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Thermochemical energy storage performance of copper oxides: Effect of support materials

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SUMMARY

Thermochemical energy storage (TCES) is one of the most promising methods for utilization of solar energy. Metal oxides can exhibit reversible redox reactions that are useful for TCES applications. Especially, transitional metal oxides can undergo reduction reactions at high temperatures while absorbing energy given to the system. Later on, when the temperature goes down below a phase-transition temperature, exothermic re-oxidation reactions can take place. Air can be used both as oxygen source and heat transfer medium during the redox reactions. Recently, several studies have been published about the utilization of metal oxides for TCES applications. Among these metal oxides, copper oxides received a great attention owing to its cyclic stability and suitable redox temperature. In this study, copper oxides are used as energy storage material in combination with ZrO₂, ZrO₂-La₂O₃, MgAl₂O₄, Mg₂Al₂O₄-La₂O₃, CeO₂, CeO₂-La₂O₃ as support materials. The best results were obtained from samples supported with MgAl₂O₄, Mg₂Al₂O₄-La₂O₃. This most likely eventuated due to the other reversible phase transformations that take place in these systems such as formation of LaAlO₃ and Cu₂Al₂O₄. Especially Mg₂Al₂O₄-La₂O₃ addition improved the system, both in terms of cyclic stability and heat capacity.

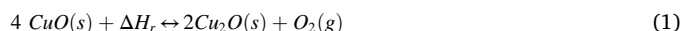
1. Introduction

Thermochemical energy storage (TCES) is a very promising technology which enables to address intermittent problems in renewable energy technologies, especially solar energy [1–5]. Continuous electricity production becomes possible with storing the energy during on-sun hours and releasing the stored energy during off-sun hours [6].

TCES exhibits better performance regarding to energy storage density (0.5–1 kWh/kg) [7], and storage within a large temperature range (25–1000°C) [8] among the other thermal energy storage techniques such as latent and sensible heat storage. In addition to this, there is no need to use complex storage procedures as the heat is stored in the form of chemical energy in TCES [9]. To store high amount of energy, redox reactions of metal oxides can be useful since their operational temperatures are relatively high [10]. For TCES applications, properties of the storage materials such as cyclic stability and agglomeration tendency are crucially [7,11,12].

From metal oxides, Co₃O₄, CuO and Mn₂O₃ exhibit sufficient performance to meet the expectations for TCES applications [13–15]. Among these oxides, Cu based oxides are also outstanding as they exhibit high energy storage capacity (811 kJ/kg for CuO) and oxygen

uncoupling ability [16,17]. From this point of view, CuO becomes very suitable as its usage would access more flexible design and process parameters for TCES applications [18,19]. The general reaction scheme for Cu oxides is given below (1).



However, occurrence of sintering due to both highly exothermic reaction nature of discharging step and comparably low Hüttig and Tammann temperatures of the copper oxide may cause the loss of cyclic stability and reactivity [8,20]. Hüttig and Tammann temperatures define as $0.3T_m$ and $0.5T_m$ (T_m is the melting point of the material) of the materials respectively and these temperatures indicate the onset of sintering temperature [21].

In the literature, high degree of reduction and high temperatures have been reported as conditions which should be avoided in case of CuO being used [22,23]. To overcome this problem, support materials have been commonly reported in similar applications in which cyclic stability and reactivity are important parameters [24,25]. Therefore, CuO could be supported by other oxide materials to increase the operational temperature of the application. Support materials are mainly used to decrease the sintering effect and supply additional vacant or

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available sites for oxygen and catalyst surface during redox reactions [26,27]. In the literature about redox of Cu based oxides there are plenty of support materials such as Al_2O_3 , ZrO_2 , SiO_2 , TiO_2 , MgAl_2O_4 and CeO_2 [21,22,28]. Among these support materials, MgAl_2O_4 was reported to provide sufficient physical stability and reactivity and ZrO_2 was reported as to decrease defluidization and agglomeration tendency [25, 29]. CeO_2 was also used as a support material with CuO resulting in high reactivity [30]. In addition, using La_2O_3 in CuO-ZrO_2 and $\text{CuO-MgAl}_2\text{O}_4$ increased the oxygen releasing ability and the agglomeration resistance [31,32].

In this study CuO with ZrO_2 , $\text{ZrO}_2\text{-La}_2\text{O}_3$, MgAl_2O_4 , $\text{MgAl}_2\text{O}_4\text{-La}_2\text{O}_3$, CeO_2 and $\text{CeO}_2\text{-La}_2\text{O}_3$ as support materials were used to investigate their thermochemical energy storage performance. Samples were tested in a fluidized bed reactor with a cyclic heating and cooling procedure. Simultaneous thermal analysis was also carried out to investigate the thermal characteristics of the samples. Phase analysis were carried out by X-Ray Diffraction to reveal the present phases both before and after the tests. Reaction mechanisms were also supported by thermodynamic equilibrium calculations.

2. Experimental procedure

The samples consisting of CuO (Trimano[®], Chemalloy[™]), ZrO_2 (Silverbond M800[®], Sibelco[™]), La_2O_3 , MgAl_2O_4 and CeO_2 used in this study were produced by VITO[™] using spray-drying. Raw materials with different composition (Table 1), organic binders and dispersants were dispersed in deionized water and mixed homogeneously. The water based suspension was then pumped into a hot drying chamber to obtain small solid loaded droplets. The spray-dried particles were sieved into 125-180 μm followed by calcination at 1100°C for 4 h in air. Table 1 also present works where these materials have been used before as oxygen carriers in chemical looping oxygen uncoupling (CLOU) applications.

Phase analysis of the samples both before and after being used in the cyclic experiments were done with XRD (Bruker[™] D8 Advance, $\text{Cu-K}\alpha$, 40 kV, 40 mA) in the 2-theta range of 15-80° with a step size of 0.01. Scanning electron microscopy (SEM-Zeiss LEO Ultra 55 FEG) was used to investigate the morphological characteristics. For secondary electron imaging 10.0 kV acceleration voltage and 10.0 mm working distance were used. To verify the experimentally observed interaction, thermodynamic equilibrium calculations were carried out using FactSage 7.2 software with the FactSage-Equilibrium Module, under the presumption of an isothermal and standard state with the use of Pure Substance (FactPS)-Oxide (FToxid) databases. The equilibrium calculations were based on the Gibbs energy minimization method.

For thermal analysis, simultaneous thermal analyzer (Netzsch[™]-STA 409 PC Luxx) was used to determine the thermal characteristics of the samples with changing the temperature between 600-1000°C as cycles. During the experiments, STA was flushed by air flow and N_2 as purge gas.

To investigate the redox and oxygen release characteristics of the samples, several heating and cooling steps were implemented in a fluidized bed reactor that has a length of 820 mm and a porous quartz plate of 22 mm in diameter placed 370 mm from the bottom [38]. The scheme of the fluidized bed setup is given in Figure S1 (Supplementary Material). For each experiment, the amount of sample was 10 g. The

Table 1
Chemical composition of the copper oxide based powders used in this study and the reference works used the same materials in the literature.

| Sample Name | Chemical Composition (wt.) | Ref. |
|-------------|---|------------------|
| CZ | 40% CuO , 60% ZrO_2 | [22,26,29,33,34] |
| CZL | 40% CuO , 55% ZrO_2 , 5% La_2O_3 | [22,31] |
| CM | 40% CuO , 60% MgAl_2O_4 | [22,25,34-36] |
| CML | 40% CuO , 55% MgAl_2O_4 , 5% La_2O_3 | [22,37] |
| CC | 40% CuO , 60% CeO_2 | [22,30] |
| CCL | 40% CuO , 55% CeO_2 , 5% La_2O_3 | [22,31] |

reduction and oxidation cycles were initiated by changing the temperature between 600°C and 1000°C for each cycle. It was reported that fixed partial pressure of O_2 , especially lower than 0.05 atm, results in different degree of oxygen deficiency in an O_2 release capable compound [21,39]. For this reason, the system was flushed by a gas mixture consisting of 5 vol.% O_2 (0.05 atm) which was obtained by N_2 diluted in air during all cycles. In one of the experiments, air was directly used without dilution of N_2 to investigate the effect of O_2 partial pressure on the system. The O_2 level in the gas were detected by a Rosemount[™] NGA 2000 multicomponent gas analyzer.

3. Results

As fluidized bed reactors have been reported as promising for thermochemical energy storage applications [18,40,41], investigation of the cyclic stability and oxygen releasing ability of the samples have in this work been carried in the fluidized bed. In the cyclic stability tests, the material was placed into the fluidized bed and cyclically heated up to 1000°C and cooled down to 600°C. Each cycle took on average 20 minutes. The temperature of the bed was measured by a thermocouple which was placed in the middle of the bed in the hot zone of the quartz reactor. When the temperature reached ~1000°C in the bed, the set point was decreased to 600°C, manually. In the cyclic stability test figures, the peaks shown on the upper part (yellow) represent the temperature changes in the bed. The peaks shown on the bottom part (blue) represent the observed O_2 concentration level in the gas analyzer connected to the reactor. The quantity of O_2 released/captured during the redox reactions is related to the reaction extent that corresponds to the amount of stored/released energy by the material [41,42]. In addition to this, the same amount of O_2 release and consumption in each cycle during the multi-cycle redox processes is consider as a sign of good cyclic stability [42-46].

Fig. 1 shows the cyclic stability test of the ZrO_2 supported CuO sample in the fluidized bed. In all experiments, the first cycle was excluded, since the first cycle is affected by the instrumental issues such as drift or temperature changes and is also influenced by how the sample was stored and treated before the experiments. Both for oxidation and reduction, very stable peaks were observed during the test. It has been reported that pure copper oxides exhibit strong agglomeration as the melting point of the material is close to the process temperature range [19]. Agglomeration is one of the main reasons for decreased redox activity of the materials [47]. From this point of view, ZrO_2 as support material for cyclic stability of CuO performed very well. During the cyclic stability test, re-oxidation peak point was observed at 900°C reaching min. peak point of 1,8vol.% O_2 in the outgoing gas while reduction was occurred at 1020°C reaching max. peak point of 10,99vol. % O_2 . The average mass change was observed via STA (simultaneous thermal analysis) for the same CZ sample with 0,8 wt.% and 1 wt.% during oxidation and reduction, respectively. In STA, the cycles were also stable, and the average re-oxidation and reduction peak temperatures were observed as 920 and 990°C, respectively. In addition to this, the peak areas which indicate the relative reaction enthalpies for oxidation and reduction were studied by using Netzsch Proteus[™] Software (Figure S2). Using the energy calibration, the specific energy content was calculated as 440 and 305 kJ/kg CZ for reduction and oxidation, respectively. XRD analysis on CZ samples, both fresh and after being tested in fluidized bed, revealed that there is no observed chemical interaction between $\text{CuO/Cu}_2\text{O}$ and ZrO_2 . This was also verified by thermodynamic equilibrium calculations (TEC).

As mentioned in the introduction, La_2O_3 has been reported as a promising support material for increasing the agglomeration resistance of CuO [22,32]. For this reason, La_2O_3 was used as a support material for CuO together with ZrO_2 (CZL). The cyclic stability of the material in the fluidized bed is shown in Fig. 2. The CZL sample, similar to CZ, also showed stable cyclic behaviour with a higher amount of average oxygen release starting at 1025°C (max. peak point of 12,11vol.% O_2) and

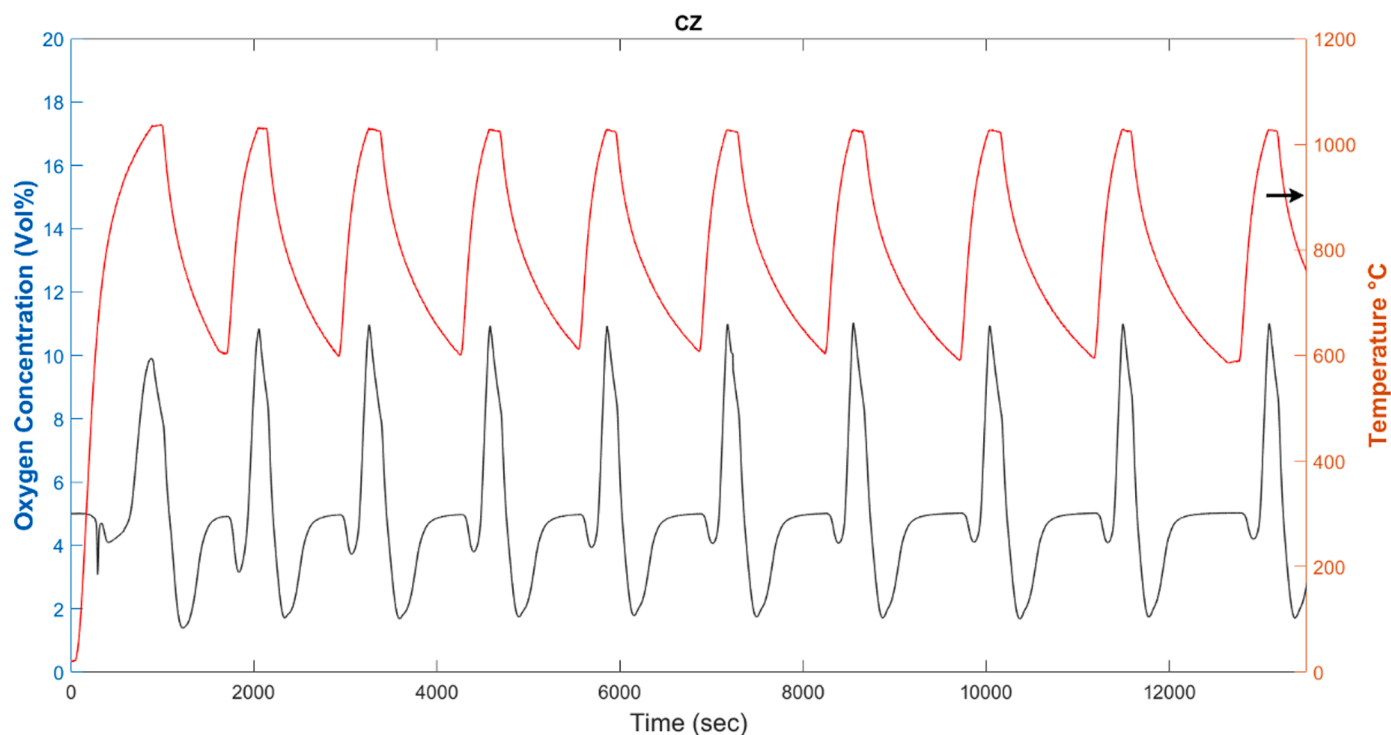


Fig. 1. Cyclic stability of the ZrO_2 supported CuO sample in the fluidized bed test.

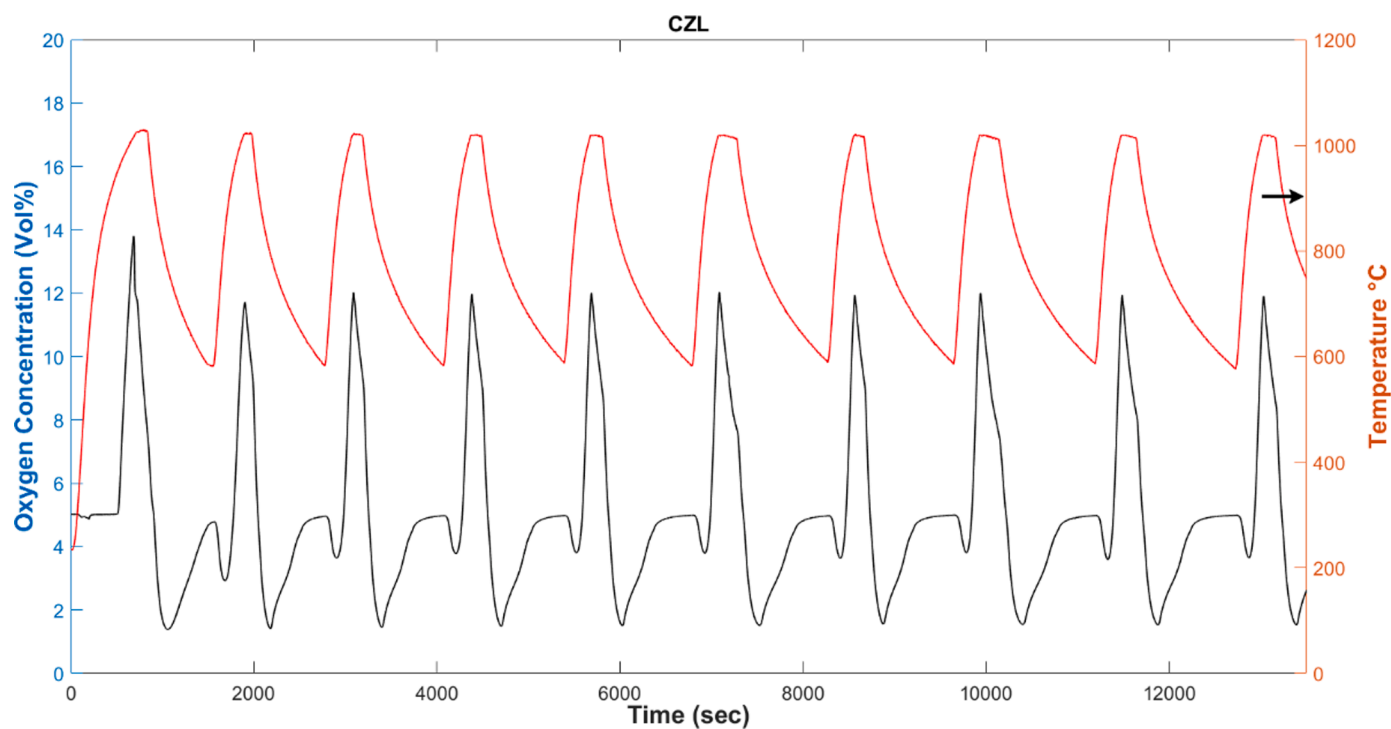


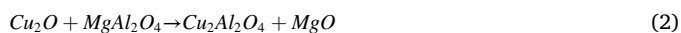
Fig. 2. Cyclic stability of the ZrO_2 - La_2O_3 supported CuO sample in the fluidized bed test.

consumption starting at 890°C (min. peak point of $1,46\text{vol.}\%O_2$). The thermal behaviour of the sample was also stable in STA and the average re-oxidation peak temperature was observed as 920°C , similar to CZ while the average reduction temperature was a little bit lower than CZ at 980°C . The average weight loss was $1,3\text{wt.}\%$ during reduction and the average weight gain was $1\text{ wt.}\%$ during oxidation. The specific energy content of the sample was calculated as 250 and 230 kJ/kg CZL for

reduction and oxidation, respectively. Similar to CZ, thermodynamic equilibrium calculations and XRD analysis after experiments also revealed that starting phases could form no new compounds during cycles.

When $MgAl_2O_4$ (CM) was used as a support material for CuO, the O_2 release performance was significantly better than CZ and CZL. The stability of the sample was outstanding both for reduction and oxidation

(Fig. 3). The average reduction peak temperature was 1010°C with the maximum 16,51 vol.%O₂ concentration while the average re-oxidation peak temperature was 870°C with the minimum 0,53 vol.% O₂ concentration in the fluidized bed tests. The cycle was also stable in STA and the average weight loss and weight gain were 1,01 and 1,06wt.%, respectively. The observed reduction peak temperature was 985°C which was similar to CZ and CZL. However, the observed average oxidation peak temperature was 930°C which is slightly higher than CZ and CZL. The specific energy content of the sample was calculated as 640 and 425 kJ/kg CM for reduction and oxidation, respectively. Thermodynamic equilibrium calculations suggested that there is a possible endothermic reaction (2) during reduction which may increase the agglomeration resistance of copper oxides as the reaction is reversible and the Cu₂Al₂O₄ has a higher melting point than pure Cu₂O [48]. Since this reaction is not thermodynamically favourable below 900°C, there was no observed phases in XRD related to this. However, oxidation of Cu₂Al₂O₄ into CuO and reforming of MgAl₂O₄ spinel is thermodynamically favourable during oxidation. From this point of view, MgAl₂O₄ is a promising candidate as support for CuO for TCES applications.



When La₂O₃ was also used as a support material together with MgAl₂O₄ spinel (CML sample), the results were even more promising. Fig. 4 shows the cyclic stability of CML in the fluidized bed test. CML performed outstanding cyclic stability with the highest O₂ release/consumption among the samples we tested in this study. For CML, the minimum and maximum O₂ concentration during oxidation and reduction were observed as 0,54 vol.% O₂ and 16,71 vol.%O₂, respectively. The average peak temperature was 820°C and 1010°C for re-oxidation and reduction, respectively. Thermal gravimetric behaviour of the material was also stable in STA analysis. Similar to the samples tested, the average peak temperature was 920°C for oxidation and 970°C for reduction which is slightly lower than others. However, the average weight loss was observed as 3,07 wt.% during reduction which is significantly higher than samples tested in this study. Similar to this, the average weight gain was obtained as 1,24 wt.%. As already mentioned

above, the weight changes during redox is related to the energy storage capacity of the system. From this point of view, CML exhibited the best performance among the tested samples as it showed the highest weight gain and weight loss in STA analysis. The specific energy content of the sample was calculated as 925 and 1050 kJ/kg CML for reduction and oxidation, respectively. However, the obtained weight changes during redox reactions were still lower than maximum theoretical weight changes for a CuO/Cu₂O redox systems with 60% support. This deviation may be caused by the effect of heating/cooling rates, particle size and distribution of the samples and the amount of used mass as it is known that thermodynamic and kinetic factors are limiting the redox reactions [16,26,49,50].

In Fig. 5, cyclic stability of CC (CeO₂ supported CuO) was shown. Among the tested samples, CC showed the poorest O₂ release/consumption capacity, even though the redox cycles were very stable. During the fluidized bed test, generally reduction reaction reached the peak point at 1020°C, while this was 840°C for re-oxidation. On average, the maximum obtained O₂ concentration was 11,2 vol.%O₂ during reduction and minimum obtained O₂ concentration was 1,76 vol.%O₂ during oxidation which is similar to other samples. However, the cyclic stability was not stable as the others in thermal analysis. For STA, the reduction took place at maximum 990°C with the average weight loss around 0,5 wt.%. In addition to this, re-oxidation in STA took place at maximum 950°C with the weight gain around 0,5 wt.%. La₂O₃ addition to CeO₂ as support material for CuO (CCL), increased the O₂ release/consumption ability compare to CC. However, the performance of CCL could not reach the performance of the other tested samples. A good thing about this sample was the almost perfect cyclic behaviour which is important for material's lifetime [51]. For CCL(Fig. 6), re-oxidation generally started to occur at 820°C and average O₂ concentration obtained was 0,25 vol.% during oxidation. These values were 1020°C and 12,58 vol.%O₂ for reduction peak and the highest obtained O₂ concentration, respectively. The cyclic stability of CCL was significantly better than CC in STA. In STA, the sample lost weight around 0,8 wt.% during reduction which generally took place at 980°C as peak maximum. For re-oxidation, the temperature was 920°C and the average weight gain was 0,7 wt.%.

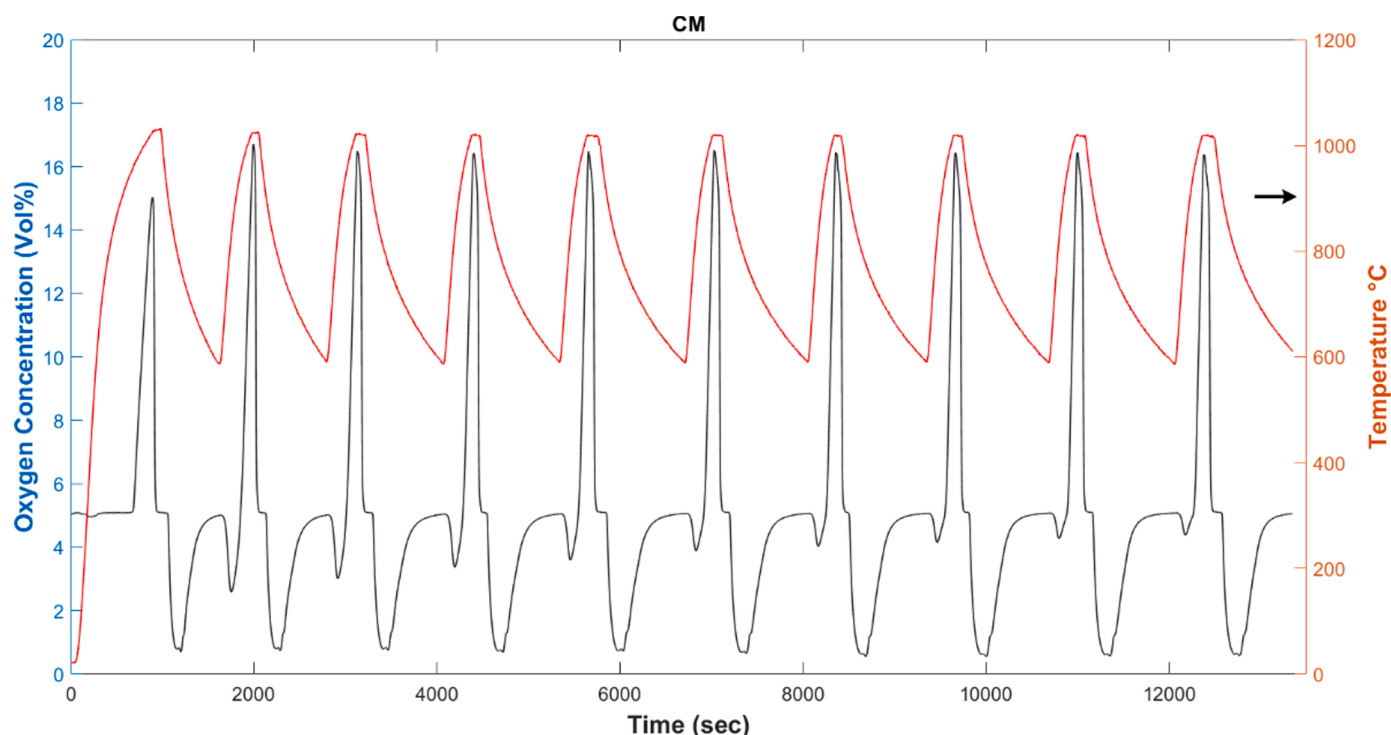


Fig. 3. Cyclic stability of the MgAl₂O₄ supported CuO sample in the fluidized bed test.

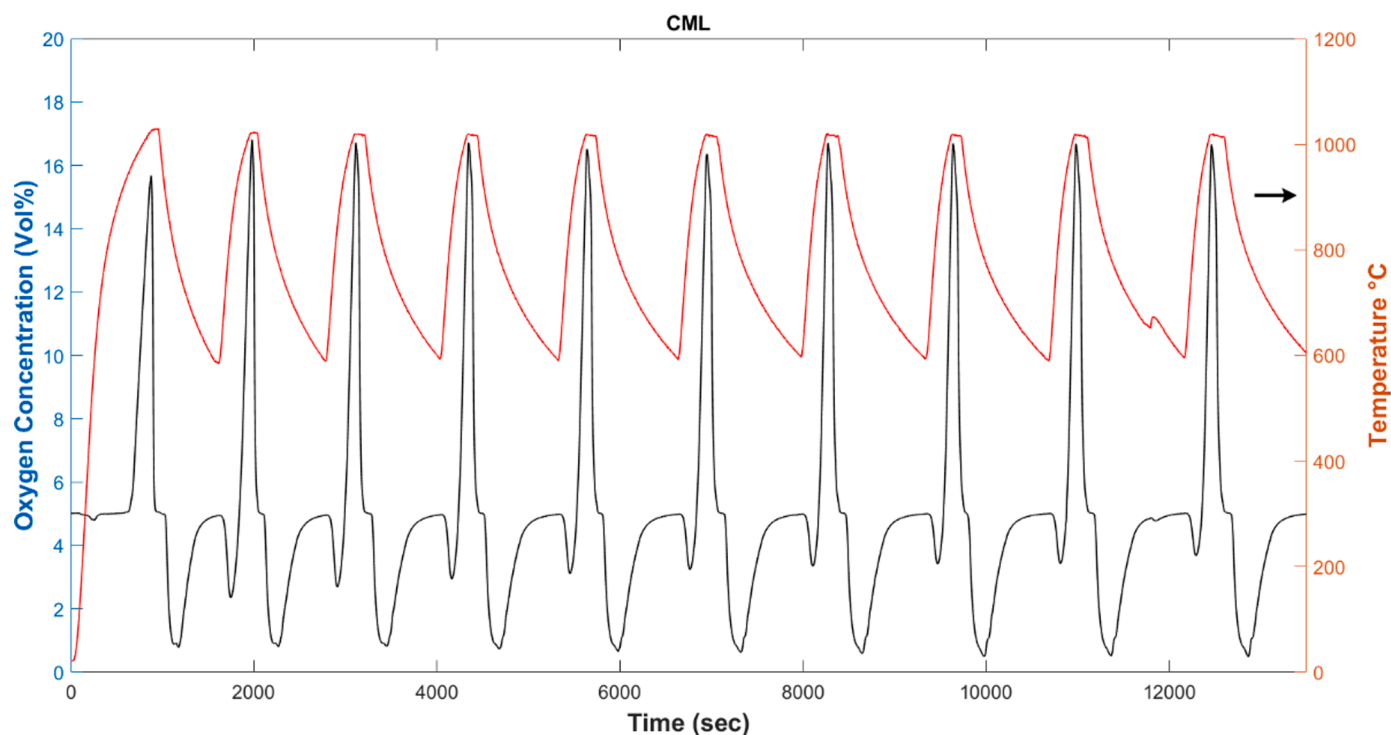


Fig. 4. Cyclic stability of the $\text{MgAl}_2\text{O}_2\text{-La}_2\text{O}_3$ supported CuO sample in the fluidized bed test.

A general overview of phase changes of the used materials can be seen in Table 2. This table includes both starting phases in the fresh samples and observed phases in the used samples after being tested in the fluidized bed is shown. Additionally, thermodynamic equilibrium calculations were also done to see how much the system differs from the thermodynamic equilibrium conditions. Formations of $\text{La}_2\text{Zr}_2\text{O}_7$, LaAlO_3 and $\text{Ce}_{0.75}\text{La}_{0.25}\text{O}_2$ were observed in the CZL, CML and CCL, respectively. To avoid the possible deactivation of active sites, support

materials should not form any compound with $\text{CuO/Cu}_2\text{O}$. However, these compounds do not show any interaction towards $\text{CuO/Cu}_2\text{O}$. On the contrary, these formations may be the reason of the slight increase of the cyclic stability for CZ-CZL, CM-CML and CC-CCL. Especially LaAlO_3 is known a potential candidate for energy storage applications [52]. In addition to this, LaAlO_3 has a high melting point (2138°C) [53]. Therefore, it does not contribute to the agglomerate formation in the system. Overall, it is clear that CML has a great potential for this study.

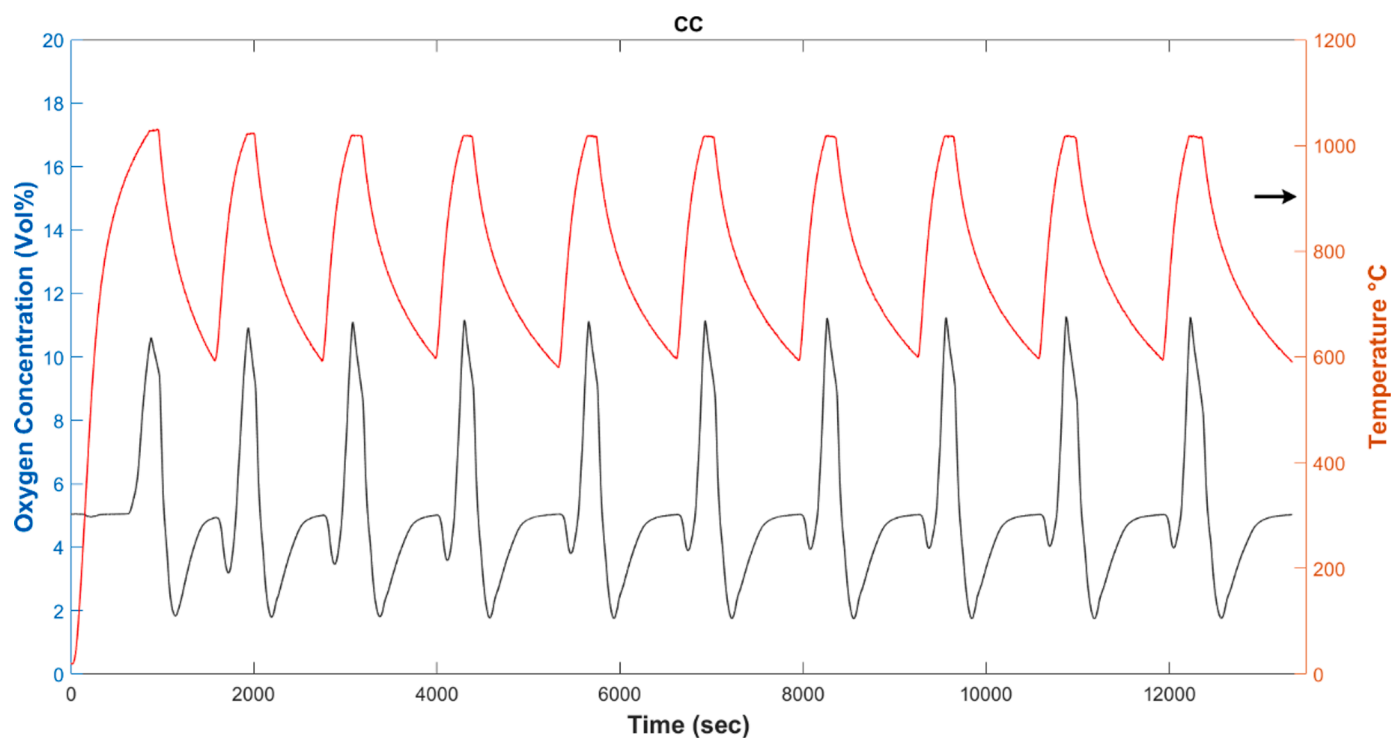


Fig. 5. Cyclic stability of the CeO_2 supported CuO sample in the fluidized bed test.

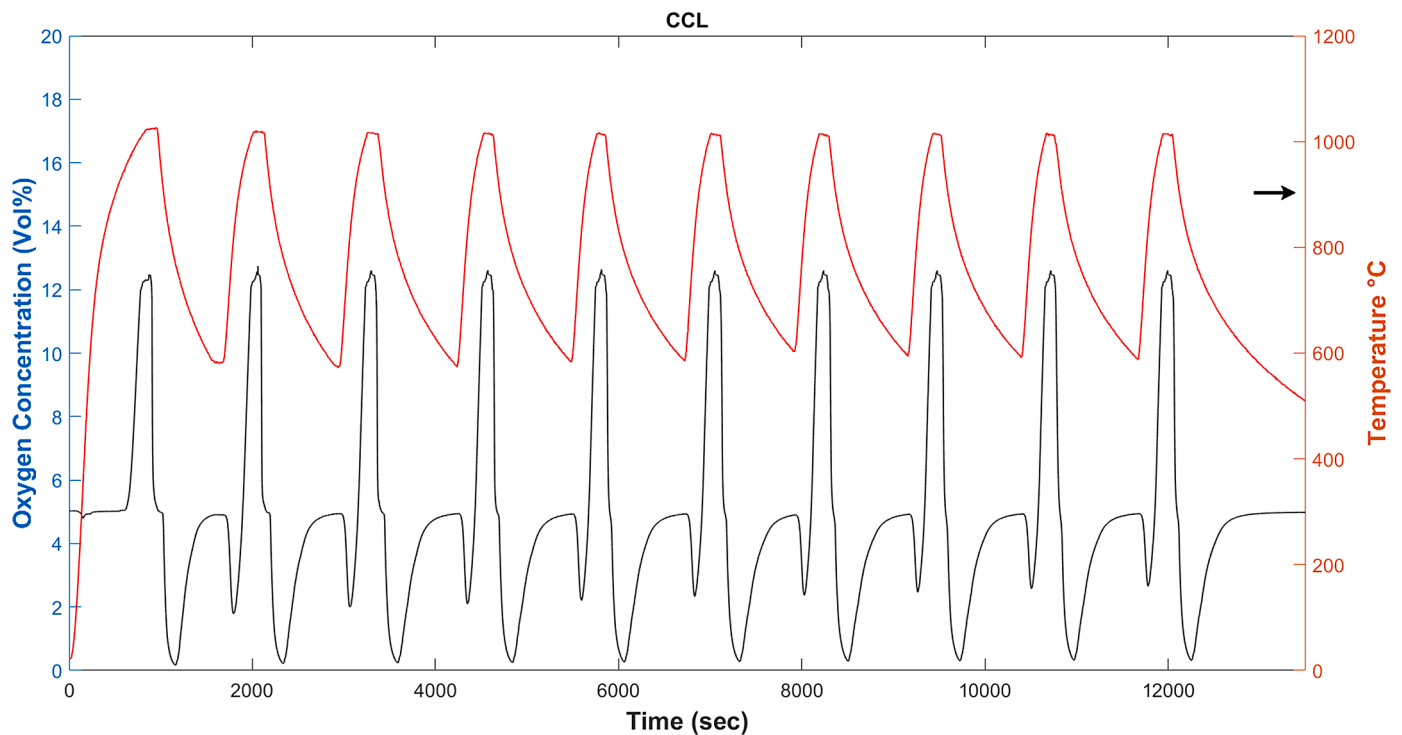


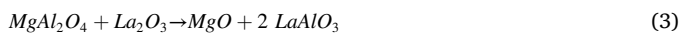
Fig. 6. Cyclic stability of the $\text{CeO}_2\text{-La}_2\text{O}_3$ supported CuO sample in the fluidized bed test.

Table 2

Phases revealed by XRD analysis in the starting samples and observed ones after the cyclic stability tests.

| Sample | Starting Phases | Observed Phases After Tests | Calculated Phases by TEC |
|--------|--|---|---|
| CZ | CuO, ZrO ₂ | CuO, ZrO ₂ | CuO, ZrO ₂ |
| CZL | CuO, ZrO ₂ , La ₂ O ₃ | CuO, ZrO ₂ , La ₂ Zr ₂ O ₇ | CuO, ZrO ₂ , La ₂ O ₃ |
| CM | CuO, MgAl ₂ O ₄ | CuO, MgAl ₂ O ₄ | CuO, MgAl ₂ O ₄ |
| CML | CuO, MgAl ₂ O ₄ , La ₂ O ₃ | CuO, MgAl ₂ O ₄ , LaAlO ₃ | CuO, MgO, MgAl ₂ O ₄ , LaAlO ₃ |
| CC | CuO, CeO ₂ | CuO, CeO ₂ | CuO, CeO ₂ |
| CCL | CuO, CeO ₂ , La ₂ O ₃ | CuO, CeO ₂ , La ₂ O ₃ , Ce _{0.75} La _{0.25} O ₂ | CuO, CeO ₂ , La ₂ O ₃ |

To investigate further the best of the materials, CML, XRD analysis was carried out for fresh and tested sample (Fig. 7). Fresh CML, consisting of CuO (PDF Card No: 48-1548), MgAl₂O₄ (PDF: 04-007-2712) and La₂O₃ (PDF: 04-015-5007), formed some perovskite-type LaAlO₃ (PDF: 04-18-6998) during fluidized bed test. Formation of LaAlO₃, which most likely formed via Reaction 3, was also confirmed by thermodynamic equilibrium calculations.



LaAlO₃ was reported as a good candidate for solar thermochemical applications due to its stability under redox conditions [52]. Most likely, together with MgAl₂O₄, it increased the physicochemical stability of the sample during redox reactions. To reveal the physicochemical condition of the particles, scanning electron microscope was used in secondary electron mode (Fig. 8). Fresh sample exhibited an even morphology consisting of decent spheres. Even though the majority of the particles preserved this morphology, some cracked and decomposed particles were observed for used particles.

To investigate the effect of partial O₂ pressure on the cyclic stability of CML, the fluidized bed tests were carried out under two different O₂ concentration (Fig. 9). When the partial O₂ pressure in the system was

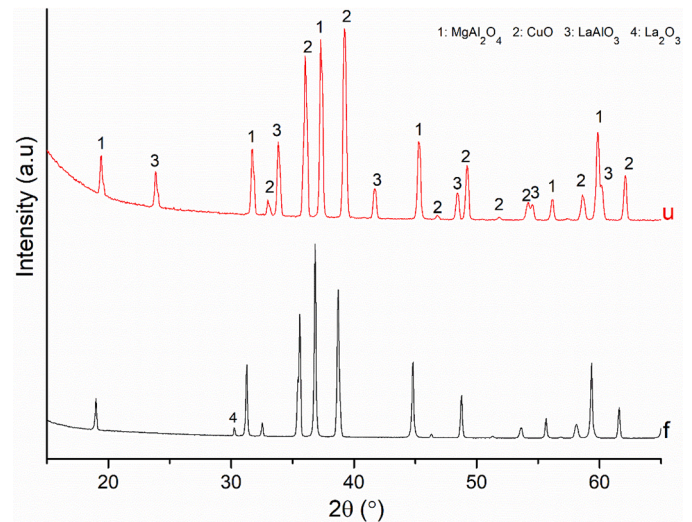


Fig. 7. XRD patterns of fresh and used $\text{MgAl}_2\text{O}_4\text{-La}_2\text{O}_3$ supported CuO sample.

0,05 atm, CML exhibited an outstanding performance compare to the case in which the partial O₂ pressure was 0,21 atm as air. This result was expected since equilibrium partial O₂ pressure of Cu-oxides is lower than 0,21 for the relevant temperature of this study [21,54]. However, 21 vol. % O₂ concentration was also used on the sample to reveal the result in case of using pure air (non-diluted with N₂) in the system.

4. Discussion

In this study, ZrO₂, La₂O₃, MgAl₂O₄ and CeO₂ supported Cu based oxide materials were tested as potential thermochemical energy storage materials. The chemical reactions and oxides systems used are similar in TCES and chemical looping oxygen uncoupling (CLOU). Therefore the

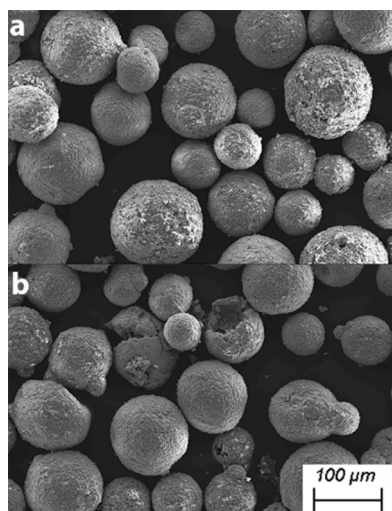


Fig. 8. Secondary electron SEM micrograph of fresh (a) and used (b) $\text{MgAl}_2\text{O}_2\text{-La}_2\text{O}_3$ supported CuO sample.

materials which have previously been used in CLOU applications [25,33, 55–62] were tested as potential energy storage materials in this study. The purpose of the support materials are to increase the operational temperature, compared to pure Cu-oxides; and decrease the agglomeration tendency [26,32]. For this reason, it was important to avoid possible new compound forming between CuO and the support material, which may decrease the performance of the CuO/Cu₂O system. After the tests, there were no formation of phases observed which may deactivate the copper oxides. However, $\text{La}_2\text{Zr}_2\text{O}_7$, LaAlO_3 and $\text{Ce}_{0,75}\text{La}_{0,25}\text{O}_2$ was formed which could not be observed during initial synthesis of the materials. Even though newly formed compounds do not directly affect the copper oxide, thermal stability of the material is important. As

outstanding thermal stability was reported for $\text{La}_2\text{Zr}_2\text{O}_7$ and LaAlO_3 , formation of these compounds may affect the active materials lifetime in a positive manner. In addition to this, no defluidization was observed during the tests which means that agglomeration of the sample was avoided. Visual inspection after the fluidized bed tests also revealed neither sintering nor agglomeration in the bed.

Table 3 is given to summarize the results. Both thermal analysis and fluidized bed reactor tests were carried out in this work. Fluidized bed reactor tests enable to use comparably large amount of samples which may be a representative of more homogeneous sampling and also show the physical behaviour of the particles such as attrition or tendency for agglomeration. On the other hand, more accurate temperature and weight changes can be obtained by thermal analysis. For this reason, these two methods are considered as supplementary. Theoretical weight gain and weight loss for reduction of CuO to Cu₂O are 11,18wt.% and 10,06wt.%, respectively. MgAl_2O_4 and $\text{MgAl}_2\text{O}_4\text{-La}_2\text{O}_3$ supported CuO showed the best O₂ release/consumption capacity in both fluidized bed reactor test and thermal analysis among the tested samples, which are closer to the theoretical weight gain and loss values. In all fluidized bed test, the intermittent oxidation was observed most likely caused by the slow kinetics of reoxidation. For each sample, the oxygen uptake after the reduction got slower during the cooling and carried over with the following heating step. In fluidized bed reactor tests, the reduction peak temperatures were only 25–40°C higher than the obtained in the thermal analysis. However, the observed oxidation peak temperatures in thermal analysis were significantly higher than fluidized bed reactor tests for CZL, CML, and CCL. This difference most likely occurred due to the formation of new compounds.

The development of supported CuO oxides is necessary regarding to reversibility, reactivity loss over cycles and sintering in order to obtain cost effective, environmentally friendly and for large scale applications. The support materials used in this study showed very good performance regarding to the cyclic stability of the active phase. Even though, the high purity support materials are just slightly cheaper than the pure copper oxides, they may increase the lifetime of the active material. In

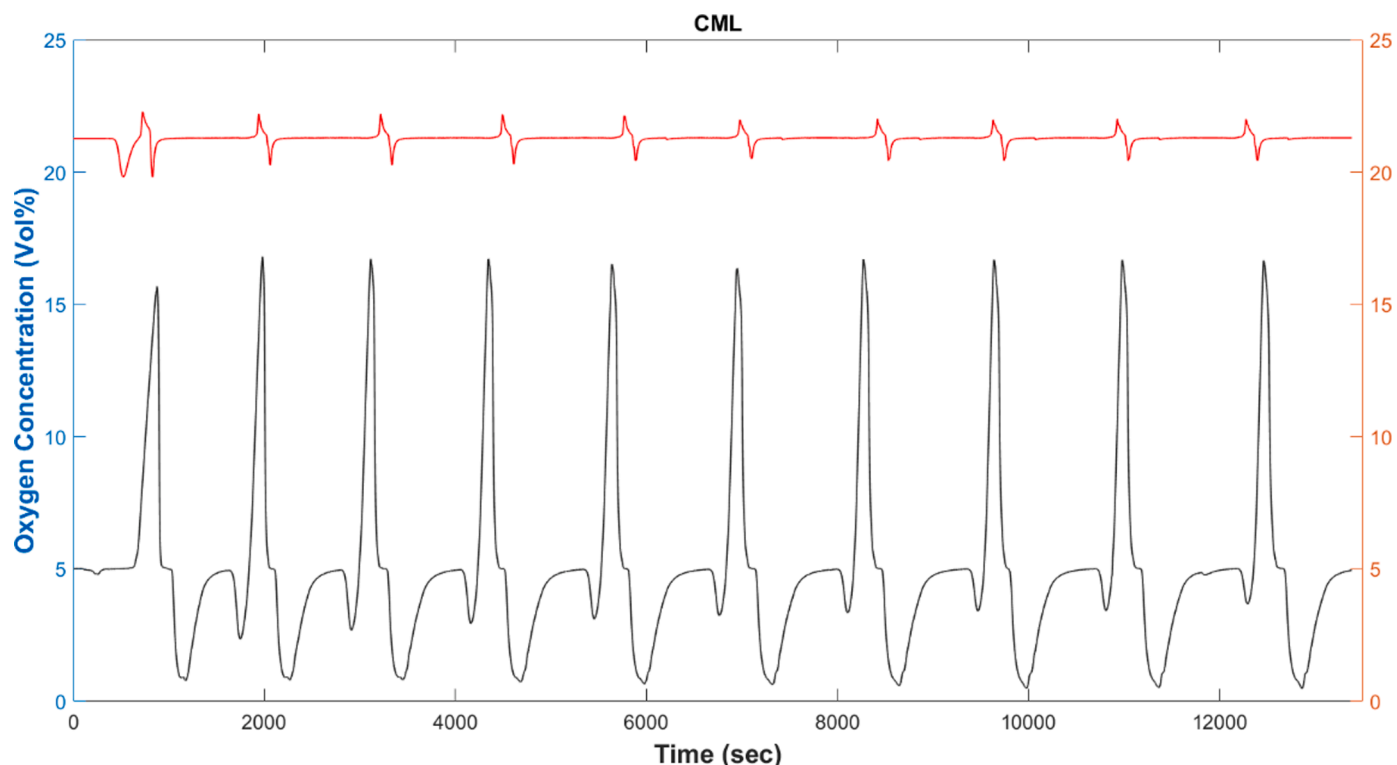


Fig. 9. Comparison of the cyclic stability of the $\text{MgAl}_2\text{O}_2\text{-La}_2\text{O}_3$ supported CuO sample under 5 vol.% O₂ and 21 vol.% O₂ atmosphere in the fluidized bed test.

Table 3

Summary of the results of the cyclic stability tests in the fluidized bed reactor and thermal analysis.

| Sample | Fluidized Bed Reactor Tests | | | | Thermal Analysis | | | |
|--------|--|--|--|--|--|---------------------|--|---------------------|
| | Reduction Peak Temperature (T_{peak}) | Peak Max. O ₂ /Release (vol. %) | Oxidation Peak Temperature (T_{peak}) | Peak Min. O ₂ /Consumption (vol. %) | Reduction Peak Temperature (T_{peak}) | Weight Loss (wt. %) | Oxidation Peak Temperature (T_{peak}) | Weight Gain (wt. %) |
| CZ | 1020 | 10,99 | 900 | 1,8 | 990 | 0,98 | 920 | 0,8 |
| CZL | 1025 | 12,11 | 890 | 1,46 | 980 | 1,30 | 950 | 1,03 |
| CM | 1010 | 16,51 | 870 | 0,53 | 985 | 1,01 | 910 | 1,06 |
| CML | 1010 | 16,71 | 820 | 0,54 | 970 | 3,07 | 920 | 2,4 |
| CC | 1020 | 11,20 | 840 | 1,76 | 990 | 0,5 | 890 | 0,5 |
| CCL | 1020 | 12,58 | 820 | 0,25 | 980 | 0,8 | 920 | 0,7 |

Table 4

Heat content of the relevant oxides [63].

| Oxides | $H_{1200}^{\circ} - H_{298}^{\circ} \left(\frac{\text{kJ}}{\text{mole}} \right)$ |
|----------------------------------|---|
| CuO | 47 |
| Cu ₂ O | 67 |
| ZrO ₂ | 64 |
| La ₂ O ₃ | 114 |
| MgAl ₂ O ₄ | 147 |
| LaAlO ₃ | 109 |

In addition to this, as the redox reactions of copper oxides take place at high temperatures, the sensible heat form of the storage of interest is also important. In Table 4, the heat content of the supporting oxides used in this study is given. From this point of view, even the non-active, support phase of the sample contributes for storing the heat. Among the tested samples in the study, CM and CML are very promising in terms of both cost effectivity and abundancy of MgO and Al₂O₃. Especially, CML showed the best performance with the highest specific energy content (925 and 1050 kJ/kg CML for reduction and oxidation, respectively) which may lead the reduction of the raw material costs.

5. Conclusion

In the present study, ZrO₂, MgAl₂O₄, CeO₂ and La₂O₃ were used as support materials to improve the cyclic stability performance of copper oxide materials for thermochemical energy storage applications. Even though CuO has excellent properties for thermochemical energy storage applications, use of pure CuO is limited due to its high decomposition temperature in air (1034°C) and low agglomeration resistance above the relevant temperature limits. Among the oxides which have been reported in the literature as support material for CuO, ZrO₂, MgAl₂O₄, CeO₂ and La₂O₃ were chosen owing to their proved success in the previous studies for other applications. The cyclic stability tests were carried out in a fluidized bed reactor. Fluidized bed tests are significantly promoted for thermochemical energy storage application studies as it enables physical testing of large amount of samples. Obtained results were also compared to results from the thermal analysis. Among the tested samples, MgAl₂O₄ and MgAl₂O₄/La₂O₃ supported CuO exhibited the best O₂ release/consumption capacity. In fluidized bed tests, CML which showed the best performance exhibited 1,5 times more O₂ release and 3 times more O₂ consumption than CZ which showed the poorest performance. For STA, CML performed averagely 6 times more weight loss and 5 times more weight gain than the poorest performance CC. As MgAl₂O₄ is comparably cheaper than other used oxides and the amount of La₂O₃ was quite low (5 wt.%) in the composition, this result was quite promising for the application. The poorest O₂ release/consumption capacity was observed for CeO₂ supported CuO which was quite unexpected since CeO₂ has been used for its catalyst effect on Cu. However, CeO₂ and La₂O₃ supported CuO samples exhibited an identical cyclic behavior and reversibility which are important for material's lifetime.

CRediT authorship contribution statement

Duygu Yilmaz: Writing - original draft, Investigation, Conceptualization. **Esraa Darwish:** Investigation, Methodology. **Henrik Leion:** Conceptualization, Writing - review & editing, Supervision.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Supplementary materials

Supplementary material associated with this article can be found, in the online version, at doi:10.1016/j.est.2020.102012.

References

- [1] M. Kubicek, A.H. Bork, J.L.M. Rupp, Perovskite oxides – a review on a versatile material class for solar-to-fuel conversion processes, *J. Mater. Chem. A* 5 (2017) 11983–12000, <https://doi.org/10.1039/C7TA00987A>.
- [2] A. Evans, V. Strezov, T.J. Evans, Assessment of utility energy storage options for increased renewable energy penetration, *Renew. Sustain. Energy Rev.* 16 (2012) 4141–4147, <https://doi.org/10.1016/j.rser.2012.03.048>.
- [3] P. Pardo, A. Deydier, Z. Anxionnaz-minvielle, S. Rougé, M. Cabassud, P. Cognet, P. Pardo, A. Deydier, Z. Anxionnaz-minvielle, S. Rougé, M. Cabassud, A review on high temperature thermochemical heat energy storage, *Renew. Sustain. Energy Rev.* 32 (2016) 591–610.
- [4] L.F. Zalba, M.J.B. Cabeza, Review on Phase changing materials to store energy, *Appl. Therm. Eng.* 23 (2003) 251–283, [https://doi.org/10.1016/S1359-4311\(02\)00192-8](https://doi.org/10.1016/S1359-4311(02)00192-8).
- [5] E. Guy, Solar heat storage using chemical reactions, *J. Solid State Chem.* 22 (1977) 51–61.
- [6] W.E. Wentworth, E. Chen, Simple thermal decomposition reactions for storage of solar thermal energy, *Solar Energy* 18 (1976) 205–214, [https://doi.org/10.1016/0038-092X\(76\)90019-0](https://doi.org/10.1016/0038-092X(76)90019-0).
- [7] L. André, S. Abanades, G. Flamant, Screening of thermochemical systems based on solid-gas reversible reactions for high temperature solar thermal energy storage, *Renew. Sustain. Energy Rev.* 64 (2016) 703–715, <https://doi.org/10.1016/j.rser.2016.06.043>.
- [8] L. André, S. Abanades, L. Cassayre, Experimental investigation of Co-Cu, Mn-Co, and Mn-Cu redox materials applied to solar thermochemical energy storage, *ACS Appl. Energy Mater.* 1 (2018) 3385–3395, <https://doi.org/10.1021/acsaem.8b00554>.
- [9] B. Michel, N. Mazet, S. Mauran, D. Stitou, J. Xu, Thermochemical process for seasonal storage of solar energy: characterization and modeling of a high density reactive bed, *Energy* 47 (2012) 553–563, <https://doi.org/10.1016/j.energy.2012.09.029>.
- [10] P. Pardo, A. Deydier, Z. Anxionnaz-Minvielle, S. Rougé, M. Cabassud, P. Cognet, A review on high temperature thermochemical heat energy storage, *Renew. Sustain. Energy Rev.* 32 (2014) 591–610, <https://doi.org/10.1016/j.rser.2013.12.014>.
- [11] S. Wu, C. Zhou, E. Doroodchi, R. Nellore, B. Moghtaderi, A review on high-temperature thermochemical energy storage based on metal oxides redox cycle, *Energy Convers. Manage.* 168 (2018) 421–453, <https://doi.org/10.1016/j.enconman.2018.05.017>.

- [12] L. Duan, D. Godino, V. Manovic, F. Montagnaro, E.J. Anthony, Cyclic oxygen release characteristics of bifunctional copper oxide/calcium oxide composites, *Energy Technol.* 4 (2016) 1171–1178, <https://doi.org/10.1002/ente.201600028>.
- [13] T. Block, M. Schmäcker, Metal oxides for thermochemical energy storage: a comparison of several metal oxide systems, *Solar Energy* 126 (2016) 195–207, <https://doi.org/10.1016/j.solener.2015.12.002>.
- [14] A.J. Carrillo, D.P. Serrano, P. Pizarro, J.M. Coronado, Manganese oxide-based thermochemical energy storage: modulating temperatures of redox cycles by Fe-Cu co-doping, *J. Energy Storage* 5 (2016) 169–176, <https://doi.org/10.1016/j.est.2015.12.005>.
- [15] M. Silakhori, M. Jafarian, M. Arjomandi, G.J. Nathan, Thermogravimetric analysis of Cu, Mn, Co, and Pb oxides for thermochemical energy storage, *J. Energy Storage* 23 (2019) 138–147, <https://doi.org/10.1016/j.est.2019.03.008>.
- [16] M. Deutsch, F. Horvath, C. Knoll, D. Lager, C. Gierl-Mayer, P. Weinberger, F. Winter, High-temperature energy storage: kinetic investigations of the CuO/Cu₂O reaction cycle, *Energy Fuels* 31 (2017) 2324–2334, <https://doi.org/10.1021/acs.energyfuels.6b02343>.
- [17] T. Mattisson, A. Lyngfelt, H. Leion, Chemical-looping with oxygen uncoupling for combustion of solid fuels, *Int. J. Greenh. Gas Control* 3 (2009) 11–19, <https://doi.org/10.1016/j.egypro.2009.01.060>.
- [18] B. Wong, Thermochemical heat storage for concentrated solar power, thermochemical system reactor design for thermal energy storage, *Ind. Eng. Chem.* (2011), <https://doi.org/10.2172/1039304>.
- [19] E. Alonso, C. Pérez-Rábago, J. Licurgo, E. Fuentealba, C.A. Estrada, First experimental studies of solar redox reactions of copper oxides for thermochemical energy storage, *Solar Energy* 115 (2015) 297–305, <https://doi.org/10.1016/j.solener.2015.03.005>.
- [20] M. Hänchen, A. Stiel, Z.R. Jovanovic, A. Steinfeld, Thermally driven copper oxide redox cycle for the separation of oxygen from gases, *Ind. Eng. Chem. Res.* 51 (2012) 7013–7021, <https://doi.org/10.1021/ie202474s>.
- [21] Q. Imtiaz, D. Hosseini, C.R. Müller, Review of oxygen carriers for chemical looping with oxygen uncoupling (CLOU): thermodynamics, material development, and synthesis, *Energy Technol.* 1 (2013) 633–647, <https://doi.org/10.1002/ente.201300099>.
- [22] M. Rydén, D. Jing, M. Källén, H. Leion, A. Lyngfelt, T. Mattisson, CuO-based oxygen-carrier particles for chemical-looping with oxygen uncoupling - experiments in batch reactor and in continuous operation, *Ind. Eng. Chem. Res.* 53 (2014) 6255–6267, <https://doi.org/10.1021/ie4039983>.
- [23] V. Frick, M. Rydén, H. Leion, Investigation of Cu-Fe and Mn-Ni oxides as oxygen carriers for chemical-looping combustion, *Fuel Process. Technol.* 150 (2016) 30–40, <https://doi.org/10.1016/j.fuproc.2016.04.032>.
- [24] M. Keller, J. Fung, H. Leion, T. Mattisson, Cu-impregnated alumina/silica bed materials for chemical looping reforming of biomass gasification gas, *Fuel* 180 (2016) 448–456, <https://doi.org/10.1016/j.fuel.2016.04.024>.
- [25] M. Arjomand, A. Azad, H. Leion, A. Lyngfelt, T. Mattisson, Prospects of Al₂O₃ and MgAl₂O₄ supported CuO oxygen carriers in chemical-looping combustion (CLC) and chemical-looping with oxygen uncoupling (CLOU), *Energy Fuels* 25 (2011) 5493–5502, <https://doi.org/10.1021/ef201329x>.
- [26] C.K. Clayton, K.J. Whitty, Measurement and modeling of decomposition kinetics for copper oxide-based chemical looping with oxygen uncoupling, *Appl Energy* 116 (2014) 416–423, <https://doi.org/10.1016/j.apenergy.2013.10.032>.
- [27] M.S.C. Chan, H.G. Baldovi, J.S. Dennis, Enhancing the capacity of oxygen carriers for selective oxidations through phase cooperation: bismuth oxide and ceria-zirconia, *Catal. Sci. Technol.* 8 (2018) 887–897, <https://doi.org/10.1039/c7cy01992k>.
- [28] L.F. de Diego, P. Gayán, F. García-Labiano, J. Celaya, A. Abad, J. Adánez, Impregnated CuO/Al₂O₃ oxygen carriers for chemical-looping combustion: avoiding fluidized bed agglomeration, *Energy Fuels* 19 (2005) 1850–1856, <https://doi.org/10.1021/ef050052f>.
- [29] M. Arjomand, A.M. Azad, H. Leion, M. Rydén, T. Mattisson, ZrO₂ supported CuO oxygen carriers for chemical-looping with oxygen uncoupling (CLOU), *Energy Procedia* 37 (2013) 550–559, <https://doi.org/10.1016/j.egypro.2014.07.009>.
- [30] A. Hedayati, A.M. Azad, M. Rydén, H. Leion, T. Mattisson, Evaluation of novel ceria-supported metal oxides as oxygen carriers for chemical-looping combustion, *Ind Eng Chem Res* 51 (2012) 12796–12806, <https://doi.org/10.1021/ie300168j>.
- [31] D. Jing, T. Mattisson, M. Ryden, P. Hallberg, A. Hedayati, J. Van Noyen, F. Snijders, A. Lyngfelt, Innovative oxygen carrier materials for chemical-looping combustion, *Energy Procedia* 37 (2013) 645–653, <https://doi.org/10.1016/j.egypro.2013.05.152>.
- [32] Y. Cao, H.Y. Zhao, S.P. Sit, W.P. Pan, Lanthanum-promoted copper-based oxygen carriers for chemical looping combustion process, *J. Therm. Anal. Calorim.* 116 (2014) 1257–1266, <https://doi.org/10.1007/s10973-013-3628-8>.
- [33] T. Mattisson, H. Leion, A. Lyngfelt, Chemical-looping with oxygen uncoupling using CuO/ZrO₂ with petroleum coke, *Fuel* 88 (2009) 683–690, <https://doi.org/10.1016/j.fuel.2008.09.016>.
- [34] M. Keller, H. Leion, T. Mattisson, H. Thunman, Investigation of natural and synthetic bed materials for their utilization in chemical looping reforming for tar elimination in biomass-derived gasification gas, *Energy Fuels* 28 (2014) 3833–3840, <https://doi.org/10.1021/ef500369c>.
- [35] M. Arjomand, M. Keller, H. Leion, T. Mattisson, A. Lyngfelt, Oxygen release and oxidation rates of MgAl₂O₄-supported CuO oxygen carrier for chemical-looping combustion with oxygen uncoupling (CLOU), *Energy Fuels* 26 (2012) 6528–6539, <https://doi.org/10.1021/ef3010064>.
- [36] I. Adánez-Rubio, P. Gayán, A. Abad, F. García-Labiano, L.F.F. de Diego, J. Adánez, Kinetic analysis of a Cu-based oxygen carrier: relevance of temperature and oxygen partial pressure on reduction and oxidation reactions rates in chemical looping with oxygen uncoupling (CLOU), *Chem. Eng. J.* 256 (2014) 69–84, <https://doi.org/10.1016/j.cej.2014.06.102>.
- [37] D. Jing, T. Mattisson, M. Ryden, P. Hallberg, A. Hedayati, J. Van Noyen, F. Snijders, A. Lyngfelt, Innovative oxygen carrier materials for chemical-looping combustion, *Energy Procedia* 37 (2013) 645–653, <https://doi.org/10.1016/j.egypro.2013.05.152>.
- [38] H. Leion, V. Frick, F. Hildor, Experimental method and setup for laboratory fluidized bed reactor testing, *Energies* (2018) 11, <https://doi.org/10.3390/en11020505>.
- [39] E.I. Leonidova, I.A. Leonidov, M.V. Patrakeev, V.L. Kozhevnikov, Oxygen non-stoichiometry, high-temperature properties, and phase diagram of CaMnO_{3-δ}, *J. Solid State Electrochem.* 15 (2011) 1071–1075, <https://doi.org/10.1007/s10008-010-1288-1>.
- [40] C. Tregambi, F. Montagnaro, P. Salatino, R. Solimene, Directly irradiated fluidized bed reactors for thermochemical processing and energy storage: application to calcium looping, in: AIP Conference Proceedings 1850, 2017, <https://doi.org/10.1063/1.4984456>.
- [41] S. Ströhle, A. Haselbacher, Z.R. Jovanovic, A. Steinfeld, The effect of the gas-solid contacting pattern in a high-temperature thermochemical energy storage on the performance of a concentrated solar power plant, *Energy Environ. Sci.* 9 (2016) 1375–1389, <https://doi.org/10.1039/c5ee03204k>.
- [42] T. Block, N. Knoblauch, M. Schmäcker, The cobalt-oxide/iron-oxide binary system for use as high temperature thermochemical energy storage material, *Thermochimica Acta* 577 (2014) 25–32, <https://doi.org/10.1016/j.tca.2013.11.025>.
- [43] C. Agrafiotis, M. Roeb, M. Schmäcker, C. Sattler, Exploitation of thermochemical cycles based on solid oxide redox systems for thermochemical storage of solar heat. Part 1: testing of cobalt oxide-based powders, *Solar Energy* 102 (2014) 189–211, <https://doi.org/10.1016/j.solener.2013.12.032>.
- [44] S. Tescari, C. Agrafiotis, S. Breuer, L. De Oliveira, M. Neises-Von Puttkamer, M. Roeb, C. Sattler, Thermochemical solar energy storage via redox oxides: materials and reactor/heat exchanger concepts, *Energy Procedia* 49 (2013) 1034–1043, <https://doi.org/10.1016/j.egypro.2014.03.111>.
- [45] G. Karagiannakis, C. Pagkoura, A. Zygogianni, S. Lorentzou, A.G. Konstandopoulos, Monolithic ceramic redox materials for thermochemical heat storage applications in CSP plants, *Energy Procedia* 49 (2014) 820–829, <https://doi.org/10.1016/j.egypro.2014.03.089>.
- [46] M. Neises, S. Tescari, L. de Oliveira, M. Roeb, C. Sattler, B. Wong, Solar-heated rotary kiln for thermochemical energy storage, *Solar Energy* 86 (2012) 3040–3048, <https://doi.org/10.1016/j.solener.2012.07.012>.
- [47] S.M. Babiniec, E.N. Coker, A. Ambrosini, J.E. Miller, ABO₃ (A = La, Ba, Sr, K; B = Co, Mn, Fe) perovskites for thermochemical energy storage, in: AIP Conference Proceedings 1734, 2016, <https://doi.org/10.1063/1.4949104>.
- [48] K.T. Jacob, C.B. Alcock, Thermodynamics of CuAlO₂ and CuAl₂O₄ and phase equilibria in the system Cu₂O-CuO-Al₂O₃, *J. Am. Ceram. Soc.* 58 (1974) 192–195.
- [49] C.K. Clayton, H.Y. Sohn, K.J. Whitty, Oxidation kinetics of Cu₂O in oxygen carriers for chemical looping with oxygen uncoupling, *Ind. Eng. Chem. Res.* 53 (2014) 2976–2986, <https://doi.org/10.1021/ie402495a>.
- [50] S.S. Jahromy, F. Birkelbach, C. Jordan, C. Huber, M. Harasek, A. Werner, F. Winter, Impact of partial pressure, conversion, and temperature on the oxidation reaction kinetics of Cu₂O to CuO in thermochemical energy storage, *Energies* (2019) 12, <https://doi.org/10.3390/en12030508>.
- [51] C. Prieto, P. Cooper, A.I. Fernández, L.F. Cabeza, Review of technology: thermochemical energy storage for concentrated solar power plants, *Renew. Sustain. Energy Rev.* 60 (2016) 909–929, <https://doi.org/10.1016/j.rser.2015.12.364>.
- [52] A.H. McDaniel, E.C. Miller, D. Arifin, A. Ambrosini, E.N. Coker, R. O'Hayre, W. C. Chueh, J. Tong, Sr- and Mn-doped LaAlO_{3-δ} for solar thermochemical H₂ and CO production, *Energy Environ. Sci.* 6 (2013) 2424–2428, <https://doi.org/10.1039/c3ee41372a>.
- [53] S.V. Ushakov, A. Navrotsky, Direct measurement of fusion enthalpy of LaAlO₃ and comparison of energetics of melt, glass, and amorphous thin films, *J. Am. Ceram. Soc.* 97 (2014) 1589–1594, <https://doi.org/10.1111/jace.12785>.
- [54] I. Barin, O. Knacke, *Thermochemical Properties of Inorganic Substances*, Springer, Berlin Heidelberg, 1973.
- [55] P. Gayán, C.R. Forero, A. Abad, L.F. De Diego, F. García-Labiano, J. Adánez, Effect of support on the behavior of Cu-based oxygen carriers during long-term CLC operation at temperatures above 1073 K, *Energy Fuels* 25 (2011) 1316–1326, <https://doi.org/10.1021/ef101583w>.
- [56] C.R. Forero, P. Gayán, F. García-Labiano, L.F. de Diego, A. Abad, J. Adánez, High temperature behaviour of a CuO/γAl₂O₃ oxygen carrier for chemical-looping combustion, *Inte. J. Greenh. Gas Control* 5 (2011) 659–667, <https://doi.org/10.1016/j.jggc.2011.03.005>.
- [57] M. Arjomand, A.-M. Azad, H. Leion, T. Mattisson, A. Lyngfelt, Evaluation of CuAl₂O₄ as an oxygen carrier in chemical-looping combustion, *Ind. Eng. Chem. Res.* (2012) 51, <https://doi.org/10.1021/ie3004027w>.
- [58] J. Adánez, A. Abad, F. García-Labiano, P. Gayán, L.F. De Diego, Progress in chemical-looping combustion and reforming technologies, *Prog. Energy Comb. Sci.* 38 (2012) 215–282, <https://doi.org/10.1016/j.pecs.2011.09.001>.
- [59] P. Gayán, I. Adánez-Rubio, A. Abad, L.F. De Diego, F. García-Labiano, J. Adánez, Development of Cu-based oxygen carriers for chemical-looping with oxygen uncoupling (CLOU) process, *Fuel* 96 (2012) 226–238, <https://doi.org/10.1016/j.fuel.2012.01.021>.
- [60] P. Moldenhauer, M. Rydén, T. Mattisson, A. Lyngfelt, Chemical-looping combustion and chemical-looping with oxygen uncoupling of kerosene with Mn-

- and Cu-based oxygen carriers in a circulating fluidized-bed 300 W laboratory reactor, *Fuel Process. Technol.* 104 (2012) 378–389, <https://doi.org/10.1016/j.fuproc.2012.06.013>.
- [61] I. Adánez-Rubio, P. Gayán, A. Abad, L.F. De Diego, F. García-Labiano, J. Adánez, Evaluation of a spray-dried CuO/MgAl₂O₄ oxygen carrier for the chemical looping with oxygen uncoupling process, *Energy Fuels* 26 (2012) 3069–3081, <https://doi.org/10.1021/ef3002229>.
- [62] A. Abad, I. Adánez-Rubio, P. Gayán, F. García-Labiano, L.F. de Diego, J. Adánez, Demonstration of chemical-looping with oxygen uncoupling (CLOU) process in a 1.5kWth continuously operating unit using a Cu-based oxygen-carrier, *Int. J. Greenh. Gas Control* 6 (2012) 189–200, <https://doi.org/10.1016/j.ijggc.2011.10.016>.
- [63] C.W. Bale, E. Bélisle, Fact-web suite of interactive programs, (n.d.). www.factsage.com.