The effect of Mg location on Co-Mg-Ru/γ-Al₂O₃ Fischer–Tropsch catalysts

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The effectiveness of Mg as a promoter of Co-Ru/γ-Al₂O₃ Fischer–Tropsch catalysts depends on how and when the Mg is added. When the Mg is impregnated into the support before the Co and Ru addition, some Mg is incorporated into the support in the form of MgₓAl₂O₃₋ₓ if the material is calcined at 550°C or 800°C after the impregnation, while the remainder is present as amorphous MgO/MgCO₃ phases. After subsequent Co-Ru impregnation MgₓCo₃₋ₓO₄ is formed which decomposes on reduction, leading to Co(0) particles intimately mixed with Mg, as shown by high-resolution transmission electron microscopy. The process of impregnating Co into an Mg-modified support results in dissolution of the amorphous Mgₓ, and it is this Mg which is then incorporated into MgₓCo₃₋ₓO₄. Acid washing or higher temperature calcination after Mg impregnation can remove most of this amorphous Mgₓ, resulting in lower values of x in MgₓCo₃₋ₓO₄. Catalytic testing of these materials reveals that Mg incorporation into the Co oxide phase is severely detrimental to the site-time yield, while Mg incorporation into the support may provide some enhancement of activity at high temperature.
1. Introduction

Fischer–Tropsch synthesis (FTS) is an increasingly important industrial reaction for the production of hydrocarbons from a mixture of carbon monoxide and hydrogen, known as syngas. FTS can be used to convert cheap gas sources, such as shale gas, to exploit stranded gas reserves, to prevent gas flaring during oil extraction or to convert feedstocks such as coal and biomass to higher value liquid hydrocarbons [1–5]. The resulting hydrocarbons after work-up comprise a mixture of liquid hydrocarbons with low sulfur and aromatic content, therefore giving access to clean-burning, synthetic fuels. Hence, FTS is a key step in gas-to-liquid, coal-to-liquid and biomass-to-liquid technologies.

Cobalt catalysts for FTS generally consist of the active metal dispersed as an oxide over an oxidic support with various quantities of metal oxide or platinum group metal promoters to improve the reducibility of the active phase. As it is widely accepted that FTS proceeds over Co metal sites [6], prior to use, catalysts are usually activated with H₂ in situ to reduce the Co oxides to Co metal.

Magnesium has been used numerous times as an additive to FTS catalysts. Indeed, some early work used Mg as a promoter in a Co/MgO/ThO₂/kieselguhr catalyst [7]. Most studies have used Mg to improve the mechanical strength and hence attrition resistance by the formation of spinels [8]. However, a study by Zhang et al. [9] found that Mg loaded at 0.8 wt% on γ-Al₂O₃ suppressed the formation of difficult to reduce Co-surface phases. The effect was thought to be due to the formation of MgAl₂O₄ spinel on the surface of the γ-Al₂O₃, which reduced the contact between Co and γ-Al₂O₃ and therefore the amount of cobalt aluminate spinel, which is not FT-active. Hence, Mg at low loadings could increase reducibility and activity. The same study found that more than 1 wt% MgO decreased the reducibility due to the formation of an MgO–CoO solid solution, which required high-temperature reduction. Chernavskii et al. [10] also found an increase in reducibility when Mg was loaded at 0.2 wt% on an γ-Al₂O₃ support. A study by Niemelä et al. [11] found that MgO-promoted Co/SiO₂-supported catalysts reduced the X-ray diffraction (XRD) particle size of Co metal particles from 14 nm to around 10 nm and decreased reducibility presumably due to the smaller particle size. It should be noted that Co/SiO₂ catalysts tend to be more reducible than Co/γ-Al₂O₃ catalysts and therefore a decrease in reducibility of MgO-promoted SiO₂ may be expected. The study by Niemelä also found that activity was higher when Mg was impregnated after Co onto the support rather than before. A more recent study by Holmen’s group observed a decrease in activity for Mg-promoted Co/γ-Al₂O₃ catalysts which could not be explained by particle size or dispersion effects [12]. The authors suggest that Mg could have a poisoning effect similar to that of alkali metals. Work by our group using proxy-based accelerated discovery methods has shown that the order of addition of the Mg has a large effect on the size of the Co particles formed and also on the activity and stability of the catalysts [13]. Mg in general reduces particle size and activity, while adding Mg before the other elements and calcining at 550°C produces catalysts with high stability.

In this work, we aim to identify the origin of the effects on particle size, activity and stability. We conduct detailed characterization of Mg-promoted catalysts, and a series of experiments aimed at controlling the location of Mg in the catalysts to better understand the effects.

2. Experimental set-up

Catalysts were synthesized by an incipient wetness impregnation process. The support, γ-Al₂O₃ (Sasol; Puralox TH100/150, 145 m² g⁻¹), was dried at 120°C overnight. The impregnation was carried out in plastic bags (Tesco; zip seal sandwich bags, 18 × 18 cm). The impregnation was a multi-step process, the first step being the impregnation of magnesium nitrate hexahydrate (Fluka; greater than or equal to 99%) followed by heat treatment at either 550°C or 800°C, then the impregnation of the cobalt nitrate hexahydrate (Acros; 99%) and ruthenium (III) nitrosyl nitrate (Alfa Aesar; min 31.3% Ru), followed by a final heat treatment at 250°C or 400°C. To the γ-Al₂O₃ was added the appropriate amount of hot aqueous nitrate solution (ca 60°C) to achieve the desired
loading. The impregnated solid was then kneaded in the plastic bag until homogeneous. For thermal treatments, the damp solid was spread out on a stainless steel tray and treated in static air in a muffle furnace. After thermal treatment, subsequent impregnations were carried out using the same procedure. The volume used for the initial impregnation was 1 ml g\(^{-1}\) for unmodified Puralox TH100/150, and the volume required was recalculated after each impregnation.

Some of the Mg-modified supports were subjected to an acid wash. Two different solutions with different pH were used to wash the Mg/γ-Al\(_2\)O\(_3\) materials—acetic acid (1 mol dm\(^{-3}\), pH 2.4) and an acetic acid–ammonium acetate buffer (0.5 mol dm\(^{-3}\), pH 4.7). Acetic acid was chosen as it has a similar pH to Co(NO\(_3\))\(_2\) (ranges from approx. 3 at 1 M concentration to approx. 2 at 4 M) and the buffer solution was chosen to determine whether a milder treatment could produce the same results. Additionally, residual species left after the washes could be removed by appropriate heat treatments, unlike for instance acetic acid–sodium acetate buffers. The washing procedure was thus:

To each Mg/γ-Al\(_2\)O\(_3\) sample (1 g) contained in a centrifuge tube (50 ml) was added the weak acid solution (40 ml) followed by shaking for 30 s. Tubes were then centrifuged for 30 min at 4000 r.p.m. before removing the supernatant with a pipette. In some cases, the Mg and Al content of the supernatant was analysed by inductively coupled plasma (ICP) analysis. The wet solid was washed with water (40 ml) via shaking by hand for 30 s before returning to the centrifuge for 30 min at 4000 r.p.m. The supernatant was removed by pipette before drying the wet solid for 18 h at 80°C. The dry solid was calcined at 550°C for 2 h in static air to remove residual acetate or ammonium ions.

XRD data on samples from Boldrin et al. [13] (supports and catalysts) and on the catalysts in the acid washing study described above were collected on a Panalytical X-Pert diffractometer fitted with a X-Celerator detector and a spinner stage set up in Bragg–Brentano geometry. X-rays were produced with a Co anode filtered by Ge giving monochromatic K\(_{\alpha1}\) radiation. Variable divergence slits were used with an illuminated length of 15 mm. The measuring time for XRD scans in the 2\(\theta\) range 18°–82° was 1 h. Samples were spiked with KCl in order to more accurately calculate the lattice parameters of the Co\(_3\)O\(_4\) phase. Two methods were used to extract quantitative information from XRD patterns: empirical peak fitting and Le Bail whole pattern fitting. Empirical peak fitting was used to calculate particle size from the Scherrer equation while Le Bail fitting was used to calculate the lattice parameters of Co\(_3\)O\(_4\) phases.

TOPAS 4.2 was used to refine the patterns of the Co\(_3\)O\(_4\)/γ-Al\(_2\)O\(_3\) samples. In addition to the background polynomial, the lattice parameters and effective particle size parameters were refined, although determination of instrumental broadening was not performed so these particle size measurements are not discussed as they would lead to a false impression of their accuracy. Where particle size measurements are quoted, these are calculated from single peaks using the Scherrer equation.

In addition to this, the acid-washed (AW) supports were characterized using the high-throughput XRD technique described by Boldrin et al. [13]. Samples were placed in well-plates, adhesive tape was stuck over the tops of all wells and the plates were upturned to stick the sample to the tape to ensure that the sample height was the same across all wells. High-throughput XRD was performed in reflection mode on a Panalytical X-Pert Pro diffractometer fitted with an XYZ stage using Co K\(_{\alpha1}\)/K\(_{\alpha2}\) (ratio = 2) radiation, Fe filter, 5 mm mask and automatic divergence slits with an illuminated length of 3 mm. The Soller slits used were 0.02° on the incident side and 0.04° on the reflection side. The measuring time for an XRD scan in the 2\(\theta\) range 38°–57° was 13 min.

Fourier transform infrared (FTIR) spectra were recorded for Mg-modified γ-Al\(_2\)O\(_3\). Spectra were collected from 600 to 4000 cm\(^{-1}\) with a resolution of 4 cm\(^{-1}\) using a Perkin Elmer Spectrum 100 spectrometer fitted with the Spectrum 100 Universal Diamond/ZnSe ATR.

Temperature-programmed reduction (TPR) was carried out on an Altamira AMI-200 unit. Samples (70 mg) were purged with flowing argon at 30 ml min\(^{-1}\) before the gas was switched to 10% H\(_2\)/Ar at 30 ml min\(^{-1}\). Samples were ramped from room temperature to 1100°C at 10°C min\(^{-1}\), holding at 1100°C for 10 min. Five injections were performed at the end of the experiment, which allowed the peak area to be related to H\(_2\) consumption.
Thermogravimetric analysis (TGA) scans were carried out on a TA Q5000IR thermal analyser using 5% H$_2$/N$_2$ as the sample gas flowing at 25 ml min$^{-1}$. Samples (ca 5 mg) were placed in aluminium pans. The dynamic high-resolution mode was used with a ramp rate of 50$^{\circ}$C min$^{-1}$, sensitivity of 1.0 and a resolution of 4.0. Samples were heated from room temperature to 600$^{\circ}$C, where the temperature was held for 5 min. The maximum temperature was limited to 600$^{\circ}$C due to the use of aluminium pans.

CO$_2$ temperature-programmed desorption (TPD) was carried out on an Altamira AMI-200 unit. Samples (250 mg) were first heated to 250$^{\circ}$C at 10$^{\circ}$C min$^{-1}$ and purged with flowing He at 30 ml min$^{-1}$ for 10 min before being cooled to 35$^{\circ}$C at 10$^{\circ}$C min$^{-1}$. Subsequently, the gas was switched to 10% CO$_2$/He at 30 ml min$^{-1}$ for 1 h at 35$^{\circ}$C followed by 30 ml min$^{-1}$ He for 30 min. Finally, the sample was heated to 800$^{\circ}$C at 10$^{\circ}$C min$^{-1}$ in He flowing at 10 ml min$^{-1}$, holding at 800$^{\circ}$C for 10 min. Five injections were performed at the end of the experiment, which allowed the peak area to be related to CO$_2$ desorption.

Co surface areas were determined by H$_2$ chemisorption at 150$^{\circ}$C in a Micromeritics ASAP 2020C by extrapolating the total gas uptakes in the H$_2$ adsorption isotherms at zero pressure. Prior to adsorption, the samples (ca 0.5 g) were pre-treated in flowing He at 120$^{\circ}$C for 1 h. Afterwards, the samples were reduced in situ by flowing pure H$_2$ and raising the temperature to 425$^{\circ}$C and maintaining this temperature for 6 h. After reduction, the samples were degassed, the temperature lowered to 150$^{\circ}$C and H$_2$ dosed over the sample in the pressure range of 100–760 mm Hg.

Transmission electron microscopy was performed to calculate Co particle size distributions and to produce elemental maps. Samples were first reduced and passivated. Powder samples were embedded in resin, cured, microtomed and placed on holey carbon-coated Cu grids. The samples were examined in the probe-corrected JEOL-2100F transmission electron microscope (TEM) at the University of Birmingham, UK, using the scanning transmission mode with high-angle annular dark-field and bright-field detectors and a voltage of 200 kV. Parallel electron energy loss spectroscopy (EELS) using a GATAN Enfina spectrometer was used to investigate the composition.

ICP analysis was performed to determine the compositions of solid samples. Following complete dissolution of the samples using microwave digestion in HCl, analysis was performed on a Perkin Elmer Optima ICP-OES.

ICP analysis was performed to determine the amount of Mg and Al ions removed from Mg-modified γ-Al$_2$O$_3$ solids by washing with acetic acid or ammonium acetate on a Ciros CCD optical emission spectrometer.

The FTS testing was carried out in a stainless steel isothermal fixed bed six-way micro-reactor connected to an online Varian CP-3800 gas chromatograph with three detectors. A Valco-valve was used to select between the gas streams for online analysis. Catalysts were reduced at 425$^{\circ}$C for 9 h in pure H$_2$ prior to testing. After reduction, the reactor was cooled to 160$^{\circ}$C and H$_2$ was replaced with syngas (H$_2$ : CO : Ar = 2 : 1 : 0.1) and pressure was raised to 20 bar. Conversion was measured by the consumption of the syngas mix (CO + H$_2$) relative to Ar. Wax and water were collected from the gas–liquid separators and analysed using an offline gas chromatogram with a flame ionization detector on a SimDist column in order to calculate the chain growth probability, $\alpha$, from the C$_{22}$–C$_{40}$ fraction.

3. Results and discussion

In our previous work [13], we described the high-throughput discovery, scale-up and testing of a series of Co-Ru catalysts supported on Mg-modified γ-Al$_2$O$_3$. The Mg was found to have several benefits: reducing the Co particle size, improving the Co surface area and increasing the stability of the catalysts during FTS. There were also detrimental effects on the reducibility of the catalysts and the initial activity. In order to investigate the origin of these effects, detailed characterization was carried out on both the initial catalysts and the modified supports.

XRD analysis of the calcined catalysts (electronic supplementary material, figure S1) showed the presence of two phases, one assigned to the γ-Al$_2$O$_3$ support and one assigned to Co$_3$O$_4$. 

Table 1. XRF and XRD data for catalysts from Boldrin et al. [13]. LP, lattice parameter; PS, particle size. Amount of Mg incorporated into Mg$_x$Co$_{3-x}$O$_4$ spinel was calculated by comparison with literature values reported by Krezhov & Konstantinov [16]. This can then be used to estimate the mass of the Mg in the Mg$_x$Co$_{3-x}$O$_4$ and, by subtraction from the total mass of Mg, the amount unaccounted for, described as 'other Mg' in the final column.

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Le Bail refinements were undertaken with peaks inserted empirically for the alumina phase due to the difficulty in assigning a structural model to this phase [14,15], with the results shown in table 1. The lattice parameter of the Co3O4 phase for the 0% Mg sample was slightly smaller than expected for bulk Co3O4. This slight contraction of the unit cell could be the result of the small size (14.7 nm) of the Co3O4 particles causing deviations from bulk properties, for instance, due to the presence of vacancies or other crystal imperfections [17]. As the Mg loading increased, the Co3O4 lattice parameter increased. By comparison with Mg0.05Co2.07O4, which is reported to have a lattice parameter of 8.1390 Å [16], and considering that Mg2+ has a larger ionic radius than Co2+ (86 pm versus 79 pm in octahedral sites [18]) this can be attributed to incorporation of Mg into the Co3O4 lattice forming a solid solution with general formula MgxCo3−xO4. This implies that, prior to Co(NO3)2 incipient wetness impregnation, there are Mg species which are sufficiently reactive to form a mixed oxide with Co ions. By comparison with the literature data on the lattice parameter of MgxCo3−xO4 compounds [17], an estimate of the amount of Mg in the MgxCo3−xO4 spinel relative to the total amount of Mg can be calculated. For the 6% Mg sample, the composition of the spinel was Mg0.76Co2.24O4, which means that 62% of the total magnesium in the sample is incorporated in this phase. The particle sizes calculated from the broadening of the Co3O4 peaks indicate that catalysts containing Mg had smaller Co3O4 particles than the catalyst without Mg, which agrees with our previous work [13].

Although a Le Bail refinement could not be performed on the γ-Al2O3 phase, information relating to the phase can be gathered by simply looking at the shifts in the peaks. Although somewhat crude, by assuming that γ-Al2O3 has the defect spinel structure, a lattice parameter was calculated by measuring the peak position of a single reflection at approximately 53.6° 2θ (table 1). It can be seen that, as the Mg loading increased, the γ-Al2O3 peaks shifted to the lower 2θ position, indicating an expansion of the interplanar distances and hence an increase in the defect spinel lattice parameter. This indicates that Mg was incorporating into the γ-Al2O3 structure to form a phase with general formula Mg_xAl2O_{3+y} since the defect spinel structure for γ-Al2O3 has a lattice parameter of 7.9 Å and the lattice parameter for the spinel MgAl2O4 is 8.0824 Å (reference pattern: ICDD 021-1152). The modified supports were also characterized by XRD to examine the crystalline phases present before Co(NO3)2 impregnation. This analysis confirmed that the shift in lattice parameter of the γ-Al2O3 phase observed in the Co-loaded catalysts was primarily due to Mg incorporation (electronic supplementary material, figure S2). All XRD patterns only contained peaks due to the Mg_yAl2O_{3+y} phase except for the sample with 6% Mg calcined at 550°C, which also contained peaks due to MgO. The emergence of the MgO peaks only at a loading of 6% Mg indicates that below 6% Mg the MgO particles were too small or insufficiently crystalline to be detected by XRD under the experimental set-up. Because peaks due to MgO were seen in this sample before Co(NO3)2 impregnation and calcination but not seen after, this indicates that MgO was highly dispersed or poorly crystalline. The proportion of MgO was calculated by measuring the peak position of a single reflection at approximately 53.6° 2θ and is therefore likely to be from an amorphous or highly dispersed Mg-containing phase.

It is evident from the XRD data that, in the pre-activated catalyst, Mg is incorporated into the Co3O4 and the γ-Al2O3 structures. From the measurement of χ in Mg_xCo3−χO4, it is possible to calculate the proportion of Mg which is present in the Mg_xCo3−χO4 phase, and, by subtraction, the remaining Mg which is not accounted for by this (given as ‘other Mg’ in table 1). If all the remaining Mg was incorporated in the γ-Al2O3 phase to form Mg_yAl2O_{3+y}, we would expect the lattice parameter of this phase to increase proportionally to the ‘other Mg’ content, and this does somewhat seem to be the case, with a linear regression giving an r^2-value of 0.966, although with only four points this is inconclusive. The absence of any peaks due to MgO or other Mg-containing phase (e.g. MgCO3) indicates that any Mg which was not incorporated into Mg_xCo3−χO4 or Mg_yAl2O_{3+y} was highly dispersed or poorly crystalline. The proportion of the total Mg which is incorporated into the Mg_xCo3−χO4 phase increases with increasing Mg loading.
Figure 1. (a) FTIR spectra for Mg/γ-Al2O3 modified supports with different Mg loadings calcined at 550°C. Unmodified γ-Al2O3 is shown for comparison. (b) FTIR spectra for Mg/Al2O3 modified supports with different Mg loadings calcined at 800°C. Insets show close-ups of the 650–1000 cm⁻¹ region, where modes due to Al–O bonds are found.

Mg/γ-Al2O3 samples were calcined at a higher temperature of 800°C to provide insight into the behaviour of the samples calcined at the lower temperature of 550°C. The XRD patterns showed a single Mg₉ₓAl₂O₃+ₓ phase for all compositions and, again, an increase in the lattice parameter with Mg loading was observed (electronic supplementary material, figure S2). However, for these samples, the increase in the lattice parameter was greater than that at the lower calcination temperature. This indicates that more Mg was incorporated into the γ-Al₂O₃ structure, presumably due to the higher temperature facilitating more diffusion into the bulk. As the lattice parameter increased more at 800°C, this implies that after calcination at 550°C there are Mg species which are XRD-silent and not incorporated into the Mg₉ₓAl₂O₃+ₓ phase. As this available Mg was not seen by XRD for samples other than 6% Mg calcined at 550°C this means that it is either poorly crystalline or highly dispersed, perhaps as a monolayer as suggested previously [19]. It is these XRD-silent Mg species which are the likely origin of the Mg in MgₓCo₃₋ₓO₄.

FTIR spectroscopy on the modified supports (figure 1) can give information about both the XRD-silent Mg species and the Mg₉ₓAl₂O₃+ₓ phase. As expected, every sample had major bands due to O–H (ca 3500 cm⁻¹), which is mainly due to adsorbed water in the pore network but may also have a contribution from hydroxide species. The presence of a peak at ca 1385 cm⁻¹ in the samples calcined at 550°C is indicative of carbonate ions. The intensity of this peak increased with Mg loading, indicating that MgCO₃ was present. This peak is absent in the samples calcined at 800°C. CHN analysis confirms the presence of carbon in all the modified supports calcined at 550°C, and the two highest loadings of Mg in the supports calcined at 800°C (electronic supplementary material, figure S3). The amount of carbon increases with Mg loading, reinforcing the hypothesis that the peaks in the FTIR are due to MgCO₃. CHN analyses conducted over time show an increasing carbon content with time since synthesis (electronic supplementary material, figure S4), suggesting that initially the XRD-silent Mg species are amorphous oxides, which convert to carbonates over time in air.

Examination of the low-energy region of the FTIR spectrum reveals peaks typically due to metal–oxygen bonds and gives information about the Mg₉ₓAl₂O₃+ₓ phase. For the samples calcined at 550°C, there is a peak at ca 824 cm⁻¹ which is attributed to tetrahedral Al–O bonds [20]. As the Mg loading increased the intensity of this band decreased. A similar trend was observed for the samples calcined at 800°C but to a greater extent (figure 1b). Furthermore, for the samples calcined at 800°C, an additional band at ca 683 cm⁻¹, attributed to octahedral Al–O bonds [21], increased in intensity as Mg loading increased. These changes in intensity with Mg loading indicate that, as the Mg loading increases, there are fewer Al³⁺ ions in tetrahedral sites and more in octahedral sites. In a normal spinel such as MgAl₂O₄, the divalent cation resides...
in tetrahedral sites and the trivalent cation is in octahedral sites. Although the structure of γ-Al2O3 is similar to spinel, it is often reported as having many defects, non-spinel sites and with a proportion of the Al3+ cations in tetrahedral sites rather than octahedral [15,21]. The FTIR data suggest that, as the Mg loading and calcination temperature was increased, the Al3+ cations’ occupation of crystallographic sites resembles more closely that of bulk MgAl2O4 spinel. Presumably, as the Mg loading increases, more Mg2+ is incorporated into the tetrahedral sites of the γ-Al2O3, which results in fewer Al3+ cations in the tetrahedral sites and more in the octahedral sites. This move towards a spinel-type MgγAl2O3+γ structure is consistent with the XRD data.

Further information about the nature of the supports can be deduced from CO2 TPD (electronic supplementary material, figure S5). The unmodified support possesses a single CO2 desorption peak at around 100°C, indicating the presence of a weakly basic site, as is expected from γ-Al2O3 [15,19]. This is also the case with all the modified supports calcined at 800°C, as well as the ones calcined at 550°C with an Mg loading of less than 2%. This suggests that incorporation of Mg into the γ-Al2O3 crystal structure does not increase the basicity of the support’s surface. The total amount of CO2 desorbed does not change with Mg loading in these samples, which reinforces this hypothesis. However, modified supports with Mg loadings of 2% and above calcined at 550°C also possessed a more strongly basic site reflected in an additional desorption peak at 300°C. This is likely to be from the XRD-silent Mg species identified previously, especially as, in these samples, the total amount of CO2 does increase with Mg loading.

Comparing the total CO2 desorption in the TPD with the amount of Mg present in each phase in the final catalyst, we can see that the correlation is better with the amount of Mg in the MgγCo3−yO4 phase than with the amount of ‘other’ Mg. Because the total amount of CO2 desorbed is mainly influenced by the XRD-silent Mg, this suggests that the XRD-silent Mg is ending up in the MgγCo3−xO4 phase rather than as ‘other’ Mg in the final catalyst, so the ‘other’ Mg is almost entirely the MgγAl2O3+γ phase (rather than being a combination of Mg in MgγAl2O4 phase and XRD-silent Mg species). This is in agreement with the XRD data, which show a good correlation between the MgγAl2O3+γ lattice parameter and the ‘other Mg’ content, and gives an indication that the cobalt impregnation solution is dissolving the majority of the amorphous Mg, which is then incorporated into the MgγCo3−yO4 phase.

The evidence detailed above strongly suggests that the modification of γ-Al2O3 with Mg has two effects: (i) the creation of an MgγAl2O3+γ phase that is intermediate between γ-Al2O3 and MgAl2O4 in crystal structure, which has a similar basicity to the unmodified support and is unchanged by the impregnation of cobalt, and (ii) the deposition of highly basic XRD-silent MgO or MgCO3 species which are later incorporated into the cobalt oxide spinel forming MgγCo3−yO4. We hypothesized that it is the MgO/MgCO3 species that is dissolved by the acidic Co(NO3)2 impregnation solution and thus forms the MgγCo3−yO4 phase. We therefore attempted to remove this second species by acid washing the modified supports before Co(NO3)2 impregnation, allowing us to separately examine the two effects. The washing solutions chosen were acetic acid (pH 2.4), which has a similar pH to concentrated cobalt nitrate solutions (pH 2–3 in the range of 3–4 M concentrations), and ammonium acetate buffer (pH 4.7) to assess if a milder treatment could achieve the same results.

ICP analysis of the supernatant allowed us to characterize the amount of Mg removed during washing, while the amount left behind is calculated by subtraction from the ICP results of the materials before washing. The results are shown in figure 2a,b (buffer wash) and the electronic supplementary material, figure S6 (acetic acid wash). Washing with either acetic acid (pH 2.4) or ammonium acetate buffer (pH 4.7) resulted in almost identical amounts of Mg being dissolved from the sample for both the supports calcined at 550°C and 800°C, indicating that the two washing solutions were equally effective in dissolving Mg. As would be expected, much more Mg was dissolved from samples calcined at 550°C than from those calcined at 800°C. Interestingly, some Al was also dissolved by both ammonium acetate buffer and acetic acid (figure 2c and electronic supplementary material, figure S7, respectively), including, in the unmodified support, raising the prospect that some Al may also be incorporated into the Co3O4 spinel, an effect which to our knowledge has not been discussed previously in the literature on γ-Al2O3-supported
catalysts. The amount of Al dissolved increases with Mg loading, suggesting that Al in mixed Mg-Al species is more easily dissolved. Less Al is dissolved by the higher pH buffer.

Comparison of the lattice parameters of the Mg$_y$Al$_{2}$O$_{3+y}$ phase from XRD of the washed and unwashed samples (figure 3a,b) shows little change on washing for all samples, indicating that this phase is largely stable towards washing with acid. There is a small decrease in the lattice parameters for the samples calcined at 550°C, and this is reflective of the ICP results given above, which suggest that, although the majority of the dissolved Mg comes from highly dispersed or amorphous Mg-containing phases, some mixed Mg-Al species are also dissolved.

A series of Co-Ru catalysts were made using the washed and unwashed supports calcined at 550°C and 800°C, and the Co$_3$O$_4$ crystallite size was measured using XRD (figure 3c,d). In samples where the support has been calcined at 550°C, Mg loading still has an effect in reducing the particle size, but the effect is reduced compared with unwashed samples. In addition, higher loadings of Mg show a bigger difference in Co particle size between washed and unwashed samples, suggesting that a large part of the particle size reduction is due to the Mg species which are removed by washing. The washed and unwashed supports calcined at 800°C give very similar particle sizes to those seen in the washed samples calcined at 550°C, which again indicates that it is the free Mg which largely causes the decrease in particle size.

TGA in 5% H$_2$ in N$_2$ was carried out on these catalysts. Samples where the supports were calcined at 550°C show clearly that washing with either acid or buffer improves reducibility, as

Figure 2. (a,b) Amount of Mg dissolved or not dissolved from Mg/γ-Al$_2$O$_3$ modified supports calcined after Mg addition at either (a) 550°C or (b) 800°C washed with ammonium acetate buffer. (c) Amount of Al dissolved from Mg/γ-Al$_2$O$_3$ modified supports calcined after Mg addition at either 550°C or 800°C washed with ammonium acetate buffer. Washed, unmodified γ-Al$_2$O$_3$ is shown for comparison.
Figure 3. (a,b) Variation with Mg loading of lattice parameter of Mg/γ-Al₂O₃ modified supports washed with either acetic acid or ammonium acetate buffer for samples calcined after Mg addition at either (a) 550°C or (b) 800°C. Lattice parameters were calculated assuming a cubic spinel structure for γ-Al₂O₃. Lattice parameters of unwashed samples are shown for comparison, (c,d) Co metal particle size calculated from high-throughput XRD [13] for catalysts synthesized using the supports from (a,b) calcined after Mg addition at either (c) 550°C or (d) 800°C. Data shown are the mean of two aliquots.

measured by the temperature maxima for the second reduction peak (Co(II)–Co(0); electronic supplementary material, figure S8). This is in line with the hypothesis that it is the free Mg which is detrimental to reducibility rather than total Mg.

Several samples washed with ammonium acetate buffer solution (henceforth AW) were scaled up for characterization and testing based on these results. Characterization of these samples is shown in table 2. Acetic acid samples were scaled up and characterized, but as there was no obvious difference between the acid- and buffer-washed samples these are not discussed further. Data from the acetic AW samples are shown in the electronic supplementary material, table S1.

XRD analysis showed that the washed catalysts still contained some MgₓCo₃−ₓO₄, indicating that the washing is not fully effective in preventing this phase from forming. It is not clear why the impregnation solution is able to dissolve Mg from the washed supports, possibly because the impregnation solution is hot, or there is some other chemical effect than simply the pH. The washes were effective in reducing the Mg in MgₓCo₃−ₓO₄ by 60% in the 3% Mg samples, and by 80% in the 6% Mg samples. The washes also removed an appreciable fraction of the Mg not incorporated into MgₓCo₃−ₓO₄—around 17% in the 3% Mg samples and 33% in the 6% Mg samples. The calcination at 800°C was more effective at reducing the formation of MgₓCo₃−ₓO₄—in the 3% Mg sample less than 1% of the total Mg was found in the MgₓCo₃−ₓO₄ phase, while in the 6% Mg sample it was 4% of the total Mg.
Table 2. Summary of XRF, XRD, TPR and \( \text{H}_2 \) chemisorption data for scaled up Co/Ru/Mg/\( \gamma \)-Al\(_2\)O\(_3\) catalysts prepared with ammonium acetate buffer washes after Mg addition. Amount of Mg incorporated into Mg\(_{x}\)Co\(_{3-x}\)O\(_4\) spinel was calculated by comparison with the literature values reported by Krezhov & Konstantinov [16]. This can then be used to estimate the mass of the Mg in the Mg\(_{x}\)Co\(_{3-x}\)O\(_4\) and, by subtraction from the total mass of Mg, the amount unaccounted for.

| Sample | nominal Mg loading (wt%) | calcination T after Mg addition (°C) | wash | XRF Mg (wt%) | XRF Al (wt%) | XRF Co (wt%) | XRF Co\(_3\)O\(_4\) lattice parameter (Å) | Co\(_3\)O\(_4\) lattice parameter x in Mg\(_x\)Co\(_{3-x}\)O\(_4\) | Mg in Mg\(_x\)Co\(_{3-x}\)O\(_4\) (%) | Co\(_3\)O\(_4\) (wt%) | TPR peak pos. (°C) | TPR peak pos. (°C) | H\(_2\) chemisorption CoSA (m\(^2\)/g Co) |
|--------|-------------------------|-------------------------------------|------|-------------|-------------|-------------|---------------------------------|---------------------------------|----------------|----------------|----------------|----------------|
| AW1    | 0                       | n.a.                                | n.a. | 0.00        | 36.57       | 17.08       | 8.0796 (3)                     | 0.00                            | 0               | 13.9           | 259            | 439            | 57.4           |
| AW2    | 3                       | 550                                 | n.a. | 2.64        | 34.89       | 17.37       | 8.0889 (4)                     | 0.21                            | 0.54            | 2.10           | 13.3           | 283            | 501            | 63.9           |
| AW3    | 3                       | 550                                 | buff | 1.94        | 35.08       | 17.08       | 8.0834 (3)                     | 0.09                            | 0.22            | 1.72           | 11.5           | 291            | 501            | 61.5           |
| AW4    | 3                       | 800                                 | n.a. | 2.64        | 35.11       | 17.21       | 8.0801 (3)                     | 0.01                            | 0.02            | 2.62           | 11.9           | 275            | 466            | 57.5           |
| AW5    | 6                       | 550                                 | buff | 2.56        | 34.84       | 16.95       | 8.0859 (4)                     | 0.15                            | 0.37            | 2.19           | 11.3           | 303            | 545            | 62.5           |
| AW6    | 6                       | 800                                 | n.a. | 5.18        | 33.09       | 17.25       | 8.0833 (3)                     | 0.09                            | 0.22            | 4.96           | 13.0           | 281            | 510            | 58.6           |
Figure 4. TPR data for Co/Ru/Mg/γ-Al₂O₃ catalysts prepared with acid washes after Mg addition. Traces are for samples (a) AW1, (b) AW2, (c) AW3, (d) AW4, (e) AW5 and (f) AW6. Black refers to catalyst prepared without acid wash; blue to buffer solution wash; and red to unwashed, calcined at 800°C after Mg addition.

The crystallite size of the MgₓCo₃₋ₓO₄ spinel phase can be calculated from XRD, and it can be seen that acid washing has a strong effect in reducing the particle size. The reason for this is unclear, and it occurs even in samples without Mg (electronic supplementary material, table S1). The relative effects of the Mg in MgₓCo₃₋ₓO₄ and Mg in MgₓAl₂O₃₊ₓ on particle size are difficult to deconvolute, because they are heavily autocorrelated (i.e. samples which have more Mg in the MgₓCo₃₋ₓO₄ phase also have more in the MgₓAl₂O₃₊ₓ phase), and the calcination at 800°C which would provide a comparator also has a separate effect on particle size. In general, the results agree with earlier results that higher Mg contents reduce particle size.

The effect of the different Mg levels on the reducibility, as seen in the TPR, is somewhat clearer (figure 4). The washing lowers the reduction temperature for the samples with the highest Mg loading of 6%, and this manifests itself in both a lowering of the peak for the Co(II)–Co(0) reduction and a sharpening of the reduction peak, consistent with the hypothesis that it is the amount of Mg in MgₓCo₃₋ₓO₄ rather than the total amount of Mg which determines the reduction temperature. Calcining the support at 800°C also lowers the reduction temperature relative to both washed and unwashed samples with the same initial Mg loading, and this also reinforces the hypothesis. This is complicated slightly by the 3% Mg samples calcined at 550°C, which show no change in peak position between washed and unwashed samples despite a large difference in the level of Mg in MgₓCo₃₋ₓO₄. However, taking these results together with the results of TGA in 5% H₂ carried out on the earlier samples (electronic supplementary material, figure S8), it becomes clear that it is the Mg in MgₓCo₃₋ₓO₄ which is detrimental to reducibility.

The cobalt surface area measurements as determined by H₂ chemisorption followed the same trend for the unwashed samples as the Mg-promoted samples scaled up previously but were slightly lower, which could be due to the higher calcination temperature. The Co surface area increased from 57.4 m² g⁻¹ for 0% Mg to 63.9 m² g⁻¹ for 3% Mg before decreasing to 59.2 m² g⁻¹ for 6% Mg. Again this is attributed to decreasing particle size coupled with decreasing reducibility as the Mg loading was increased.

AW2–AW6 were tested in the Fischer–Tropsch reaction to examine the effect of acid washing and high-temperature calcination prior to Co addition, along with a control sample, which was the 0% Mg sample discussed at the beginning of the Results section and in our previous work. To avoid confusion, we will refer to this sample as 0% Mg. The main difference between this sample and AW1 is that AW1 had a final heat treatment at 450°C rather than at 250°C, and a slightly lower Co surface area of 9.8 m² g⁻¹ compared with 11.2 m² g⁻¹. The testing procedure involved
examining the activity and selectivity at 210°C, 230°C and 240°C, before returning to 210°C over a total of 245 h time online at testing temperature.

Figure 5 shows the testing data for three samples—AW2, AW3 and AW4—with these three samples all containing 3% Mg, either without (AW2) or with (AW3) an acid wash, or with a calcination of 800°C after Mg impregnation (AW4). It can be clearly seen that AW4 has the highest activity at all temperatures, followed by AW3, with AW2 having the lowest activity. This clearly shows the beneficial effect of acid washing and higher temperature calcinations on the activity. Plots of testing data for all samples are shown in the electronic supplementary material, figures S9 (activity) and S10 (selectivity).

The trends in activity at 210°C (shown in table 3) cannot be explained by the Co surface area data measured by chemisorption (table 2). For example, aside from the 0% Mg standard, the highest Co surface area was measured for AW2, but this had the lowest activity. Despite the correlation between the Mg content in \( \text{Mg}_x\text{Co}_{3-x}\text{O}_4 \) and reducibility discussed above, it appears that the trends in activity cannot be explained simply by reducibility; for example, AW5 has a second reduction peak 35°C higher than any other catalyst but still has higher activity than AW2 or AW6.

However, calculating the site-time yields (activity per H chemisorption site, measured \textit{ex situ}) of each catalyst at 210°C and plotting them against the lattice parameter of the \( \text{Mg}_x\text{Co}_{3-x}\text{O}_4 \) phase measured in the as-calcined state provides insight into the trends in activity just discussed. Figure 6a shows the site-time yields with respect to the lattice parameter for all six catalysts. It is clear that the level of Mg substitution into the \( \text{Co}_3\text{O}_4 \) spinel has a marked effect on the intrinsic activity of the catalysts—as the \( \text{Mg}_x\text{Co}_{3-x}\text{O}_4 \) phase becomes more Mg-rich, the site-time yield decreases approximately linearly with increasing lattice parameter.

Because a single-mixed Co-Mg phase is formed after calcination, it is highly likely that, on reduction in the reactor, Mg which was mixed intimately with Co remains in close proximity to the reduced Co metal atoms. Hence the more Mg-rich the mixed spinel, the greater the amount of Mg in close proximity to the Co metal active sites in the activated catalyst. In order to investigate this, EELS was applied to a reduced and passivated 3% Mg catalyst (figure 7). The catalyst was reduced in pure \( \text{H}_2 \) at 425°C for 9 h followed by passivation in 1% \( \text{O}_2 \) in Ar at room temperature. The results indicate that Mg remains in close proximity to Co after reduction because the concentration distribution of Mg roughly maps that of Co. Presumably, on reduction, Co(III) and Co(II) ions are reduced to Co(0), destroying the \( \text{Mg}_x\text{Co}_{3-x}\text{O}_4 \) structure but remaining in close proximity to the Mg ions which they were previously intimately mixed with.
Table 3. Fischer–Tropsch activity data for Co/Ru/Mg/γ-Al₂O₃ catalysts at 210°C. Stabilities are calculated from the change in activity or stability between the initial period of testing at 210°C and the final period at 210°C after the middle high-temperature period. Errors shown are standard deviations. Site-time yield (STY) calculated using Co surface area from ex situ H₂ chemisorption. Chain growth probability, α, calculated from offline analysis of C₂⁺–C₄₀ fractions. GHSV, gas hourly space velocity.

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<th>sample</th>
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<th>final interval (h)</th>
<th>GHSV (L g⁻¹ h⁻¹)</th>
<th>initial syngas conversion (%)</th>
<th>initial selectivity</th>
<th>stability</th>
<th>CH₄ (%)</th>
<th>C₂–C₄ (%)</th>
<th>C₅⁺ (%)</th>
<th>conversion (%)</th>
<th>CH₄ (%)</th>
<th>C₅⁺ (%)</th>
<th>α (×10⁻³ s⁻¹)</th>
<th>STY</th>
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<td>87 ± 19</td>
<td>99 ± 4</td>
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Figure 6. (a,b) Relationship between site-time yield and Mg$_x$Co$_{3-x}$O$_4$ at (a) 210°C and (b) 240°C. (c) Relationship between percentage change in site-time yield comparing initial and final periods at 210°C. (d) Selectivities towards C$_5^+$ (squares) and CH$_4$ (diamonds) for all samples at all temperatures. The standard is black; filled symbols have 3% Mg; unfilled symbols have 6% Mg; blue symbols are calcined at 550°C and washed; red symbols are calcined at 550°C and unwashed; green symbols are calcined at 800°C.

Hence, it is proposed that Mg ions have a poisoning effect on the intrinsic activity of Co metal sites and the degree of poisoning depends on the concentration of Mg ions close to the Co sites. The decrease in activity with increasing Mg incorporation into Co$_3$O$_4$ appears not to be simply due to the total amount of Mg present in the sample, exemplified by the fact that AW6 has almost twice the total amount of Mg as AW2 but approximately 150% of the activity at 210°C.

Poisoning by Ca has been observed previously with the reduction in activity being attributed to electronic effects causing a decrease in surface hydrogen concentrations and increased CO adsorption and dissociation [22]. Another study investigating the poisoning effect of alkali and alkaline earth metals in the context of impurity elements in biomass-derived syngas implied that Mg had a poisoning effect but that it was less than for Ca. The authors suggested that a physical site-blocking effect could be the cause [23].

After approximately 65 h at high temperature, the catalysts were returned to 210°C. Interestingly, all the samples except the 0% Mg standard had higher activity after the high-temperature run than before, as shown in table 3 by stability values more than 100%. This suggests that, rather than deactivating the catalysts, the high-temperature period led to changes in the catalysts which actually activated them. The percentage increase in activity is strongly related
to the level of Mg in the $\text{Mg}_x\text{Co}_{3-x}\text{O}_4$ (figure 6c), suggesting that the higher temperatures are causing Mg to migrate away from Co, causing a reduction in the poisoning effect of Mg.

During the high-temperature phase of the test at 230°C and then 240°C, the site-time yields of all the catalysts except AW5 increase relative to the 0% Mg standard sample, to the extent that, by 240°C, AW4 has a higher site-time yield than the standard, while AW3 is the same within error (figure 6b). Again, the relative increase in site-time yield is somewhat correlated to the level of Mg in the $\text{Mg}_x\text{Co}_{3-x}\text{O}_4$, although as the yields fall back below those of the standard when returning to 210°C, this effect cannot be entirely due to a high-temperature activation process and must be at least somewhat an intrinsic property of Mg promotion. Arrhenius plots for the 0% Mg standard, AW3 and AW4 (electronic supplementary material, figure S11) show no deviation from a linear trend at the highest temperature, which also indicates that there is no large change in the number of active sites. The fact that it is AW4, in which $x$ in $\text{Mg}_x\text{Co}_{3-x}\text{O}_4$ is only 0.01, which shows the largest increase in site-time yield with temperature raises the interesting prospect that the improved high-temperature performance is due to Mg incorporated into the alumina phase.

Looking at the selectivity of the samples (figure 6d), there is a consistent correlation between selectivity for all products and conversion among all samples at all temperatures, strongly suggesting that Mg in any form is not having a detectable influence on the selectivity or on the relative rates of possible competing reactions such as water gas shift.

4. Conclusion

The fact that magnesium is incorporated into the cobalt oxide spinel phase as well as the alumina phase despite the high-temperature treatment before impregnation has important implications for the use of structural promoters. We have shown that, while incorporation into the alumina phase may have beneficial effects on the high-temperature performance, incorporation into the cobalt phase has severely deleterious effects on the site-time yield, which may hold for other structural promoters. We have also shown that the impregnation solution has leaching effects greater than its pH implies, and also that the solution is potentially capable of dissolving measurable amounts of aluminium from the support, which presumably is also incorporated into the cobalt spinel phase, with unknown effects.
Quantification of the extent and effect of this doping of promoters and elements from the support into different structures of a catalyst is a difficult problem requiring detailed investigation with XRD, high-resolution TEM and elemental analysis, as well as a carefully designed experimental strategy, but should be a fruitful avenue of research in assessing new materials and reassessing older materials.

Data accessibility. The datasets supporting this article are available (doi:10.17638/datacat.liverpool.ac.uk/69).

Authors’ contributions. J.R.G. and P.B.: samples synthesis, data acquisition and analysis; D.O., D.I.E. and P.R.E.: data acquisition and analysis; all authors were involved in the conception and design of the experiments, drafting and revising the manuscript, and final approval.

Competing interests. We have no competing interests.

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