXRD, Agote-Arán *et al.* were able to reveal that during calcination the majority of the Mo ions adopt an isolated tetrahedral Mo-oxo arrangement anchored to the straight MFI channels.⁷ Furthermore, the lack of spectra feature in the post-edge region is in line with the assignment of highly dispersed Mo species that do not possess long-range order. As such, the XANES spectra indicate that similar to the *in situ* calcined Mo/H-ZSM-5 catalyst, the Mo ions in the collapsed ZSM-5s are also highly dispersed and mostly tetrahedrally coordinated.

Analysis of the energy position of the absorption edge allows the identification of the formal oxidation state (Fig. S5, ESI[†]) and it shows the Mo ions in all four samples are preserved as +6. Linear combination fitting (LCF) of the XANES spectra (Fig. S5, ESI[†]) listed in Table S2 (ESI[†]) demonstrate MoZ-700 is the dominant species in all samples, again confirming the majority of the Mo⁶⁺ ions adopting a tetrahedral coordination geometry. The result also suggests that the rest of the Mo

ions adopt a distorted octahedral geometry similar to that of hydrated Mo species in calcined Mo/H-ZSM-5 zeolites. It is proposed that tetrahedral coordinated Mo-oxo species formed inside the zeolite during the calcination to 700 °C remain stable and do not undergo thermally induced reduction during further calcination up to 1350 °C and complete collapse of the zeolite framework. As the ZSM-5 framework collapses, the denser new phases trap the Mo-oxo species and seal off their access to moisture, preventing them from reverting back to the octahedral coordination.^{4,29} The Mo-oxo species on the surface, however, do have access to air/moisture and they undergo rehydration and contribute to the XANES spectra similarly to that in Mo-RT.

The accompanying EXAFS allows independent validation of the XANES results, specifically on local coordination around Mo ions. The k^2 -weighted Fourier transform (FT)-moduli (Fig. 9b, k -space EXAFS spectra shown in Fig. S7, ESI[†]) of the collapsed catalysts are dominated by a first-shell contribution and is assigned to near neighbor O ions. First shell fitting of the EXAFS spectra of the thermally collapsed Mo/H-ZSM-5 catalysts (Fig. S8, ESI[†]) shows the fitted average Mo-O coordination number (CN) of all four collapsed Mo zeolite samples is slightly above 4 (Table 2), consistent with a tetrahedrally coordinated Mo-oxo species. The Mo-O CNs for the two @C samples are higher than those fitted for the AC samples, suggesting a higher fraction of Mo ions present with a coordination number greater than 4. This result is in agreement with the XANES analysis, although it is worth noting the difference in CNs is within the error for determining these values (*i.e.* 10%). The lack of intensity in FT-moduli beyond the first coordination shell suggests that the structures around the Mo ions lack long-range order. These results from the FT-EXAFS are consistent with isolated [MoO₄]²⁻ cations being bonded to double Al sites in the ZSM-5 framework.

The analysis of second-shell contribution in FT-EXAFS proved to be complicated. The relatively weak scattering intensity and the possible destructive interference among scatterers,³⁰ combined

Table 1 Collapse temperature (t_c (°C)) for samples with increasing [Mo] derived from the DSC traces displayed in Fig. 4

Sample	H-ZSM-5	Mo2-Z5	Mo4-Z5	Mo6-Z5
t_c (°C)	1218	1141	1198	1199



Table 2 Structure parameters obtained from the structural model fitted to the EXAFS spectra

Sample	Scattering path	Coordination number (N)	Bond length (\AA)	ΔE (eV)	σ^2 (\AA^2)	R -factor
MoZ-700	M=O	2.0 (6)	1.69 (4)	-2.7 (2.7)	0.003 (5)	0.015
	M-O	2.0 (6)	1.82 (4)			
Mo2-Z5-@C	M-O	4.6 (1.5)	1.76 (3)	-3.6 (4.8)	0.012 (5)	0.02
Mo4-Z5-@C	M-O	4.7 (1.4)	1.75 (3)	-2.2 (4.3)	0.012 (4)	0.028
Mo2-Z5-AC	M-O	4.2 (1.2)	1.77 (2)	0.9 (4.0)	0.008 (4)	0.005
Mo2-Z5-AC	M-O	4.2 (1.0)	1.73 (2)	-8.6 (3.4)	0.004 (3)	0.039

with multiple possible structural models all add complexity to FT-EXAFS-based analysis. To resolve the structure of the Mo species beyond the first coordination shell, we employed wavelet transform (WT) technique for the analysis of EXAFS. WT-EXAFS has the ability to decompose a given EXAFS signal in k -space that allows the resolution of overlapping contributions from different elements. Fig. 10 illustrates the contour plots of the Morlet WT-EXAFS moduli. All the contours contain an intense first-shell peak located at $R = 1.21 \text{ \AA}$ (phase uncorrected) and around $3.7\text{--}3.9 \text{ \AA}^{-1}$ in $\chi(k)$, which is assigned to a first-shell Mo–O contribution. The second-shell contribution (located around $R = 3.0 \text{ \AA}$) in the contour of MoZ-RT consists of two peaks. For reference, the second shell Mo–Al/Si peak in Mo/H-ZSM-5 is believed to be located around 2.9 \AA ,³⁰ while the Mo–Mo peaks are in the range of 2.9 to 3.2 \AA .^{20,31} The position of the low $\chi(k)$ peak (at 3.7 \AA^{-1}) is almost identical to that of oxygen, although the presence of multiple scattering contributions cannot be ruled out. An intense peak centered around 8.3 \AA^{-1} is also observed. The significantly higher $\chi(k)$ position designated this peak to contribution from a heavy element, and it is assigned to a Mo–Mo single scattering contribution. The presence Mo–Mo and possible multiple scattering contributions evidence that at RT Mo ions in ZSM-5 are not isolated and are most likely a di/poly-molybdate species.³⁰ This finding echoes the result from Kerr-gated Raman spectra. The contour plot of MoZ-700 shows a single second-shell peak located at $\chi(k) = 4.7 \text{ \AA}^{-1}$. The $\chi(k)$ position of this second-shell peak indicates it originates from a backscatterer with Z greater than an O anion but not as heavy as the Mo anion. For Mo/H-ZSM-5, it can only be Al (or Si). As such, the WT-EXAFS analysis indicates that Mo anions calcined to $700 \text{ }^\circ\text{C}$ exists as an isolated species anchored to the ZSM-5 zeolite framework. These observations are fully consistent with the result reported by Agote-Arán *et al.*⁷ The contour plots of Mo4-Z5-@C and Mo4-Z5-AC are almost identical to that of MoZ-700 with a single second-shell peak located at $\chi(k) = 4.9 \text{ \AA}^{-1}$, while the peak at high $\chi(k)$ indicative of an Mo–Mo contribution is absent. As such, the WT-EXAFS indicates that the Mo species in the collapsed Mo/H-ZSM-5 catalysts resemble the local structure of the isolated tetrahedral Mo–oxo species attached to the channels of ZSM-5. This result indicates during the collapse of the zeolite framework not only the oxidation state and coordination geometry of the Mo cation are preserved, but also possibly the fragments/building units of the zeolite structure immediately surrounding Mo (*i.e.*, the 10-membered ring where the Mo cation is anchored to) are retained.⁷

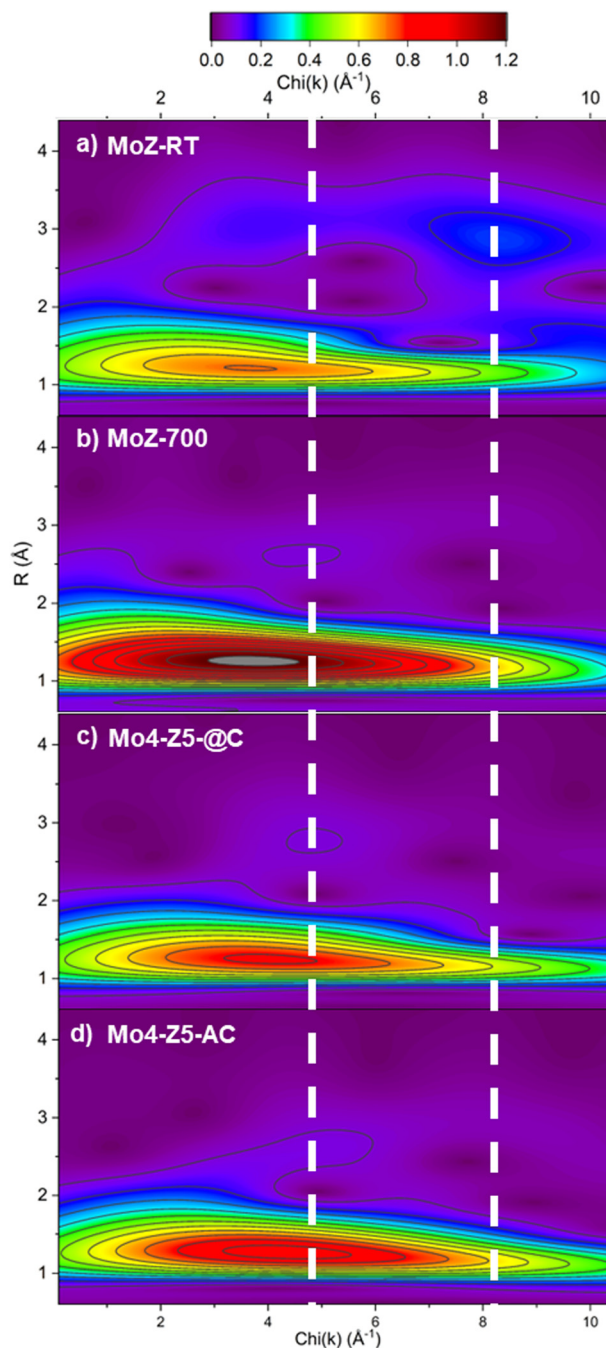


Fig. 10 k^2 weighted Morlet WT-EXAFS contour plot of (a) MoZ-RT, (b) MoZ-700, (c) Mo4-Z5-@C, and (d) Mo4-Z5-AC. The white dotted lines mark the k -space position of the Mo–Al/Si and Mo–Mo single scattering path at 4.8 \AA^{-1} and 8.3 \AA^{-1} . All four contours are plotted to the same color scale.



Methane dehydroaromatisation activity

Evidence from both XANES and EXAFS analysis converges on the notion that the Mo⁶⁺ ions after the high temperature calcination are preserved in a state similar to the Mo⁶⁺ ions in the freshly calcined Mo/H-ZSM-5 catalyst before methane activation, while chemical analysis confirmed more than 25% of the original Mo loading is preserved (Table S3, ESI†). This opens up the question of whether these Mo ions are still catalytically active toward the MDA reaction. The resulting benzene response from the MDA reaction are plotted in Fig. S9 (ESI†). Both catalysts were able to activate methane. The observed C₆H₆ evolution on the collapsed Mo/H-ZSM-5 catalysts is in agreement with studies by Kosinov *et al.*,³² and Agote-Arán *et al.*⁷ that suggest that Brønsted acidity is not required for the aromatisation to occur, since the framework that hosts the Brønsted acid sites has already collapsed. As shown, the benzene response of the two catalysts differs significantly. Relative to Mo/H-ZSM-5, the at collapsed catalyst affords 80% of the original response while the collapsed only managed 15%.

Summary and conclusions

This study brings new insights into the thermal stability, the collapse mechanism and nature of the Mo species of the benchmark MDA catalyst Mo/H-ZSM-5. *Ex situ* Kerr-gated Raman spectra at room temperature suggesting the presence of both monomeric and polymeric molybdate species, the latter species increasing with [Mo]. A consequence of this is the formation of more 'hydrated' and non-crystalline Al₂(MoO₄)₃ although interestingly this did not impinge on the thermal stability of the ZSM-5 as the temperature at which crystallinity is lost did not decrease with [Mo] loading remaining on par with the parent H-form.

The structural evolution during the collapse of Mo/H-ZSM-5 was then investigated from the perspective of both the zeolite framework and Mo ions. The MFI structure was almost in a state of complete collapse at a temperature just below *t_c*; we note however that previous work has shown that although the XRD pattern indicates that long range crystalline ordering has been lost, many of the structural units (*i.e.* Secondary Building Units (SBUs) or common fragments (cages)) are often retained immediately after collapse in what has been termed a low density amorphous (LDA) phase.^{25,33} Further heating leads to complete collapse of the aluminosilicate framework to form a high density amorphous oxide and an aluminium rich dense ceramic phase mullite, while the dense silicate phase α -cristobalite recrystallizes at temperatures above the *t_c*. Further investigation on the mechanism of collapse with a model Mo/H-SSZ-13 crystals demonstrates that (as a result of the Al zoning effect) within a single crystal of zeolite the Al rich and Si rich regions not only amorphized (collapse) at different temperatures, but also with different modalities (illustrated in Fig. 5). The Al rich region amorphised *via* framework collapse at a temperature below *t_c* and forms a rigid amorphous solid first,

while the Si rich shell amorphised in a fashion typical to classical solid melting. These findings can have a profound implication on studies of zeolite in general since zeolites are often characterised by its macroscopic properties, while microscopic properties within a particle of zeolite have often been overlooked. This result offers a possible explanation for the seemingly contradictory *t_c* value observed for Mo/H-ZSM-5s. The observed formation of EFAl and aluminium molybdate species and the resulting framework damage are mostly concentrated in the Al-rich region, while the Si rich region is mostly unaffected. Hence the Si rich region would have thermal stability on par to that of the H-ZSM-5.

On the other hand, the exchanged Mo⁶⁺ ions do not undergo autothermal reduction during the high temperature calcination up to 1350 °C. Most of the Mo⁶⁺ ions in the collapsed zeolite adopt tetrahedral coordination similar to freshly calcined Mo/H-ZSM-5. These results suggest that the Mo ions were trapped by the collapsed zeolite and do not directly participate in the collapse process and indeed these observations would also be consistent with the retention of ZSM-5 structural units in an LDA which retains many of the key structural features necessary for catalytic activity. WT-EXAFS analysis indicates that the Mo⁶⁺ ions can be considered isolated after treatment at reaction temperatures (700 °C) and after the zeolite framework collapse, and more importantly, they form an aluminate species that bear similarities to the freshly calcined tetrahedral Mo⁶⁺ species that anchor to the channels of ZSM-5. We note that this suggests an evolution of the Mo⁶⁺ species after high temperature treatment from polynuclear to mononuclear and also emphasizes the importance of performing measurements under *in situ* or ideally under *operando* conditions.

These results suggest that in Mo-based ZSM-5 catalysts the thermal stability of the zeolite support alone during the reaction and regeneration is important for ensuring good catalytic performance. Interestingly we observe that in already collapsed Mo-zeolites, the newly formed dense phase should not suffer from the stability issue experienced by the zeolite and can act as a kinetically more stable support for the Mo ions. Furthermore, since MDA is an endothermic reaction with aromatisation limited by thermodynamics,⁸ Mo ions supported on a more stable material may be able to facilitate higher operation temperature, which could shift the equilibrium and achieve higher conversion.

From these observations one can conclude that the macroscopic thermal stability that has been determined from techniques such as PXRD or DSC can be somewhat irrelevant to zeolite catalysis *per se*, since an Al distribution gradient can exist in zeolite particles in the form of zoning, and the thermal stability of the catalytic relevant framework Al mostly depends on the microscopic Al concentration within the zeolite crystal. In the most extreme cases such as the one demonstrated here in the model SSZ-13, the catalyst may already be deactivated by the irreversible structural change introduced by the collapse of framework Al, as many of the active sites of zeolite catalysts require either (or both) the framework Al or the charge balancing cation anchored to the framework Al to adopt very specific



coordination, while the crystallinity of the catalyst is still mostly maintained due to the Si rich part of the zeolite able to survive through the high temperature condition.

Data availability

Processed data (spectra and patterns) are available from the following repository: <https://rdr.ucl.ac.uk/>.

Conflicts of interest

Andrew M. Beale reports a relationship with Finden Ltd. that includes co-ownership of the company in the form of equity and grant funding. All other authors have no known competing financial interest or personal relationship that could have appeared to influence the work reported in this paper.

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