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Article

Ga/HZSM-5 Catalysed Acetic Acid Ketonisation for Upgrading of Biomass Pyrolysis Vapours

Hessam Jahangiri ^{1,2,3} , Amin Osatiashtiani ³ , Miloud Ouadi ¹, Andreas Hornung ^{1,4,5}, Adam F. Lee ^{6,*}  and Karen Wilson ^{6,*}

- ¹ School of Chemical Engineering, University of Birmingham, Birmingham B15 2TT, UK; h.jahangiri@bham.ac.uk (H.J.); m.ouadi@bham.ac.uk (M.O.); andreas.hornung@umsicht.fraunhofer.de (A.H.)
² School of Water, Energy and Environment, Cranfield University, Cranfield MK43 0AL, UK
³ European Bioenergy Research Institute, Aston University, Birmingham B4 7ET, UK; a.osatiashtiani@aston.ac.uk
⁴ Fraunhofer UMSICHT, Fraunhofer Institute for Environmental, Safety and Energy Technology, An der Maxhütte 1, 92237 Sulzbach-Rosenberg, Germany
⁵ Friedrich-Alexander University Erlangen-Nuremberg, Schlossplatz 4, 91054 Erlangen, Germany
⁶ Applied Chemistry & Environmental Science, RMIT University, Melbourne, VIC 3000, Australia
* Correspondence: adam.lee2@rmit.edu.au (A.F.L.); karen.wilson2@rmit.edu.au (K.W.); Tel.: +61-399-252-623 (A.F.L.); +61-(03)-9925-2122 (K.W.)

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Abstract: Pyrolysis bio-oils contain significant amounts of carboxylic acids which limit their utility as biofuels. Ketonisation of carboxylic acids within biomass pyrolysis vapours is a potential route to upgrade the energy content and stability of the resulting bio-oil condensate, but requires active, selective and coke-resistant solid acid catalysts. Here we explore the vapour phase ketonisation of acetic acid over Ga-doped HZSM-5. Weak Lewis acid sites were identified as the active species responsible for acetic acid ketonisation to acetone at 350 °C and 400 °C. Turnover frequencies were proportional to Ga loading, reaching $\sim 6 \text{ min}^{-1}$ at 400 °C for 10Ga/HZSM-5. Selectivity to the desired acetone product correlated with the weak:strong acid site ratio, being favoured over weak Lewis acid sites and reaching 30% for 10Ga/HZSM-5. Strong Brønsted acidity promoted competing unselective reactions and carbon laydown. 10Ga/HZSM-5 exhibited good stability for over 5 h on-stream acetic acid ketonisation.

Keywords: pyrolysis; ketonisation; bio-oil; turnover frequencies (TOFs)

1. Introduction

Hydrocarbons sourced from non-edible or waste lignocellulosic or algal biomass are an attractive source of sustainable liquid transportation fuels to mitigate current dependence on fossil fuels and associated anthropogenic climate change [1,2]. However, biomass-derived fuels are incompatible with existing distribution infrastructure and vehicle engines without (catalytic) upgrading to improve their physicochemical properties [2,3]. A range of thermochemical technologies exist for bio-oil production, including hydrothermal liquefaction [4,5] and pyrolysis [6,7], or gasification [8,9] and subsequent Fischer–Tropsch synthesis [10,11]. Pyrolysis routes have gained particular attention over the past 30 years, offering a high liquid (bio-oil) yield which can be used directly as a drop-in fuel, blended with conventional diesel, or as an efficient energy vector [12,13]. Pyrolysis bio-oils are mixtures of oxygenated compounds which typically comprise phenolics, furanics, carboxylic acids and other small oxygenates whose composition varies with biomass source and processing [6,14,15]. The high oxygen content of crude bio-oils results in a heating value half that of petroleum-derived fuels, while the presence of carboxylic acids renders the oils corrosive (pH 2–3) [15] and chemically unstable due to

presence of small reactive oxygenates (e.g., unsaturated aldehydes) which may undergo acid-catalysed polymerisation [2]. Crude bio-oils must therefore be upgraded to remove corrosive components and improve stability prior to subsequent hydrodeoxygenation to improve their calorific value.

A range of catalytic routes exist for upgrading pyrolysis bio-oils, including esterification [16], hydrodeoxygenation (HDO) [17], aldol condensation [18] and ketonisation [19]. Each route has advantages and disadvantages. For example, esterification operates at low temperature in the liquid phase but requires an external source of short-chain alcohols, and produces water by-product which must be separated [20]. HDO is effective for the production of cyclic and aliphatic alkanes as liquid fuels, but requires a renewable H₂ input and precious metal catalysts that are stable in (often acidic) bio-oils and coke-resistant [21]. Aldol condensation stabilises bio-oils by converting some reactive oxygenates over solid base catalysts, but does not address the intrinsic acidity of bio-oils that can deactivate HDO catalysts [22]. Ketonisation affords an intermediate deoxygenation step that can be close-coupled to a pyrolysis reactor to upgrade vapours before then condense into a bio-oil, thereby improving bio-oil acidity, and achieving partial deoxygenation [19], although it is also accompanied by a small loss of carbon as CO₂. Ketonisation [22–24], takes place through the condensation of two carboxylic (e.g., acetic) acid molecules to form a ketone (e.g., acetone), CO₂ and H₂O (Figure 1). An important advantage of ketonisation over esterification for acid neutralisation is that the former can be performed in the vapour phase without additional reactants, thus enabling close-coupling to a pyrolysis reactor to upgrade bio-oil vapours prior to their condensation as a bio-oil [24,25]. Ketonisation also facilitates bio-oil deoxygenation and concomitant hydrocarbon chain growth [26], and hence improves the calorific value of the resulting condensate (in addition to its pH and stability).

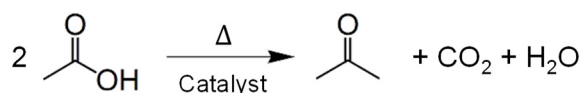


Figure 1. Acetic acid ketonisation.

Ketonisation is widely studied in organic synthesis [25,27], being catalysed by diverse heterogeneous catalysts including alkaline earth metal oxides such as BaO, MgO [28,29], transition metal oxides including MnO₂ [24,30,31], TiO₂ [32,33], Fe₃O₄ [22,24,34], CeO₂ [35,36] and ZrO₂ [37,38] and actinide oxides such as ThO₂ [39]. The mechanism of ketonisation and corresponding sensitivity to catalyst properties remains the subject of ongoing debate [25,40]. Ketonisation over basic and reducible oxide catalysts proceeds via two distinct pathways depending on the lattice energy of the metal oxide, with lower energy lattice (stronger bases) forming stable carboxylates that thermally decompose at elevated temperature (>420 °C) [41] to yield ketones, whereas higher energy lattices favour a lower temperature surface catalysed route [25]. However, there are fewer reports of carboxylic acid ketonisation over zeolites, being limited to HZSM-5, HZSM-11, HZSM-34, HZSM-35, mordenite, erionite and zeolite Beta. Of these, HZSM-5(100) is the most favourable for acetic acid ketonisation to acetone [42], being very selective to xylenols and acetone at moderate reaction temperatures (320 °C) and forming acetone as the major product >350 °C [42]. Zeolite modification by transition metals and lanthanides such as Ce, Co, Ni and Ga increases aromatic product yields during catalytic pyrolysis [43–45], however to our knowledge Ga-promoted zeolites have never been investigated for carboxylic acid ketonisation. Gallium can be introduced into zeolites by incipient wetness impregnation and ion exchange [46], although the choice of preparation method had little impact on aromatic products from Ga/HZSM-5-catalysed fast pyrolysis [46].

HZSM-5 is an attractive catalyst for acetic acid ketonisation, owing to its corrosion resistance, high surface area, commercial availability and ability to stabilise undercoordinated cations and hence tune surface composition and resulting catalytic performance [25]. The presence of strongly acidic protons in zeolites reportedly promotes the formation of surface acyl species, rather than carboxylates, following acid adsorption. Subsequent coupling of a carboxylic acid and surface acyl yields an acid anhydride, which in turn dissociates to liberate CO₂ and a ketone as illustrated in Figure 2 [25,47],

although Chang et al propose that ketonisation proceeds by nucleophilic attack of an acylium ion by adsorbed carboxylate [48]. The latter is similar to a ketene intermediate pathway, in with the acylium ion is directly formed by acid protonation and water loss [48]. The relationship between Lewis/Bronsted acid character and ketonisation activity/selectivity over zeolites remains poorly established.

1. Adsorption of carboxylic acid with elimination of water,
2. Surface acyl formation,
3. Surface adsorption of second carboxylic acid,
4. Coupling of adsorbed carboxylic acid with a surface acyl group to form an acid anhydride,
5. Adsorbed anhydride decomposition to form CO_2 and acetone.

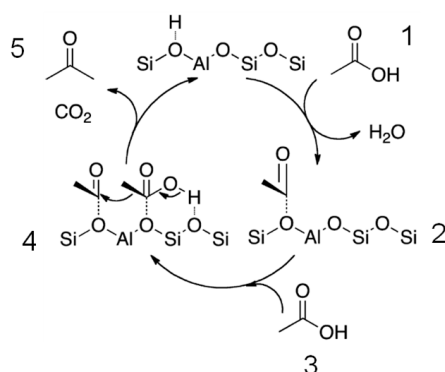


Figure 2. Proposed mechanism for acetic acid ketonisation to acetone over zeolites. Reproduced with permission from reference [47], Copyright © 2016, Elsevier.

Herein, we report the impact of Ga doping on the surface acidity of HZSM-5 and associated reactivity for the continuous vapour phase ketonisation of acetic acid to acetone. Activity and selectivity to acetone were proportional to Ga loading, reflecting the formation of weak Lewis acid sites and suppressed coking.

2. Results and Discussion

2.1. Catalyst Characterisation

Crystalline phases were characterised by powder X-ray diffraction (XRD). Figure S1 shows diffraction patterns for HZSM-5 as a function of Ga loadings to HZSM-5, and pure bulk Ga_2O_3 . No reflections associated with gallium oxide phases were observed for any loadings, indicating either the presence of highly dispersed Ga_2O_3 nanoparticles throughout the HZSM-5 pore network, or the exchange of Ga^{3+} with Al^{3+} ions in the framework (or protons at the surface of the zeolite). It is well documented that protons play an important role in regulating the interaction of metal oxides with zeolite surfaces [49,50]. Fang and co-workers report that impregnation favours the formation of Ga_2O_3 and small amounts of GaO^+ at surface of ZSM-5, thereby introducing weak Lewis acid sites [51]. In contrast ion-exchanged Ga/HZSM-5 prepared by refluxing the zeolite in aqueous $\text{Ga}(\text{NO}_3)_3$ at 70–100 °C [46,52] appears to favour framework dealumination through Ga ion-exchange. Although the latter syntheses resemble our impregnating conditions we cannot conclude whether Ga resides as surface GaO^+ clusters or within the zeolite framework. The reference gallium oxide was phase-pure monoclinic (m- Ga_2O_3) with reflections at $2\theta = 19.0^\circ, 30.4^\circ, 30.5^\circ, 31.8^\circ, 33.5^\circ, 35.2^\circ, 37.4^\circ, 38.5^\circ, 42.8^\circ, 45.9^\circ, 48.7^\circ$ and 57.5° [53,54]. Crystallite sizes of the parent HZSM-5 (Table 1) were independent of Ga loading, and significantly smaller than the zeolite. Nitrogen porosimetry revealed type IV isotherms for xGa/HZSM-5 (Figure S2), with the observed mesoporosity attributed to interparticle voids [55]. Corresponding Brunauer-Emmett-Teller (BET) surface areas, total pore volumes and micropore volumes continuously decreased with increased Ga loading (Table 1), attributed to partial

pore blockage, possibly as a result of extra-framework gallium deposition within the micropores [56]. Bulk Ga_2O_3 exhibited a very low surface area $<10 \text{ m}^2 \cdot \text{g}^{-1}$.

Table 1. Elemental analysis and physicochemical properties of catalysts.

| Catalysts | Ga Loading ^a /wt% | S_{BET} / $\text{m}^2 \cdot \text{g}^{-1}$ | V_{Total} ^b / $\text{mL} \cdot \text{g}^{-1}$ | V_{micro} ^c / $\text{mL} \cdot \text{g}^{-1}$ | Crystallite Size ^d /nm | Acid Loading ^e / $\text{mmol} \cdot \text{g}^{-1}$ | Weak:Strong Acid Site Ratio ^f |
|-------------------------|------------------------------|-----------------------------------------------------|-------------------------------------------------------------------|-------------------------------------------------------------------|-----------------------------------|---------------------------------------------------------------|------------------------------------------|
| HZSM-5 | 0 | 427 | 0.294 | 0.141 | 65.7 | 1.13 | 0.29 |
| 0.5 Ga/HZSM-5 | 0.3 | 420 | 0.290 | 0.140 | 55.2 | 1.09 | 0.38 |
| 3 Ga/HZSM-5 | 3.0 | 338 | 0.249 | 0.106 | 50.3 | 1.03 | 0.53 |
| 10 Ga/HZSM-5 | 9.0 | 313 | 0.210 | 0.099 | 62.6 | 0.80 | 0.83 |
| Ga_2O_3 | 75 | 7.6 | 0.08 | – | 26.4 | 0.14 | 0.40 |

^a ICP-OES, ^b total pore volume at P/Po = 0.98, ^c t-plot method, ^d XRD, ^e propylamine desorption, ^f XPS.

Elemental analysis revealed the surface Ga content was consistently lower than the bulk determined by XPS and ICP-AES analysis respectively (Table S1), consistent with the selective incorporation of gallium inside the HZSM-5 pore network. The formation of large Ga_2O_3 particles on the external surface of zeolite crystallites can be discounted due to the absence of associated XRD patterns. Note that the lower Ga surface versus bulk content for the m- Ga_2O_3 reference reflects oxygen termination of gallium surfaces [57]. O 1s XP spectra of HZSM-5 revealed a single broad peak with a 533 eV binding energy associated with Si–O–Si and Si–O–Al environments [58,59] (Figure 3a), which was unaffected by low levels of Ga doping, but shifted to lower binding energy for 10Ga/HZSM-5, approaching that of Ga_2O_3 at 530.7 eV [60,61]. A similar trend was observed for the Ga 2p_{3/2} XP spectra (Figure 3b), which exhibited a single broad peak at 1119.0 eV for low Ga loadings, whose binding energy decreased towards that of m- Ga_2O_3 at 1117.9 eV for 10Ga/HZSM-5 [62]. These data demonstrate that the local environment of gallium in HZSM-5 is chemically distinct from that in bulk Ga_2O_3 , consistent with either highly dispersed Ga_2O_3 nanoparticles, or ion-exchange of Ga^{3+} into the zeolite framework [62,63]. Corresponding Al and Si 2p XP spectra of xGa/HZSM-5 (Figure S3a,b) each evidenced a single chemical environment with respective binding energies of approximately 75.1 eV and 103.8 eV, consistent with the literature for HZSM-5 [58,59]. Al and Si 2p peaks shifted to lower binding energy for 10Ga/HZSM-5 indicative of significant ion-exchange and concomitant formation of extra-framework alumina.

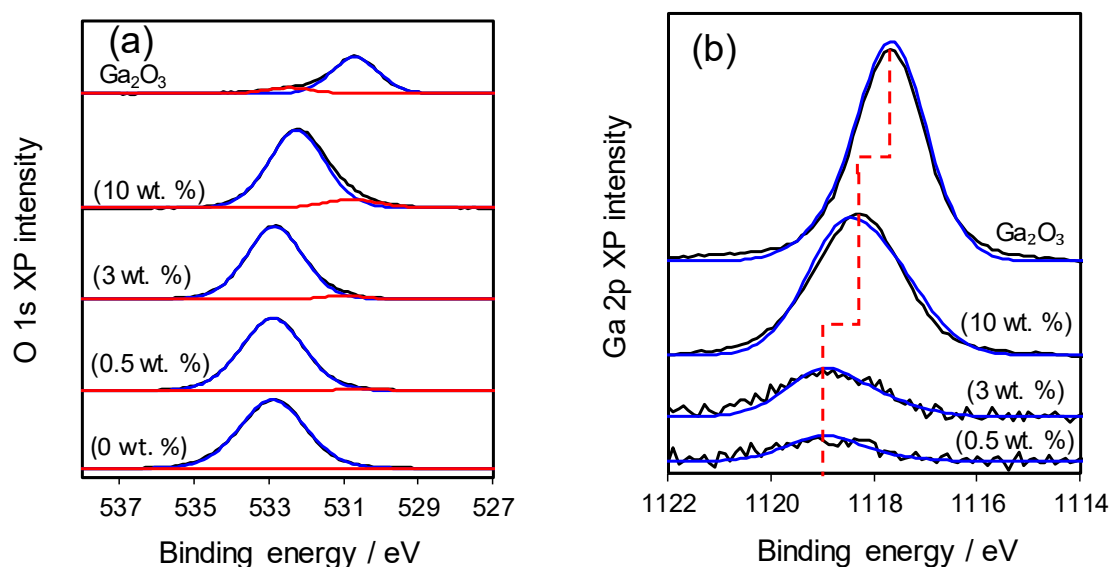


Figure 3. (a) O 1s and (b) Ga 2p XP spectra of xGa/HZSM-5 and Ga_2O_3 .

The acid properties of xGa/HZSM-5 and m- Ga_2O_3 were first investigated by diffuse reflectance Fourier transform infrared spectroscopy (DRIFTS) following pyridine adsorption (Figure S4a).

Strong bands at 1444 cm^{-1} and 1545 cm^{-1} were assigned to pyridine bound to Lewis and Brønsted acid sites respectively, the intense band at 1490 cm^{-1} to pyridine bound to both acid sites and the weak 1600 cm^{-1} band to pyridine bound to Lewis acid sites [56]. The relative Lewis/Brønsted acid character was quantified from the ratio of 1444 cm^{-1} and 1545 cm^{-1} band intensities (Figure S4b). The Lewis:Brønsted ratio exhibited a small increase for 10Ga/HZSM-5, in accordance with literature reports [56,64,65]. Corresponding DRIFTS for pyridine on $m\text{-Ga}_2\text{O}_3$ revealed two weak bands at 1452 cm^{-1} and 1614 cm^{-1} indicative of pure Lewis acid character as previously reported [66,67]. Acid strength was subsequently probed by temperature-programmed reaction spectroscopy (TPRS) of propylamine. Reactively-formed propene (arising from propylamine decomposition over acid sites) was evolved in two desorptions at $\sim 480\text{ }^\circ\text{C}$ and $\sim 540\text{--}555\text{ }^\circ\text{C}$ associated with strong and weak acid sites respectively (Figure 4); the former possibly arising from high-index facets or defects [68,69]. The desorption temperature of both peaks was independent of Ga loading, however the ratio of weak:strong acid sites decreased monotonically reaching ~ 0.83 for 10Ga/HZSM-5. The decreased acid strength was consistent with ion-exchange of less electronegative Ga^{3+} for Al^{3+} into the zeolite surface, which is expected to decrease hydroxyl polarisation and hence Brønsted acid strength [70]. Acid site loadings and weak:strong acid site ratio respectively decreased and increased with Ga loading (Table 1 and Figure S5), however the total acid site density was approximately constant at $\sim 2.6\text{ }\mu\text{mol}\cdot\text{m}^{-2}$. The acid site density of $m\text{-Ga}_2\text{O}_3$ was significantly higher at $18.4\text{ }\mu\text{mol}\cdot\text{m}^{-2}$, with a weak:strong acid site ratio of 0.40 akin to 0.5Ga/HZSM-5, however the absolute Ga loading was far lower than any of the xGa/HZSM-5 materials.

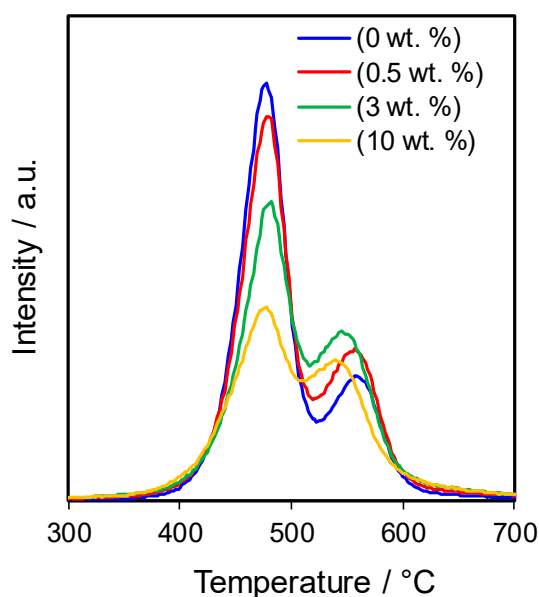


Figure 4. Reactively-formed propene from propylamine temperature-programmed reaction spectroscopy (TPRS) over xGa/HZSM-5.

2.2. Catalytic Activity in Ketonisation

Vapour phase acetic acid ketonisation was subsequently studied over xGa/HZSM-5 in a fixed-bed continuous flow reactor. Turnover frequencies (TOFs) were derived by normalising the steady state rate of acetic acid conversion to the acid site loadings from Table 1. At $350\text{ }^\circ\text{C}$, TOFs were almost independent of Ga loading, exhibiting only a small increase for 10Ga/HZSM-5. Increasing the reaction temperature to $400\text{ }^\circ\text{C}$ increased TOFs for all catalysts as previously reported [19,71], with a monotonic rise with Ga loading now apparent (Figure 5). Catalytic reactivity mirrored the weak:strong acid site ratio for both reaction temperatures, indicating that ketonisation preferentially occurs over weak acid sites within xGa/HZSM-5. Limited deactivation ($<15\%$) was observed for 5 h on-stream for all xGa/HZSM-5 catalysts (Figure S6), attributed to pore/site-blocking by coke or strongly bound

bidentate carboxylate species [72], or structural changes, whereas the Ga₂O₃ reference exhibited minimal deactivation. Powder XRD revealed negligible change zeolite structure following the reaction (Figure S7), however elemental analysis confirmed the presence of surface carbon post-reaction for all xGa/HZSM-5 catalysts (falling from 12 wt% for the parent HZSM-5 and xGa/HZSM-5 samples to only 1 wt% for Ga₂O₃, Table S2).

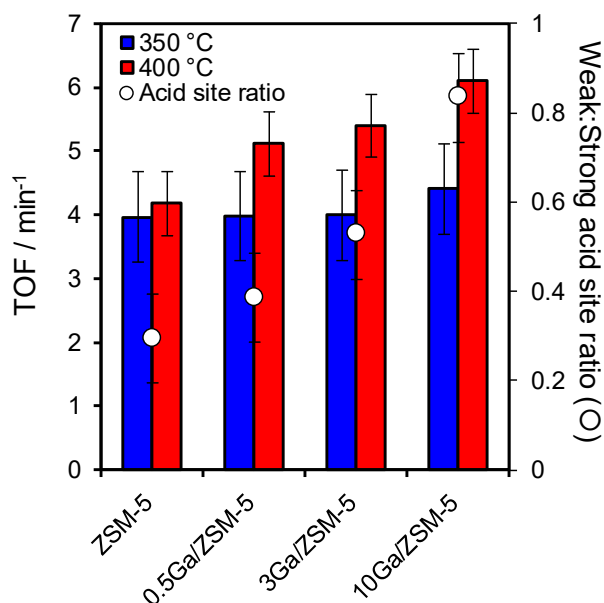


Figure 5. Turnover frequencies (TOFs) for acetic acid ketonisation over xGa/HZSM-5 and corresponding weak:strong acid site ratio. Reaction conditions: 200 mg catalyst, 0.2 mL·min⁻¹ acetic acid, 50 mL·min⁻¹ N₂, 1 bar.

Acetone selectivity at iso-conversion increased with Ga loading at both 350 °C and 400 °C (Figure 6), concomitant with the rise in weak:strong acid site ratio and Lewis acidity [73]. Vervecken also reported an increase in acetone selectivity >350 °C for acetic acid ketonisation over HZSM-5(100) [42], attributed to a higher activation energy for ketonisation that competing aromatisation (which forms xylenols, phenolics and other aromatics). The maximum acetone selectivity for 10Ga/HZSM-5 was 30%; the principal by-products were CO₂, xyleneol, phenol and hydrocarbons [42]. The observation that weak Lewis acid sites and/or related acid-base pairs are the active species for vapour phase acetic acid ketonisation (Figure S8) is consistent with previous experimental [74–77] and computational studies [72]. As discussed in the Introduction, acidic protons in zeolites promote the formation of surface acyl species, which may couple with carboxylate species formed over weaker acid sites to yield an acid anhydride intermediate which in turn decomposes to liberate CO₂ and acetone [25]. However, Chang et al report that ketonisation over HZSM-5 occurs via nucleophilic attack of an acylium ion by carboxylate species [48]; the acylium ion being formed by acid protonation and dehydration [48]. In the case of xGa/HZSM-5, Ga loadings >10 wt% may further increase acetone productivity (and to a lesser extent selectivity) at lower reaction temperature. Although all xGa/HZSM-5 catalysts were stable for 5 h on-stream at 400 °C, future extended ageing and recycling tests are required to optimise formulation and performance.

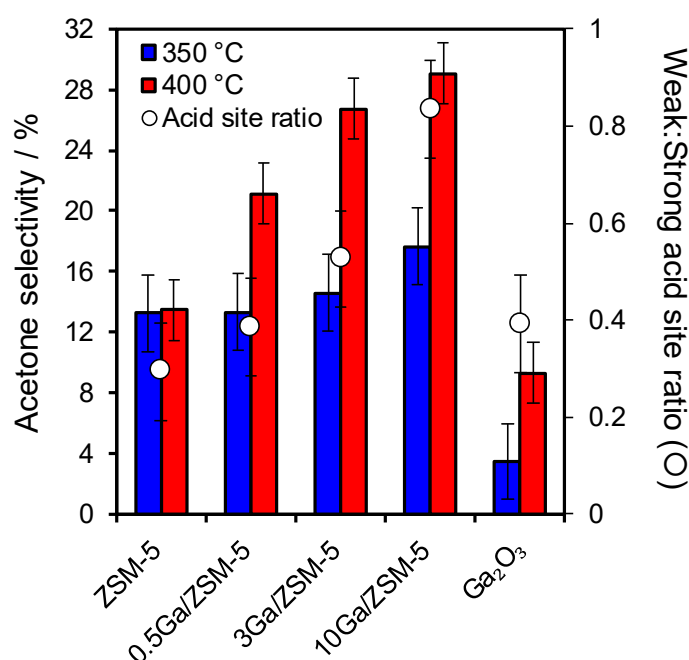


Figure 6. Correlation between acetone selectivity from acetic acid ketonisation at iso-conversion (23% and 29% at 350 °C and 400 °C, respectively) and weak:strong acid site ratio for xGa/HZSM-5. Reaction conditions: 200 mg catalyst, 0.2 mL·min⁻¹ acetic acid, 50 mL·min⁻¹ N₂, 1 bar.

3. Materials and Methods

3.1. Catalyst Synthesis

A commercial HZSM-5 (SiO₂:Al₂O₃ = 30, Zeolyst International, CBV 3024E) was first calcined in air at 550 °C for 4 h to remove any surface residues. Then 3 g of calcined HZSM-5 was subsequently wet-impregnated with appropriate concentrations of a 20 mL aqueous solution of Ga(NO₃)₃·xH₂O (crystalline, 99.9% trace metals basis, Sigma-Aldrich, UK) to prepare Ga gallium zeolites with nominal loadings of 0.5 wt% (0.005 M), 3 wt% (0.03 M) and 10 wt% (0.1 M). In each case the resulting slurry was stirred for 6 h at ambient temperature, dried overnight at 90 °C, and finally calcined at 500 °C under static for 4 h. The resulting catalysts were designated xGa/HZSM-5 where x is the nominal Ga loading. Bulk Ga₂O₃ (≥99.99% trace metals basis, Sigma-Aldrich, UK) was also calcined at 500 °C for 4 h as a reference material.

3.2. Catalyst Characterisation

The bulk Ga loading was determined by elemental analysis using a Thermofisher iCAP 7000 ICP-OES (Thermofisher, UK). Identification of crystalline phases was performed using a Bruker D8 Advance powder X-ray diffractometer (Bruker, UK) with Cu K_α radiation for angles between 2θ = 10–80° with a step size of 0.04°. Volume averaged particle sizes were estimated from the Scherrer equation using the peak width of characteristic HZSM-5 and Ga₂O₃ reflections at 2θ = 14.8° and 35.2° respectively. Surface areas, pore size distributions and mesopore volumes were determined by N₂ porosimetry using a Quantasorb Nova 4000 e porosimeter and Novawin 11.03 software (Quantachrome, UK). Samples were outgassed in vacuo at 300 °C for 18 h according to Quantachrome recommendations for microporous zeolites prior to analysis, with specific surface areas calculated by applying the Brunauer–Emmet–Teller (BET) model over the range P/P₀ = 0.02–0.07 of the adsorption isotherm. Micropore volumes were determined using the t-plot method developed by Lippens and de Boer over the range P/P₀ = 0.2–0.5. X-ray photoelectron spectroscopy (XPS) measurements were performed using a Kratos Axis HSi photoelectron spectrometer (Kratos Analytical, UK) equipped with a charge neutraliser and a monochromated Al K_α X-ray source (hν = 1486.7 eV). Spectra were recorded using

an analyser pass energy of 20 eV and X-ray power of 225 W at a normal emission. Spectral fitting was performed using CasaXPS version 2.3.14 (Casa Software Ltd, UK), with binding energies corrected to the C 1s peak at 284.6 eV; high resolution C and O 1s, Ga, Al and Si 2p XP spectra fitted using a common Gaussian/Lorentzian line shape. Spectra were Shirley background-subtracted and surface compositions quantified by application of element- and instrument-specific response factors. Errors in surface composition were estimated by varying the background subtraction procedure across reasonable limits and re-calculating fits. The carbon content of spent catalysts was measured using a Thermo Scientific Flash 2000 organic elemental analyser (ThermoFisher, UK) calibrated to sulfanilamide, fitted with a Cu/CuO CHNS quartz tube and a thermal conductivity detector. Samples were prepared by adding ~10 mg catalyst and ~2 mg V₂O₅ to tin crucibles.

Acid site loadings and strength were determined by n-propylamine (Sigma Aldrich, UK, ≥99%) temperature-programmed reaction spectroscopy (TPRS) using a Mettler Toledo TGA/DSC 2 STARE system (Mettler Toledo, UK) connected to a Pfeiffer Vacuum ThermoStar GSD 301 T3 (Pfeiffer, UK) benchtop mass spectrometer (MS). Propylamine adsorption was performed by adding liquid n-propylamine to pre-weighed samples (1 mL per 20 mg) and placing in an alumina crucible. Excess physisorbed propylamine was removed by drying in vacuo at 30 °C for 1 h prior to analysis. Samples were heated in the thermogravimetric analyser (TGA) from 40 °C to 800 °C at a ramp rate of 10 °C.min⁻¹ under flowing N₂ (40 mL.min⁻¹), with evolved gases analysed by MS to monitor reactively formed propene. Lewis/Brønsted character was determined by diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) of samples following pyridine adsorption over diluted samples (10 wt% in KBr). Excess physisorbed pyridine was removed in vacuo at 30 °C overnight prior to room temperature measurement using a Nicolet Avatar 370 MCT (ThermoFisher, UK) with Smart Collector accessory and liquid nitrogen cooled mercury cadmium telluride (MCT-A) detector. DRIFTS spectra were background-subtracted, and the ratio of the transmitted intensities of the 1450 cm⁻¹ and 1540 cm⁻¹ peaks used to quantify the ratio between Lewis and Brønsted acid sites.

3.3. Catalytic Ketonisation

Acetic acid ketonisation was performed in a bespoke continuous flow packed-bed reactor with online gas chromatography (GC) analysis. The reactor comprised a 1 cm o.d., quartz tube, within which the catalyst bed was placed centrally and retained by quartz wool plugs. A constant catalyst bed volume of 4 cm³ was used in all experiments, comprised of 200 mg of catalyst diluted with fused silica granules. The reactor tube was positioned in a temperature-programmable furnace with a thermocouple placed in contact with the catalyst bed. Acetic acid (Sigma-Aldrich, UK, ACS reagent ≥ 99.7%) was fed in a down-flow fashion into the reactor using an Agilent 1260 Infinity Isocratic Pump (Agilent, UK) and N₂ as the carrier gas (50 mL.min⁻¹). All reactor lines were heated to 130 °C to prevent condensation, and a 1 cm diameter metal tube packed with fused silica granules was used to ensure acetic acid vaporisation before the reactor. Products were analysed online by a Varian 3800 GC (Varian, UK) with heated gas-sampling valve, equipped with a BR-Q PLOT column (30 m × 0.53 mm i.d.). Acetone and acetic acid were detected using a flame ionisation detector (FID). The GC was calibrated for acetic acid and acetone by triplicate injections of 50 µl standard solutions through a split/splitless injector. Acetic acid conversion and selectivity were calculated according to Equations (1) and (2).

$$\text{Conversion} = \frac{n_{\text{AcOH}_0} - n_{\text{AcOH}}}{n_{\text{AcOH}_0}} \times 100 \quad (1)$$

$$\text{Selectivity} = \frac{2 \times n_{\text{Acetone}}}{n_{\text{AcOH}_0} - n_{\text{AcOH}}} \times 100 \quad (2)$$

where n_{AcOH_0} is the initial moles of acetic acid, n_{AcOH} is the final moles acetic acid and n_{Acetone} represents the moles of produced as acetone. Acetone was the primary product over all catalysts. No acetic acid conversion was observed in the absence of a catalyst.

4. Conclusions

A family of Ga-doped HZSM-5 materials were synthesis by wet impregnation as solid acid catalysts for the vapour phase ketonisation of acetic acid, a potential route to upgrading pyrolysis bio-oil vapours. XRD indicates that Ga is either incorporated into the parent zeolite framework or highly dispersed across the zeolite surface as GaO^+ clusters for loadings spanning 0.3–10 wt% Ga doping has little impact on the zeolite textural properties but increased the Lewis acid character concomitant with a decrease in acid strength relative to HZSM-5. Turnover frequencies for acetic acid ketonisation, and acetone selectivity at iso-conversion, were both proportional to the weak:strong acid site ratio, evidencing that ketonisation over xGa/HZSM-5 preferentially occurs over weak (Lewis) acid sites. The most active catalyst was 10 wt% Ga/HZSM-5, which was stable for 5 h on-stream despite significant carbon laydown, with an acetone selectivity of 30%. Acetic acid ketonisation is an attractive route to upgrading biomass pyrolysis vapours through close-coupling with Ga/HZSM-5 catalysts derived from earth abundant elements.

Supplementary Materials: The following are available online at <http://www.mdpi.com/2073-4344/9/10/841/s1>, Figure S1: XRD patterns of xGa/HZSM-5 and bulk Ga_2O_3 ; Figure S2: N_2 adsorption-desorption isotherms of xGa/HZSM-5 and Ga_2O_3 ; Figure S3: (a) Al 2p and (b) Si 2p XP spectra of xGa/HZSM-5; Figure S4: (a) DRIFT spectra of pyridine-saturated xGa/HZSM-5 and Ga_2O_3 and (b) corresponding Lewis:Bronsted acid site ratio (1444 cm^{-1} : 1545 cm^{-1} bands) for xGa/HZSM-5; Figure S5: Density of strong and weak acid sites for xGa/HZSM-5 from propylamine TPRS; Figure S6: Acetic acid conversion over xGa/HZSM-5, and Ga_2O_3 vs time on stream. Reaction conditions: 200 mg catalyst, at $400\text{ }^\circ\text{C}$, $0.2\text{ mL}\cdot\text{min}^{-1}$ acetic acid, $50\text{ mL}\cdot\text{min}^{-1}$ N_2 , 1 bar; Figure S7: XRD patterns of (a) fresh and (b) post-reaction xGa/HZSM-5 and Ga_2O_3 ; Figure S8: Correlation between acetone selectivity from acetic acid ketonisation at iso-conversion (23% and 29% at $350\text{ }^\circ\text{C}$ and $400\text{ }^\circ\text{C}$, respectively) and acid strength from propylamine temperature-programmed reaction spectroscopy (higher temperatures indicate weaker acidity) over xGa/HZSM-5, and Ga_2O_3 ; Table S1: Surface and bulk composition of xGa/HZSM-5 and Ga_2O_3 ; Table S2: Carbon content of used xGa/HZSM-5 and Ga_2O_3 after 5 h reaction.

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