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# Enhancement of iron-based oxygen carriers through alloying with tungsten oxide for chemical looping applications including water splitting

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**Abstract:** Chemical looping applications offer a variety of options to decarbonise different industrial sectors, such as iron and steel and hydrogen production. Chemical looping with water splitting (CLWS) is a chemical looping technology, which produces  $H_2$  while simultaneously capturing  $CO_2$ . The selection of oxygen carriers (OCs) available to be used in CLWS is finite, due to the thermodynamic limitations of the oxidation with steam for different materials at the relevant process temperatures. Iron-based materials are one of the most widely studied options for chemical looping combustion (CLC), touted for their relative abundance and low cost; likewise, for CLWS, iron is the most promising option. However, when the reduction of iron oxide ( $Fe_2O_3$ ) is extended to wüstite (FeO) and iron (Fe), agglomeration and sintering problems are the main challenge for fluidisation. This work presents iron and tungsten mixed oxides as the OCs for a family of chemical looping applications. The OCs were produced via co-precipitation; performance assessment was conducted in a thermogravimetric analyser and a lab-scale fluidised bed reactor over continuous redox cycles. The use of tungsten combined with iron results in a solid solution of tungsten within the  $Fe_2O_3$  matrix that produced a more mechanically stable material during operation, which performed well during multiple redox cycles with no apparent decrease in the oxygen transport capacity and showed no apparent agglomeration. Furthermore, materials containing tungsten showed a resistance to carbon deposition, whereas the reference  $Fe_2O_3$  showed peaks of CO and  $CO_2$  during the oxidation period, thus indicating carbon deposition. © 2023 Society of Chemical Industry and John Wiley & Sons, Ltd.

**Keywords:**  $CO_2$  capture; chemical looping combustion; iron oxide; mixed oxides

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## Introduction

Global carbon dioxide emissions from fossil fuels combustion are the main contribution causing global warming. To achieve the global reduction targets in the near term, carbon capture and storage (CCS) technologies need to be placed to capture CO<sub>2</sub> from industry, which is the largest emitting sector of this greenhouse gas, with over 40% of the global emissions in 2019 (when heat and electricity emissions are allocated to final sectors).<sup>1</sup>

Chemical looping combustion (CLC) is part of the CCS technologies portfolio, in which redox reactions of metal oxides are used to transfer oxygen from the air to the fuel for combustion processes. Since direct contact between the fuel and air is avoided, a concentrated CO<sub>2</sub> stream without combustion impurities is produced, which then can be separated (through steam condensation), transported and stored.<sup>2</sup> The choice of metal oxide to be used as oxygen carrier (OC) depends on the CLC process, but mainly these materials need to maintain a high reactivity and a constant oxygen transport capacity (Ro) and remain both mechanically and chemically stable over extended periods of time.

Hydrogen production is achieved mainly via steam methane reforming (SMR), and accounts for around 3% of the global CO<sub>2</sub> emissions from the industry sector.<sup>3</sup> Chemical looping with water splitting (CLWS) is a CLC variant used for H<sub>2</sub> production, which is a promising option to reduce emissions while producing high purity H<sub>2</sub>.<sup>4</sup> The most common configuration for CLWS is a three-reactor system, where a third reactor (steam reactor) is added to the standard CLC configuration (fuel and air reactors) (Fig. 1).<sup>5</sup>

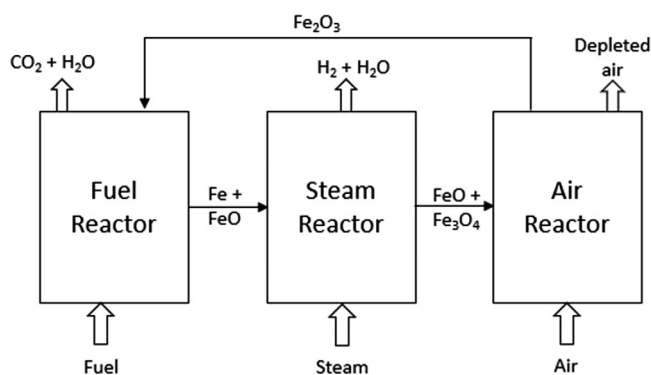


Figure 1. Chemical looping with water splitting process, where the oxide system of Fe<sub>2</sub>O<sub>3</sub>/Fe<sub>3</sub>O<sub>4</sub>/FeO/Fe is used as oxygen carrier.

Thermodynamic limitations of the redox pairs that can be used for CLWS limit the options to just a few choices: Fe, W and Ce oxides.<sup>6</sup>

Fe-based oxygen carriers for CLC have been widely studied and represent an advantage when compared to the other options for CLWS due to its abundance, low cost and being environmental benign.<sup>7,8</sup> For most CLC applications, Fe-based oxygen carriers have been used in conditions to restrict the reduction of Fe<sub>2</sub>O<sub>3</sub> to Fe<sub>3</sub>O<sub>4</sub>. Further reduction to FeO and Fe represent slower reaction kinetics and lower fuel combustion efficiencies when compared to reduction to only Fe<sub>3</sub>O<sub>4</sub>.<sup>9,10</sup> These limitations are due to the severe chemical and mechanical stresses to which the material is subjected, leading to an increased reactive/thermal sintering, which reduces the available reactive surface area.

In this work we present a new iron-tungsten mixed oxide for extended reduction in CLC as a first step towards its use in CLWS to produce hydrogen. We compare the performance of this new material with two different Fe<sub>2</sub>O<sub>3</sub> sample materials that serve as a reference to quantify the improvements.

## Experimental

### Material synthesis

#### Synthesis of oxygen carriers via co-precipitation

The oxygen carriers were synthesised using an automated rig for co-precipitation, detailed elsewhere.<sup>11</sup> For the iron-tungsten (Fe-W) mixed oxide (to avoid confusion, the material was synthesized as a mixed oxide, which we believe then partially alloyed at reaction temperatures), two solutions were prepared for the synthesis: (1) Fe(NO<sub>3</sub>)<sub>3</sub>·9H<sub>2</sub>O and Na<sub>2</sub>WO<sub>4</sub>·2H<sub>2</sub>O were dissolved in deionised water (DI water) to obtain a 250 ml solution with a total ion concentration of 1 M (with a molar Fe-W ratio of 0.95:0.05 and 0.98:0.02) and (2) a 1 M solution of NaOH to regulate the pH.

The rig controlled the flow rates of the two solutions to maintain a constant pH value of 12 ± 0.1 during the co-precipitation under constant stirring. After addition of solution 1 was completed, the stirring was stopped, and the mixture was aged for 1 h. The precipitate was then washed, by adding DI water, followed by centrifugation at 3000 rpm for 5 min to separate out the phases. The washing process was repeated until the measured conductivity was  $\sigma < 100 \mu\text{S}$ . The precipitate was then dried overnight at 60°C and calcined for 6 h

**Table 1. Summary of oxygen carriers prepared for this study.**

Sample name	Tungsten content (mol%)	Preparation method
FeW-5-CP	5	Co-precipitation
FeW-2-CP	2	Co-precipitation
Fe <sub>2</sub> O <sub>3</sub> -MM	-	Mechanically mixing
Fe <sub>2</sub> O <sub>3</sub> -CP	-	Co-precipitation

at 900°C in air (2 L•min<sup>-1</sup>) using a horizontal tube furnace with a heating rate of 15°C•min<sup>-1</sup>.

Two reference samples of Fe<sub>2</sub>O<sub>3</sub> were prepared for comparison. The first sample was prepared following the same co-precipitation procedure but without the addition of Na<sub>2</sub>WO<sub>4</sub>•2H<sub>2</sub>O.

### Synthesis of oxygen carriers via mechanical mixing

A second reference sample was prepared from a commercially available material. A batch of mechanically mixed (MM) oxygen carriers was prepared by mixing 0.2 mol of Fe<sub>2</sub>O<sub>3</sub> (powder <5 μm, > 96% from Sigma-Aldrich) with 30 ml of DI water. The slurry was loaded into a planetary ball mill (PM-400, RETSCH) and was milled together with steel balls for 6 h at 250 rpm. The sample was dried and calcined under the same conditions as the co-precipitated materials.

A summary and nomenclature of the samples prepared for this study is presented in Table 1.

### Oxygen carrier characterization and testing

Powder X-ray diffraction (XRD) analysis was carried out in a D2 Phaser (Bruker) using Cu-Kα radiation with λ = 1.541 Å. Scanning electron microscopy (SEM) in a JEOL JSM 6010LA at 20 kV was used to observe the surface morphology of the fresh and reacted particles. The crushing strength of the oxygen carriers was measured using a 50 N force gauge (Mark-10). Nitrogen adsorption analysis was done using a TriStar 3000 (Micromeritics) sorption analyser with N<sub>2</sub> at 77K, to determine the surface areas (BET), pore size distributions and volume of the micro and mesopores (size range up to ~200 nm - BJH). Mercury intrusion porosimetry (MIP) was carried out using an AutoPore IV 9500 (Micromeritics) porosimeter to

determine the pore volume in the region outside of the resolution of N<sub>2</sub> adsorption, between 10 nm and 10 μm. Helium pycnometry was carried out using an AccuPyc II 1340 (Micromeritics) pycnometer to determine the skeletal density of the oxygen carriers.

A thermo-gravimetric analyser (TGA) model Q5000 (TA Instruments) was used to analyse the materials. Isothermal redox cycling was performed to evaluate the conversion and redox capability of the oxygen carriers (particle size range of 150–300 μm). A platinum crucible (*h* = 1.5 mm, *d* = 9.8 mm) was used to hold a layer of OC (5 ± 0.3 mg) in each experimental run. The total gas flow rate used in all experiments was kept constant at a value of 200 ml•min<sup>-1</sup> (SATP). A purge stage of N<sub>2</sub> was used to separate each reduction and oxidation stage and thus avoid gas mixing in the reaction chamber. The observed rate of reaction (in mol•g<sup>-1</sup>•s<sup>-1</sup>), was calculated from the weight loss (for the reduction) and weight gain (for the oxidation) periods of the experiment. The weight profiles obtained from the TGA were differentiated with respect to the time and divided through the initial mass of the fully oxidised sample. Redox-cycling experiments with a higher sample mass of OCs in the TGA (40 mg) was carried out to measure the crushing strength and perform XRD-analysis of the cycled particles. The experiments with a higher sample mass were performed with increased cycle times, for both the reduction and oxidation periods, to match the experimentally obtained Ro of the experiments with the low sample mass.

Tests were complemented with redox cycling experiments performed in a lab-scale fluidised bed reactor (FBR).<sup>12</sup> An inert experiment was carried out initially with 40 g of quartz sand (425–500 μm). Then, 0.5 g of OCs in the particle size range of 300–425 μm was added to the bed and the cycling experiment was run with the bed containing the sample. The change of gas concentration was measured continuously during the experiment and the data was deconvoluted to account for gas mixing and response time of the gas analyser following a method described elsewhere.<sup>13</sup>

## Results and discussion

### Characterisation results

Tungsten addition into the materials increased the porosity measured using both techniques. The BET surface area increased for the FeW-5-CP and FeW-2-CP samples to values of 3.77 and 3.39 m<sup>2</sup>•g<sup>-1</sup>

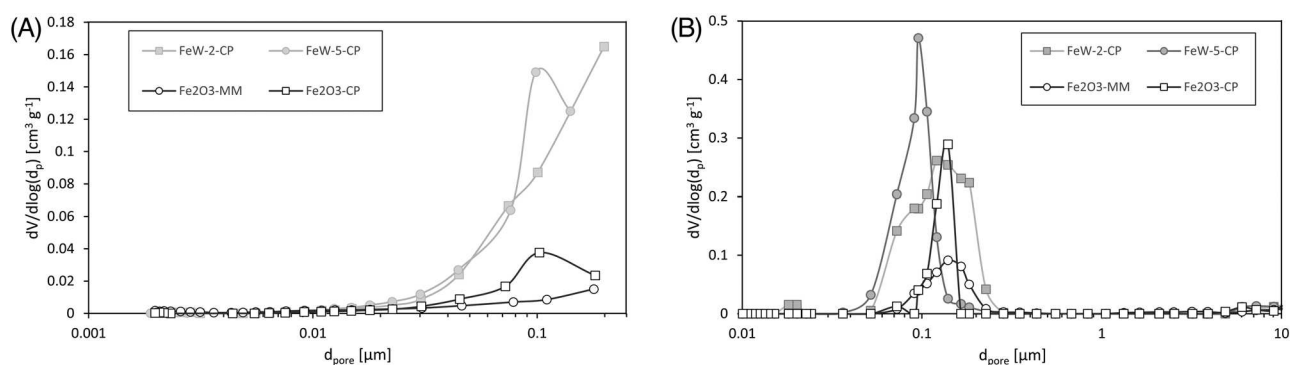


Figure 2. Pore size distributions determined by (A) BJH  $N_2$ -adsorption analysis, and (B) MIP analysis on the entire pore size range of the fresh oxygen carrier particles for all the samples prepared in this study.

**Table 2. Summary of MIP,  $N_2$  adsorption and He pycnometry measurements to determine the pore size distributions and porosity for the different OCs prepared for this work.**

	FeW-5-CP	FeW-2-CP	Fe <sub>2</sub> O <sub>3</sub> -MM	Fe <sub>2</sub> O <sub>3</sub> -CP
BET surface area [ $m^2 \cdot g^{-1}$ ]	3.77	3.39	2.48	1.94
Cumulative pore vol. (BJH) [ $cm^3 \cdot g^{-1}$ ]	0.065	0.045	0.009	0.022
Cumulative pore vol. (MIP) [ $cm^3 \cdot g^{-1}$ ]	0.089	0.092	0.030	0.042
Average $d_p$ (4 V/A) (MIP) [nm]	466	420	1,200	760
Skeletal density (He pyc.) [ $g \cdot cm^{-3}$ ]	5.07	5.24	5.05	5.00
Porosity (MIP)	0.628	0.634	0.485	0.458

respectively, when compared to the reference samples, with values of 2.48 and 1.94  $m^2 \cdot g^{-1}$  for the Fe<sub>2</sub>O<sub>3</sub>-MM and Fe<sub>2</sub>O<sub>3</sub>-CP samples, respectively. The total volume of pores measured for the reference samples was smaller than that for the Fe-W mixed oxide samples. In addition, the measured skeletal density by He pycnometry showed a slight increase with the addition of W in the analysed samples (size range 300–425  $\mu m$ ).

The total porosity of the materials measured by MIP showed an increase by the W addition into the material to values of around 60%, when compared to the Fe<sub>2</sub>O<sub>3</sub> reference samples (below 50%). The pore size distributions of the OCs were found to shift towards smaller pore diameters (Fig. 2B).

Pore size distributions determined by  $N_2$  adsorption and MIP are presented in (Fig. 2A and B), respectively. The results of the porosity measurements are summarised in Table 2.

### Thermogravimetric screening and testing of the redox capability of the OCs

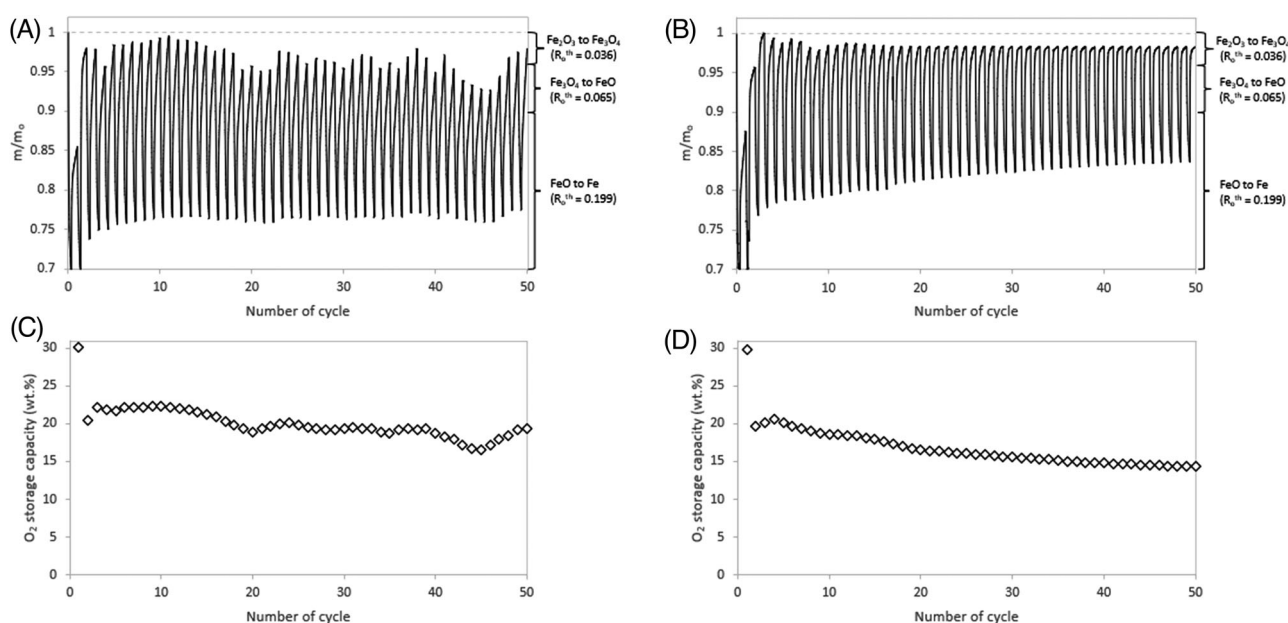
The redox capability of the oxygen carrier samples was tested in the TGA. A comparison between the two Fe<sub>2</sub>O<sub>3</sub> reference samples and the co-precipitated Fe-W

oxygen carriers was carried out over 50 redox-cycles at 850°C under a reducing environment of 15 vol% CO +  $N_2$  for 10 min and oxidised in 4 vol% O<sub>2</sub> +  $N_2$  for 20 min. These preliminary experiments were conducted with CO to avoid the complexity of operating a TGA with steam addition. The O<sub>2</sub> concentration on the oxidation step was chosen to be low to avoid any localised hot spots due to temperature gradients in the material since the oxidation of Fe-based materials is highly exothermic.

### Reference Fe<sub>2</sub>O<sub>3</sub> samples

The reactivity of the Fe<sub>2</sub>O<sub>3</sub>-MM and Fe<sub>2</sub>O<sub>3</sub>-CP samples was screened in the TGA to serve as reference and be able to compare the improvements made by adding W to the materials. The results of these experiments are presented in Fig. 3.

An analysis of the relative weight change and the Ro of both samples, (Fig. 3A and C) for the Fe<sub>2</sub>O<sub>3</sub>-MM and (Fig. 3B and D) for Fe<sub>2</sub>O<sub>3</sub> co-precipitated, shows that all the lattice oxygen available for reaction was used on the first reduction cycle for both materials. Despite this, the oxidation does not recover back to a



**Figure 3.** Results of the TGA experiments using the  $\text{Fe}_2\text{O}_3$  reference samples as oxygen carrier over fifty redox cycles at  $850^\circ\text{C}$ : (A) The relative weight change,  $m/m_o$ , recorded during the redox cycling experiment, and (C) Ro for the  $\text{Fe}_2\text{O}_3$ -MM and (B) The relative weight change,  $m/m_o$ , recorded during the redox-cycling experiment, and (D) Ro for the  $\text{Fe}_2\text{O}_3$ -CP. The values for  $\text{Ro}^{\text{th}}$  represent the theoretical weight change for each phase transition change of iron oxide: transition from  $\text{Fe}_2\text{O}_3$  to  $\text{Fe}_3\text{O}_4$  has an  $\text{Ro}^{\text{th}} = 0.036$ , the next transition from  $\text{Fe}_3\text{O}_4$  to  $\text{FeO}$  has an  $\text{Ro}^{\text{th}} = 0.065$  and the final transition from  $\text{FeO}$  to  $\text{Fe}$  has an  $\text{Ro}^{\text{th}} = 0.199$ .

value of  $m/m_o = 1$  in any of the samples in the first oxidation cycle (which would have meant full oxidation back to  $\text{Fe}_2\text{O}_3$ ), and this was only achieved in cycle 3 for the  $\text{Fe}_2\text{O}_3$ -CP sample. The  $\text{Fe}_2\text{O}_3$ -MM sample did not reach full oxidation to  $\text{Fe}_2\text{O}_3$  during the whole experiment. The relative weight change profile of the  $\text{Fe}_2\text{O}_3$ -MM samples shows an erratic behaviour, with oxidation after each cycle stopping at different  $m/m_o$  values and not reaching oxidation back to  $\text{Fe}_2\text{O}_3$  in at least 15 cycles, which means that the oxidation only reached  $\text{Fe}_3\text{O}_4$  in 30% of the experimental runs. The experimental value of the Ro measured for the  $\text{Fe}_2\text{O}_3$ -MM sample followed this same erratic trend and showed an overall decrease with cycling (Fig. 3C).

The value of Ro measured for the  $\text{Fe}_2\text{O}_3$ -CP sample decreased steadily after the 3<sup>rd</sup> cycle with continuous cycling, as can be seen in (Fig. 3D), reaching a final value of 14.5 wt.%. The oxidation step in the experiment appears to stabilise after 15 cycles and stopping at a value of  $m/m_o = 0.98$  continuously until the end of the run. These results for both reference samples might suggest that sintering of the pores worsens with continuous cycling. Therefore, the surface morphology of the particles was studied.

Results of the SEM analysis for the reference samples is shown in Fig. 4.

SEM images for both reference fresh samples show a dense particle with small grains in the surface of the particles of the  $\text{Fe}_2\text{O}_3$ -MM (Fig. 4A and B) and the  $\text{Fe}_2\text{O}_3$ -CP (Fig. 4C and D). The recovered oxygen carrier samples after redox-cycling exhibited sintering of the particles after the experiment, for both the  $\text{Fe}_2\text{O}_3$ -MM sample (Fig. 4E and F) and the  $\text{Fe}_2\text{O}_3$ -CP sample (Fig. 4G and H). A clear formation of a grain structure, in which the grains started to fuse within each other can be seen in (Fig. 4E and F). Less sintering within the grains can be observed for the  $\text{Fe}_2\text{O}_3$ -CP sample (Fig. 4G and H) compared to the  $\text{Fe}_2\text{O}_3$ -MM sample, but a fully developed grain structure within the surface of the  $\text{Fe}_2\text{O}_3$ -CP sample is observed. This sintering will result in an increase in the diffusion resistance through the surface of the particles, which results in the consistent decrease on the experimental Ro observed in (Fig. 3A and B).

## Fe-W mixed oxides

The results of the experiments for the FeW-2-CP and FeW-5-CP samples are presented in Fig. 5 and 6,

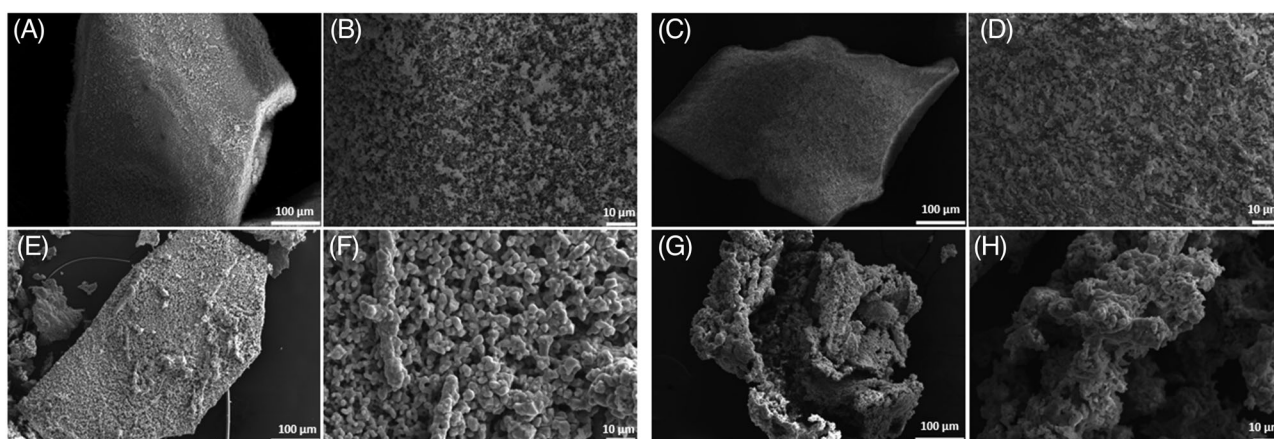


Figure 4. Scanning electron micrographs of fresh and cycled  $\text{Fe}_2\text{O}_3$  reference samples. The micrographs show (A, B) the fresh  $\text{Fe}_2\text{O}_3$ -MM sample, (C, D) the fresh  $\text{Fe}_2\text{O}_3$ -CP sample, (E, F) the  $\text{Fe}_2\text{O}_3$ -MM sample after 50 cycles, and (G, H) the  $\text{Fe}_2\text{O}_3$ -CP sample after 50 cycles, SE micrographs with (A, C, E, G)  $\times 200$ , and (B, D, F, H)  $\times 1200$  magnification.

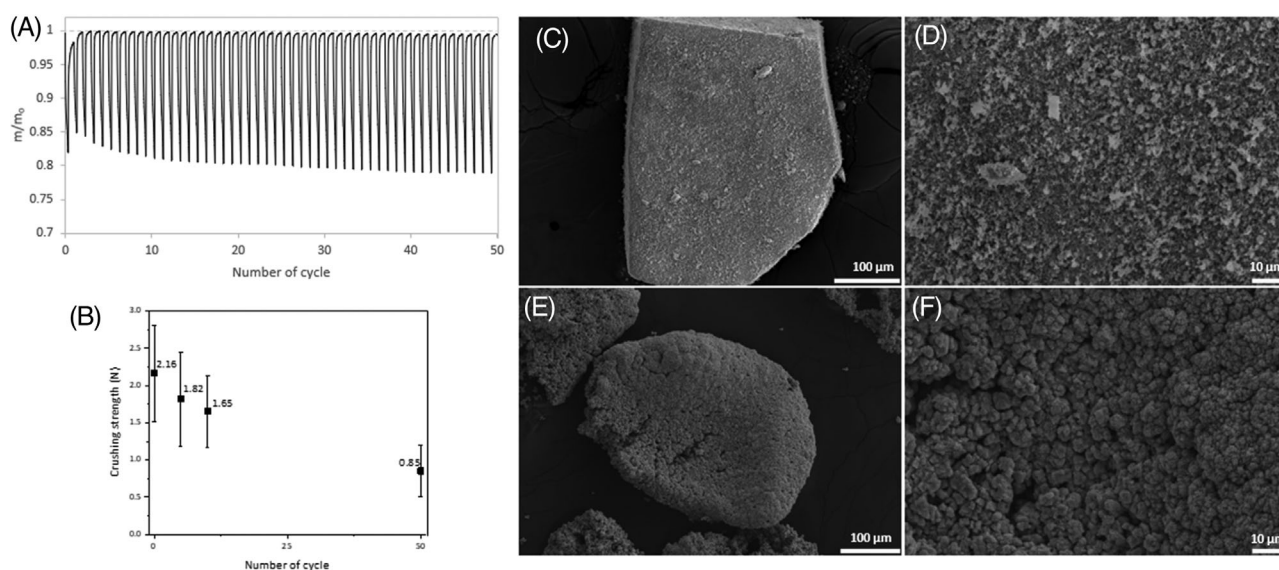


Figure 5. Results of the TGA experiments using the FeW-2-CP sample as oxygen carrier over 50 redox cycles at  $850^\circ\text{C}$ . (A) The relative weight change,  $m/m_0$ , recorded during the redox cycling experiment, (B) the crushing strength against cycle number, SEM images of (C, D) the fresh sample, and (E, F) the sample after 50 cycles, SE micrographs with (C, E)  $\times 200$ , and (D, F)  $\times 1200$  magnification.

respectively. The relative weight change of the FeW-2-CP sample (Fig. 5A) showed a stable redox-cycling capability and a measured Ro of 19 wt.% during the last 20 cycles of the experiment. (Fig. 6A) shows an analysis on the relative weight change of the FeW-5-CP sample. The measured Ro increased during the experiment, with a maximum value of 17 wt.% at the end of the experiment. Both samples prepared using tungsten showed full reoxidation back to a value of  $m/m_0 = 1$  during continuous cycling, representing a

significant improvement from the reference  $\text{Fe}_2\text{O}_3$  samples.

The crushing strength of both samples decreased after the experimental run, most likely due to an increased porosity developed after cycling. The crushing strength of the FeW-2-CP (Fig. 5B) and the FeW-5-CP (Fig. 6B) decreased to 40% and 52% of their initial values, respectively. SE micrographs of the particles recovered after cycling showed the formation of a grain structure in the surface of the FeW-2-CP

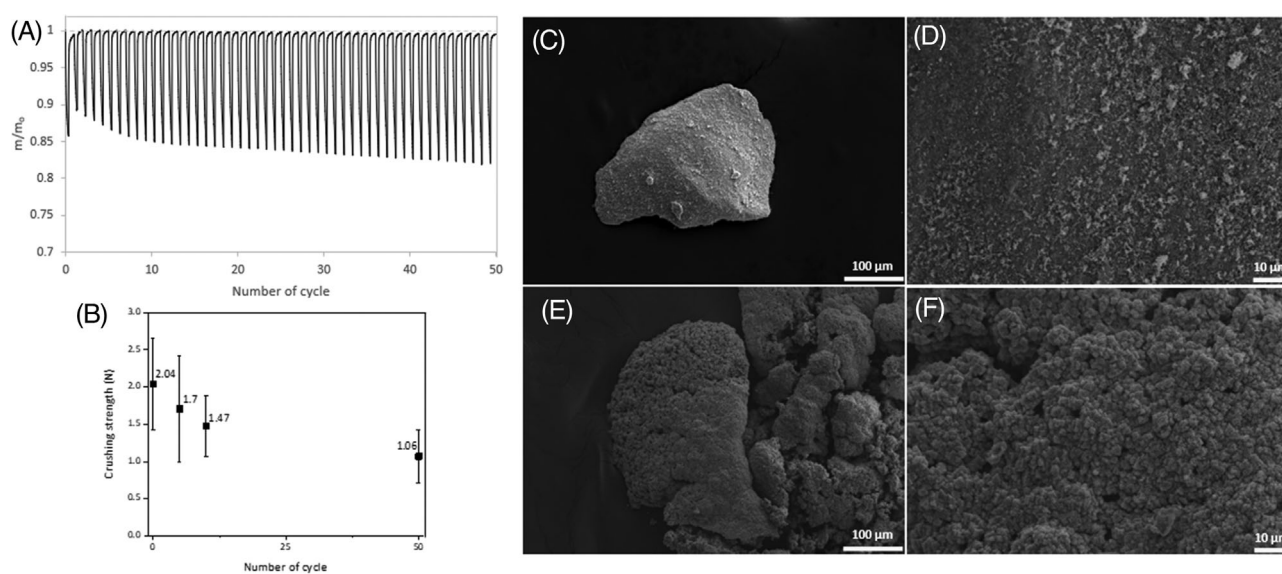


Figure 6. Results of the TGA experiments using the FeW-5-CP sample as oxygen carrier over 50 redox cycles at 850°C. (A) The relative weight change,  $m/m_0$ , recorded during the redox cycling experiment, (B) the crushing strength against cycle number, SEM images of (C, D) the fresh sample, and (E, F) the sample after 50 cycles, SE micrographs with (C, E)  $\times 200$ , and (D, F)  $\times 1200$  magnification.

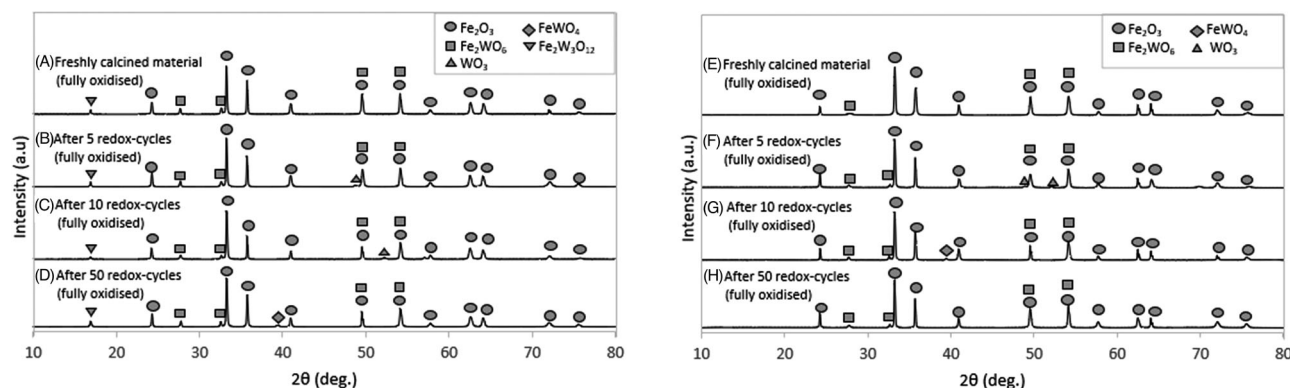


Figure 7. XRD plot of the FeW-5-CP sample in its fully oxidised state (A) before cycling, (B) after 5 redox cycles, (C) after 10 redox cycles, and (D) after 50 redox cycles, as well as the XRD plot of the FeW-2-CP sample in its fully oxidised state (E) before cycling, (F) after 5 redox cycles, (G) after 10 redox cycles, and (H) after 50 redox cycles.

(Fig. 5E and F) and the FeW-5-CP (Fig. 6E and F). The recovered particles showed no sign of agglomeration and some mild intraparticle sintering at the surface can be observed in Figs. (5F) and (6F) for both samples.

The X-ray diffraction analysis of the Fe-W samples (Fig. 7) was performed to study the change of the oxidised phases during cycling. XRD analysis confirmed that the major phase identified in all samples in the study is  $\text{Fe}_2\text{O}_3$  (JCPDS 04-008-7627). The analysis showed the formation of different iron-tungstates structures:  $\text{Fe}_2\text{WO}_6$  (JCPDS 00-042-0492),  $\text{FeWO}_4$  (JCPDS 01-071-2390) and  $\text{Fe}_2\text{W}_3\text{O}_{12}$  (JCPDS 00-024-0538), as well as  $\text{WO}_3$

(JCPDS 04-007-2426). These previous results are in agreement with previous findings from Maiti et al.,<sup>14</sup> which found that the reduction and oxidation of the iron tungstate ( $\text{Fe}_2\text{WO}_6$ ) goes through a series of intermediary phases. In the XRD patterns of the FeW-2-CP samples (Fig. 7E–H), the formation of  $\text{Fe}_2\text{W}_3\text{O}_{12}$  was not identified, whereas it was formed in all the FeW-5-CP samples analysed (Fig. 7A–D). This was most likely because of the increased W available to form the intermediary phases. An analysis of the phase evolution of the W present in the FeW-2-CP during cycling showed an evolution to form the phase  $\text{Fe}_2\text{WO}_6$ , which appears to be the more stable form of

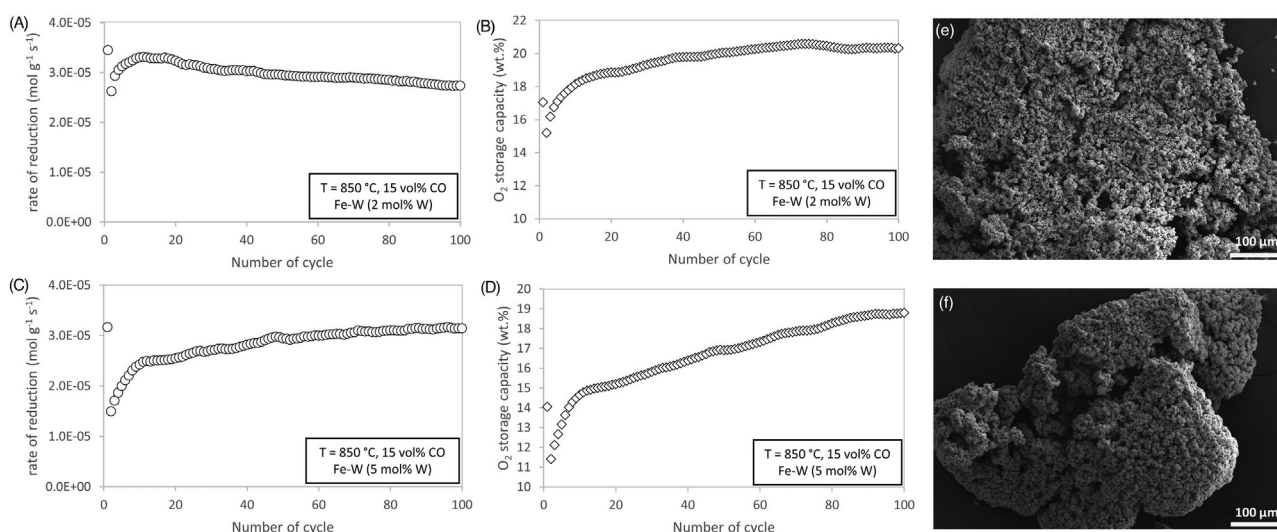


Figure 8. The maximum observed rate of reduction and Ro measured in the TGA at 850 C over 100 redox cycles using 15 vol% CO + N<sub>2</sub> for the reduction and 4 vol% O<sub>2</sub> for the oxidation: (A) observed rate and (B) Ro for the FeW-2-CP oxygen carrier and (C) observed rate and (D) Ro for the FeW-5-CP oxygen carrier, and SEM images of an oxygen carrier particle recovered after 100 redox cycles of the (E) FeW-2-CP and (F) FeW-5-CP, SE micrographs with (E, F) ×200 magnification.

the W present in the material. This phase became less apparent as the number of redox-cycles increase, most likely because the tungsten had formed a solid solution within the Fe<sub>2</sub>O<sub>3</sub> matrix in the material.

### Long-term redox cycling performance of the Fe-W OCs with CO

Long-term TGA experiments with CO were carried out to assess the CLC performance of the Fe-W oxygen carriers over 100 cycles. The samples were reduced in 15 vol% CO + N<sub>2</sub> for 10 min and oxidised in 4 vol% O<sub>2</sub> + N<sub>2</sub> for 20 min. The maximum observed rates of CO consumption and oxygen storage capacity of the materials are presented in Fig. 8.

(Figure 8A) shows that the maximum rate observed for the FeW-2-CP decreased from its initial value of  $3.45 \times 10^{-5} \text{ mol g}^{-1} \text{ s}^{-1}$  and quickly recovered over within the first 10 cycles, reaching a reactivity of almost the same value it had initially. The observed rate of reduction started to decrease steadily after cycle 20, from  $3.25 \times 10^{-5} \text{ mol g}^{-1} \text{ s}^{-1}$  to  $2.75 \times 10^{-5} \text{ mol g}^{-1} \text{ s}^{-1}$  at the end of the 100 redox-cycles. (Fig. 8C) shows that the FeW-2-CP particle started to disintegrate after 100 redox-cycles. The micrograph also shows that a highly porous structure had formed. It is possible that this was enhanced by bulk diffusion of metal ions outwards from the particle centre, as has been observed before.<sup>15</sup>

The maximum observed rate for the FeW-5-CP decreased from its initial value of  $3.15 \times 10^{-5} \text{ mol g}^{-1} \text{ s}^{-1}$  and steadily recovered over the duration of the experiment, reaching again the initial reactivity in the last ten cycles (Fig. 8C). Similarly, (Fig. 8D) shows that the Ro for this material had a steady increase until reaching stability at the last ten cycles with an average value of 18.7 wt.%. The decrease and following recovery of the maximum rate of reduction was most likely to structural rearrangements, which then stabilize after around 100 cycles. (Fig. 8F) shows a FeW-5-CP particle recovered after 100 redox-cycles, which appears to have developed a highly porous structure while maintaining most of its integrity and shape. Given the additional cost of adding W to the system, and the apparent maximum in reactivity and Ro, it does not appear likely that adding additional W beyond 2 mol% is worthwhile.

Although an experiment testing 100 redox cycles may provide an indication of the long-term performance of the OCs, a much higher number of cycles would be required to estimate indicators such as lifetime of the particles.

### Carbon deposition

Thermodynamically it is possible that carbon was deposited on the iron oxide particles when reduction to

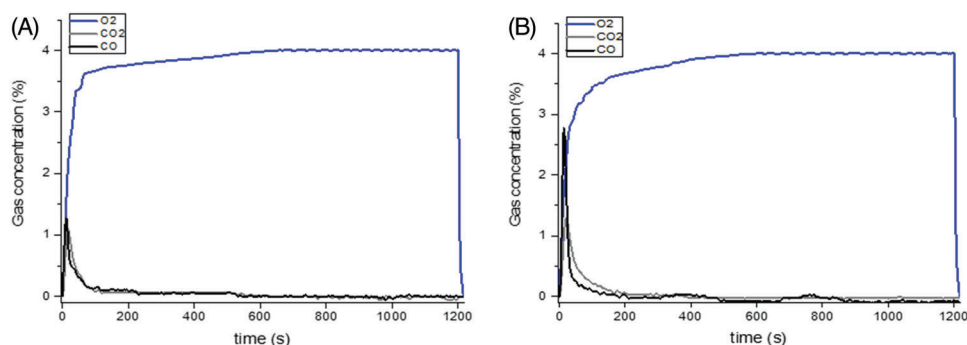


Figure 9. Concentration profile of the 5<sup>th</sup> oxidation cycle in the FBR cycling experiment at 850°C using 15 vol% CO + N<sub>2</sub> for the reduction and 4 vol% O<sub>2</sub> + N<sub>2</sub> for the oxidation: (A) Fe<sub>2</sub>O<sub>3</sub>-MM sample and (B) Fe<sub>2</sub>O<sub>3</sub>-CP sample. The gas flow rate was kept constant to maintain a value of  $u/u_{mf}$  of 3.

Fe was done with CO.<sup>16</sup> Since any residual carbon deposited on the surface of the particles would react with the oxygen to produce CO<sub>2</sub> and CO during the oxidation step, the extent of the carbon deposition was quantified from the deconvoluted CO<sub>2</sub> and CO gas concentration profiles measured during the oxidation step in the fluidised bed experiments (Fig. 9). CO<sub>2</sub> and CO produced was assumed to be formed via reactions 1 and 2, respectively.



Carbon deposition on the surface of the reference samples accounted for 2.25 wt.% and 5.45 wt.% of the initial sample mass of OC for the Fe<sub>2</sub>O<sub>3</sub>-MM and Fe<sub>2</sub>O<sub>3</sub>-CP, respectively. No carbon deposition was observed in the experiments with the Fe-W materials under the same conditions, a significant improvement most likely due to the high degree of metal dispersion and the resulted higher surface area of the Fe-W mixed oxides that may have improved the thermal and chemical stability of the material, which has been observed before.<sup>17–19</sup> Of course, increasing the amount of CO<sub>2</sub> in the feed can also reduce the amount of carbon deposited.

## Conclusions

This work demonstrated that the synthesis of Fe-W mixed oxides via co-precipitation to be used as oxygen carrier in CLWS as a promising option. The alloying of Fe with small amounts of W showed an improvement of the reactivity over continuous CLC operation. We

have compared the performance of the mixed oxides with two reference sample materials (Fe<sub>2</sub>O<sub>3</sub>) and the Fe-W oxides showed no sign of agglomeration and a resistance to carbon deposition, whereas both reference materials showed carbon deposition and a decrease in the measured Ro, and redox capability that can be attributed to the observed sintering of the particles recovered after cycling.

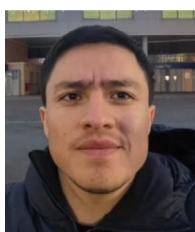
The stable reactivity and performance of the new Fe-W OCs can be attributed to the formation of a solid solution between the iron oxide and the W in the material, which increases the melting point of the reduced phases of the material, therefore avoiding sintering and agglomeration problems. The results of the XRD analysis suggest that the W present in the material goes through different intermediary phases during redox-cycling. Further analysis should focus on the phase evolution of the Fe-W mixed oxide to determine the influence of the transition phases during the first few redox cycles and the later stabilisation of the OC performance. Further studies with lower W content could further explain how the stabilisation of the Ro and reaction rate is influenced by the intermediary iron-tungstate phases.

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