Oxygen storage capacity and thermal stability of brownmillerite-type 
\(\text{Ca}_2(\text{Al}_{1-x}\text{Ga}_x)\text{MnO}_{5+\delta}\) oxides

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(Prof. John T.S. Irvine)
Abstract
Understanding the oxygen uptake/release mechanism in oxygen storage materials is of great importance in the design of energy-related materials and their corresponding applications. In this work, the effects of Ga doping amount on the oxygen storage capacity and thermal stability of Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_{5+\delta}$ (0 ≤ x ≤ 1) with a brownmillerite-type structure were investigated. Ca$_2$AlMnO$_{5+\delta}$ can reversibly store/release a large amount of excess oxygen (~3.0 wt%) at low temperature (between 300 and 600 °C) under oxidative atmospheres. With the increasing Ga doping amount in Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_{5+\delta}$, these materials uptake less oxygen at higher temperature which can be attributed to the difficulty in the oxidation of tetrahedral GaO$_4$ blocks into octahedral GaO$_6$ blocks under 1 atm O$_2$. However, with the increasing of Ga-substitution amount, these Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_{5+\delta}$ (0 ≤ x < 1) can start to uptake oxygen at lower temperatures during the cooling process under flowing O$_2$ due to the distorted structure. The results demonstrated that Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_{5+\delta}$ (0 ≤ x < 1) can reversibly store/release large amounts of oxygen via just controlling the surrounding temperature and/or oxygen partial pressure but without using reductive gases, which would enable them great potentials in many applications.

Keywords: Oxygen storage materials; brownmillerite-type structure; Doping effect; Reversible behaviour
1. Introduction

Metal oxides based on valence-changeable constituent cations have the ability to exhibit oxygen nonstoichiometry, which is often related to oxygen uptake/release capacity.[1-4] Those oxides with remarkable oxygen uptake/release capability can be regarded as oxygen storage materials (OSMs), but excellent OSMs should also possess some other properties, such as quick and reversible adsorption/desorption of oxygen controlled by oxygen partial pressure (PO$_2$) and/or temperature.[5, 6] One of the most important applications of OSMs is used as three-way catalysts (TWC) for the contaminant removal in automobile exhausts,[7, 8] in which the OSM helps to maintain appropriate oxygen partial pressure so that both oxidation reaction (of CO, H$_2$, and carbohydrates) and reduction reaction (of NOx) can occur effectively. The OSMs incorporate oxygen in higher pO$_2$ and release oxygen in lower pO$_2$, therefore for TWC, materials that can work in PSA mode (pressure swing absorption) are recommended.[7, 9, 10] For some other specific applications (e.g., Li-oxygen batteries), pure fresh oxygen gas released from oxides may be safer than that from the pressurized oxygen tank. Even though some peroxides, such as CaO$_2$, MgO$_2$, can provide fresh oxygen gas with the increase of temperature under oxidative gas, their irreversibility limits their wide applications.[11, 12] Therefore, it is important to develop special reversible OSMs which can release fresh oxygen gas with the changes of temperature under inert or oxidative atmosphere.

As a well-known structure, perovskite oxides have been recognized as one of the most important types of OSMs with excellent redox properties, high oxygen ion mobility and highly stable structure with an unusual valence state of elements or a high extent of oxygen deficiency.[13-20] Especially, those perovskite oxides based on Mn are potential OSMs because of the easy conversion between Mn$^{3+}$ and Mn$^{4+}$.[21-
Recently, YMnO$_{3+\delta}$ has been reported to have large and reversible oxygen content at a low temperature range under oxidative atmosphere because of the easy phase transition between hexagonal $P6_3cm$ phase and more oxidized structures (e.g., $Pca2_1$, $R3c$).[24]

Brownmillerite-type perovskite oxides with a general formula of $A_2B_2O_5$ can be viewed as the anion-deficient perovskite with alternately stacked tetrahedral BO$_4$ and octahedral BO$_6$ layers.[22, 25, 26] They can uptake oxygen to form perovskite oxides $ABO_{3+\delta}$, thus having the potential applications as OSMs. In addition, the reversible phase change between brownmillerite-type perovskite with $A_2B_2O_5$ and perovskite with $ABO_{3+\delta}$ could improve the reversibility oxygen release/storage. Especially, perovskite-type oxides based on Mn have attracted remarkable attention and have been widely researched as OSMs.[27-34] For example, BaYMn$_2$O$_{5+\delta}$ exhibits remarkable oxygen storage capacity (OSC, 3.7 wt\%) at low temperatures below 500 °C with an excellent reversible behaviour in alternative reductive ($H_2$) and oxidative ($O_2$) conditions, but it is not suitable in pure oxidative or alternative oxidative and inert gas conditions.[5, 22, 30] Thus, excellent OSMs based on brownmillerite-type perovskite oxides still need more research. Recently, $Ca_2AlMnO_{5+\delta}$ was reported to have a high OSC (~3 wt\%) in response to changes in surrounding temperature and pressure of $O_2$ in a highly reversible behaviour, accompanied with the valence change between Mn$^{3+}$ and Mn$^{4+}$.[27, 34-36] Another similar brownmillerite-type oxide, $Ca_2GaMnO_5$ has been investigated about its magnetic structure and oxygen content under different preparation conditions.[37, 38] Nevertheless, the detailed understanding on the phase structure and oxygen content in the solid solutions of $Ca_2(Al_{1-x}Ga_x)MnO_{5+\delta}$ is still not clear. Herein, we prepared some brownmillerite-type oxides based on $Ca_2(Al_{1-x}Ga_x)MnO_{5+\delta}$ with various substituting amounts (0 ≤ x ≤ 1),
and investigated their oxygen storage/release properties and thermal stability against temperature and the surrounding PO$_2$ with the purpose to better understand their oxygen storage/release mechanism.

2. Experimental Section

2.1 Synthesis

Brownmillerite-type oxides with the formula of Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$ ($x = 0, 0.25, 0.5, 0.75, 1$) were prepared via solid state method using CaCO$_3$, Al$_2$O$_3$, MnO$_2$ and Ga$_2$O$_3$ as precursors. Stoichiometric amounts of precursors were homogeneously mixed by grounding in a mortar. The obtained powders were pressed into cylindrical pellets, which were further fired in a tube furnace at 1250 °C under static air for 24 h and under flowing pure argon for another 12 h. Then these obtained pellets were ground into fine powder for XRD characterization and OSC test.

2.2 Characterization

Powder X-ray diffraction (XRD) patterns were obtained on a PANalytical Empyrean Reflection Diffractometer using Cu Ka radiation ($\lambda = 1.541$ Å). The oxygen uptake/release ability of these samples was tested by thermogravimetric analysis (TGA) on a NETZSCH TG 209 instrument (NETZSCH-Geraetebau GmbH, Selb, Germany) with a TASC 414/3 controller. Oxygen uptake property was obtained up to 900 °C with heating/cooling rates of ±1 °C/min.

3. Results and Discussion

Fig. 1 shows the XRD patterns of as-prepared Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$ (i.e., post-annealing at 1250 °C under flowing argon gas for 12 h). The XRD pattern for as-prepared Ca$_2$AlMnO$_5$ in Fig. 1a can be ascribed to orthorhombic structure with $Ibm2$ space group, similar to reported results.[39] The lattice parameters for as-prepared
Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$ after Rietveld refinement were summarized in Table 1. The lattice parameters of our as-prepared Ca$_2$AlMnO$_5$ are determined to be $a = 5.4705(1)$, $b = 15.0050(3)$, and $c = 5.2431(1)$ Å.

![XRD patterns](image)

**Fig. 1.** XRD patterns of as-prepared Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$ (i.e., post-annealing at 1250 °C under flowing argon for 12 h): (a) Ca$_2$AlMnO$_5$, (b) Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_5$, (c) Ca$_2$(Al$_{0.5}$Ga$_{0.5}$)MnO$_5$, (d) Ca$_2$(Al$_{0.25}$Ga$_{0.75}$)MnO$_5$, (e) Ca$_2$GaMnO$_5$.

The XRD patterns for the as-prepared Ga-substituted samples after post-annealing at 1250 °C under Ar for 12 h can be indexed to orthorhombic structure with $Pnma$ space group.[40] The crystal structures of brownmillerite-type Ca$_2$AlMnO$_5$ with $Ibm2$ space group and Ca$_2$GaMnO$_5$ with $Pnma$ space group are schematically shown in Fig. 2a and Fig. 2b via Diamond software,[41] respectively, based on the structure reported in literature.[37, 42, 43] As shown in Fig. 2a, Ca$_2$AlMnO$_5$ comprises tetrahedral AlO$_4$ and octahedral MnO$_6$ blocks, where the MnO$_6$ chain orientations are fully ordered.[39] As shown in Fig. 2b, the chain orientations of MnO$_6$ in Ca$_2$GaMnO$_5$ are disordered, where the R-type (right-hand chains) and L-type (left-hand chains) Ga-O chains are ordered with -L-R-L-R- alternation along the $b$ axis.[37] The main
difference between the structures of $\text{Ca}_2\text{AlMnO}_5$ and $\text{Ca}_2\text{GaMnO}_5$ is the ordering of tetrahedron and octahedron in the orthorhombic structure, resulting in different space groups.

Fig. 2. Schematic illustration of the crystal structures with single cell of (a) brownmillerite-type $\text{Ca}_2\text{AlMnO}_5$, (b) brownmillerite-type $\text{Ca}_2\text{GaMnO}_5$, (c) the fully oxygenated form $\text{Ca}_2\text{AlMnO}_{5.5}$. The illustration was obtained from Inorganic Crystal Structure Database (ICSD) web and prepared via Diamond Software [41, 44] based on the structure reported in literature.[37, 42, 43]

The lattice parameters for the as-prepared $\text{Ca}_2(\text{Al}_{1-x}\text{Ga}_x)\text{MnO}_5$ in Table 1 indicate that with the increasing substitution of Ga amount ($0 < x \leq 1$), the $b$ and $c$ lattice parameters increased obviously and the cell volume also slightly increased, which can be attributed to the increased thickness of the $(\text{Al},\text{Ga})\text{O}_4$ layer due to the larger ionic radius of $\text{Ga}^{3+}$ (0.62 Å) than that of $\text{Al}^{3+}$ (0.535 Å). However, there is some slight decrease in the $a$ lattice parameters with the increasing of Ga amount in the brownmillerite-type $\text{Ca}_2(\text{Al}_{1-x}\text{Ga}_x)\text{MnO}_5$ ($0 < x \leq 1.0$), which can be attributed to the tilting of the oxygen position in the MnO$_6$ octahedron blocks.[45, 46]
Table 1. Lattice parameters of as-prepared Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$ and oxygenated Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_{5+\delta}$.

<table>
<thead>
<tr>
<th>Samples</th>
<th>Space group</th>
<th>$a$ (Å)</th>
<th>$b$ (Å)*</th>
<th>$c$ (Å)</th>
<th>Cell volume ($\text{Å}^3$)*</th>
<th>Rwp</th>
<th>Rp</th>
<th>$\chi^2$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ca$_2$AlMnO$_5$</td>
<td>Ibm2</td>
<td>5.4705(1)</td>
<td>15.0050(3)</td>
<td>5.2431(1)</td>
<td>430.38(2)</td>
<td>10.38%</td>
<td>7.02%</td>
<td>4.99</td>
</tr>
<tr>
<td>Ca$<em>2$AlMnO$</em>{5+\delta}$</td>
<td>Iemma</td>
<td>5.3732(4)</td>
<td>14.7160(2)</td>
<td>5.2534(4)</td>
<td>415.45(1)</td>
<td>16.73%</td>
<td>11.50%</td>
<td>5.96</td>
</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.75}$Ga$_{0.25}$)MnO$_5$</td>
<td>Pnma</td>
<td>5.4698(2)</td>
<td>15.0802(6)</td>
<td>5.2488(4)</td>
<td>432.95(4)</td>
<td>16.26%</td>
<td>11.77%</td>
<td>6.11</td>
</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.75}$Ga$<em>{0.25}$)MnO$</em>{5+\delta}$</td>
<td>Iemma</td>
<td>5.3892(2)</td>
<td>14.7810(1)</td>
<td>5.2669(2)</td>
<td>419.56(7)</td>
<td>11.34%</td>
<td>8.28%</td>
<td>2.85</td>
</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.5}$Ga$_{0.5}$)MnO$_5$</td>
<td>Pnma</td>
<td>5.4700(2)</td>
<td>15.1643(6)</td>
<td>5.2561(2)</td>
<td>435.99(4)</td>
<td>11.50%</td>
<td>11.76%</td>
<td>5.98</td>
</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.5}$Ga$<em>{0.5}$)MnO$</em>{5+\delta}$</td>
<td>Iemma</td>
<td>5.4072(2)</td>
<td>14.8300(1)</td>
<td>5.2793(2)</td>
<td>423.35(1)</td>
<td>13.18%</td>
<td>10.38%</td>
<td>3.66</td>
</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.25}$Ga$_{0.75}$)MnO$_5$</td>
<td>Pnma</td>
<td>5.4617(2)</td>
<td>15.3012(6)</td>
<td>5.2629(2)</td>
<td>439.82(4)</td>
<td>16.31%</td>
<td>11.89%</td>
<td>5.87</td>
</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.25}$Ga$<em>{0.75}$)MnO$</em>{5+\delta}$</td>
<td>Pnma</td>
<td>5.4630(1)</td>
<td>15.1820(2)</td>
<td>5.2634(6)</td>
<td>436.60(1)</td>
<td>11.31%</td>
<td>9.15%</td>
<td>3.12</td>
</tr>
<tr>
<td></td>
<td>Iemma</td>
<td>5.4204(4)</td>
<td>14.9110(2)</td>
<td>5.2955(4)</td>
<td>428.00(1)</td>
<td>13.12%</td>
<td>10.21%</td>
<td>4.23</td>
</tr>
<tr>
<td>Ca$_2$GaMnO$_5$</td>
<td>Pnma</td>
<td>5.4611(1)</td>
<td>15.3303(3)</td>
<td>5.2664(1)</td>
<td>440.92(2)</td>
<td>15.05%</td>
<td>11.41%</td>
<td>5.00</td>
</tr>
<tr>
<td>Ca$<em>2$GaMnO$</em>{5+\delta}$</td>
<td>Pnma</td>
<td>5.4629(1)</td>
<td>15.3023(3)</td>
<td>5.2707(1)</td>
<td>440.61(2)</td>
<td>13.70%</td>
<td>10.27%</td>
<td>4.31</td>
</tr>
</tbody>
</table>

* b parameter and cell volume of Iemma structure was divided by 2 to compare with those in Ibm2 and Pnma.

Based on the TGA test results under O$_2$ atmosphere (Fig. 4), all the samples obtained the highest weight at around 450 °C. Therefore, these as-prepared Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$ samples were treated in air atmosphere at 450 °C for 12 h in a furnace for oxygenation in order to check the phase structure changes of oxygenated samples. The XRD patterns of oxygenated samples are shown in Fig. 3 and their lattice parameters were also summarized in Table 1. The XRD patterns of oxygenated Ca$_2$AlMnO$_{5+\delta}$ (Fig. 3a) can be ascribed to the orthorhombic structure with Iemma space group ($a = 5.3732(4)$, $b = 29.4320(2)$ and $c = 5.2534(4)$ Å), similar to reported...
results. The crystal structure of the full oxygenated Ca$_2$AlMnO$_{5.5}$ is shown in Fig. 2c based on the structure reported in literature. After oxygenation, the fully oxygenated Ca$_2$AlMnO$_{5.5}$ was composed of alternating tetrahedral AlO$_4$ and octahedral AlO$_6$ blocks, which are separated by octahedral MnO$_6$. The structure differences between the pristine Ca$_2$AlMnO$_5$ and the oxygenated Ca$_2$AlMnO$_{5.5}$ imply that the oxygen uptake process mainly occur in the $b$-lattice direction with the partial oxidation of the tetrahedral AlO$_4$ to octahedral AlO$_6$, but still half of the AlO$_4$ tetrahedron are intact in the oxygenated Ca$_2$AlMnO$_{5.5}$, which has been considered to be linked with its reversible oxygen uptake/release process.

The XRD patterns of oxygenated Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_{5+δ}$ and Ca$_2$(Al$_{0.5}$Ga$_{0.5}$)MnO$_{5+δ}$ can also be ascribed to the orthorhombic structure with Imma space group. In contrast, the XRD patterns of oxygenated Ca$_2$(Al$_{0.25}$Ga$_{0.75}$)MnO$_{5+δ}$ can be indexed to intermixed orthorhombic Pnma/Imma space group, indicating the formation of octahedral AlO$_6$ blocks from tetrahedral AlO$_4$ blocks during the oxidation process while the GaO$_4$ blocks are still intact. Comparing the lattice parameters of the pristine materials (i.e., Ca$_2$AlMnO$_5$, Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_5$, Ca$_2$(Al$_{0.5}$Ga$_{0.5}$)MnO$_5$ and Ca$_2$(Al$_{0.25}$Ga$_{0.75}$)MnO$_5$) and their oxygenated counterparts (i.e., Ca$_2$AlMnO$_{5+δ}$, Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_{5+δ}$, Ca$_2$(Al$_{0.5}$Ga$_{0.5}$)MnO$_{5+δ}$ and Ca$_2$(Al$_{0.25}$Ga$_{0.75}$)MnO$_{5+δ}$), there is a large increase (a bit smaller than double) in the $b$ lattice parameter, indicating the oxygenation reaction takes place topotactically with the formation of octahedral AlO$_6$ from tetrahedral AlO$_4$. With the increase of Ga-substituting amount ($0 \leq x \leq 0.75$) in the oxygenated Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_{5+δ}$, the lattice parameters ($a$, $b$ and $c$) and cell volumes increased due to the larger ionic radius of Ga$^{3+}$ (0.62 Å) than that of Al$^{3+}$ (0.535 Å). When completely substituting Al with Ga (i.e., $x = 1.0$) in Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_{5+δ}$, the oxygenated Ca$_2$GaMnO$_{5+δ}$ (Fig. 3e) still maintains the same orthorhombic Pnma space group as
its pristine Ca$_2$GaMnO$_5$. Comparing the lattice parameters of the pristine samples (i.e., Ca$_2$(Al$_{0.25}$Ga$_{0.75}$)MnO$_5$ and Ca$_2$GaMnO$_5$) and their oxygenated counterparts (i.e., Ca$_2$(Al$_{0.25}$Ga$_{0.75}$)MnO$_{5+\delta}$ and Ca$_2$GaMnO$_{5+\delta}$) with Pnma space group, there are slight increases in $a$ and $c$ lattice parameter and obvious decrease in $b$ lattice parameters, which can be attributed to the slightly compressed oxygen tetrahedra around the Ga atoms along the $b$ axis during the oxidation process in air atmosphere at 450 °C for 12 h.[37, 39] However, it has been reported that oxygenated Ca$_2$GaMnO$_{5.39}$ can be obtained by treating Ca$_2$GaMnO$_5$ under the oxygen partial pressure of 80 atm O$_2$ at 415 °C for 15 h.[37] The XRD results in the pristine materials and their oxygenated counterparts indicate that Ga-substitution in Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$ would affect their oxygen uptake ability, which will be further investigated by TGA technique in following paragraphs.

![XRD patterns of oxygenated samples](image)

**Fig. 3.** XRD patterns of oxygenated samples after treated at 450 °C under static air for 12 h: (a) Ca$_2$AlMnO$_{5+\delta}$, (b) Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_{5+\delta}$, (c) Ca$_2$(Al$_{0.5}$Ga$_{0.5}$)MnO$_{5+\delta}$, (d) Ca$_2$(Al$_{0.25}$Ga$_{0.75}$)MnO$_{5+\delta}$, (e) Ca$_2$GaMnO$_{5+\delta}$. 


Fig. 4 shows the oxygen uptake/release properties of as-prepared samples under pure O\textsubscript{2} with 1 °C/min heating and cooling rate between room temperature and 900 °C. The results show that upon heating in O\textsubscript{2} atmospheres, these samples firstly remain almost stable between room temperature and 300 °C, then sharply gain weight from 300 °C to reach the maximum peaks at about 370-470 °C, which are different according to different Ga substituting amounts. The maximum weight gain (ΔW) and its corresponding temperature, OSC, oxygen content and the oxygenated samples were summarized in Table 2, in which the Mn valence in the as-prepared samples is assumed to be Mn\textsuperscript{3+}. Ca\textsubscript{2}(Al\textsubscript{0.75}Ga\textsubscript{0.25})MnO\textsubscript{5} shows the maximum weight at 450 °C with a 3.16 wt% weight gain (Fig. 4b), corresponding to the OSC of 1975 μmol-O·g\textsuperscript{-1}, while Ca\textsubscript{2}AlMnO\textsubscript{5} in Fig. 4a shows the second recorded highest OSC at 370 °C with a 2.97 wt% weight gain, corresponding to the OSC of 1856 μmol-O·g\textsuperscript{-1}, indicating substituting Al with 0.25 atomic Ga would improve the oxygen uptake ability maybe due to the enhanced oxygen mobility by the increased lattice parameters. It is noteworthy that even though Ca\textsubscript{2}(Al\textsubscript{0.75}Ga\textsubscript{0.25})MnO\textsubscript{5} showed the highest weight increase during the initial heating process from room temperature to 400 °C, it can’t lose weight to its original value at room temperature during the consequent heating process from 550 °C to 900 °C, which can be attributed to the difficulty in the transformation of octahedral GaO\textsubscript{6} to GaO\textsubscript{4} tetrahedron. Further increasing the Ga substituting amount to x ≥ 0.50 on the Al-site would result in reduced OSC, which is maybe due to the difficulty in the oxidation of GaO\textsubscript{4} tetrahedron in brownmillerite-type structure to octahedral GaO\textsubscript{6} in air atmosphere at 450 °C, as well as the higher atomic weight of Ga than Al.[39] As a result, Ca\textsubscript{2}GaMnO\textsubscript{5} in Fig. 4e can only uptake about 0.62 wt% weight gain, corresponding to the OSC of 388 μmol-O·g\textsuperscript{-1}. It has been reported that the electron mobility to the MnO\textsubscript{6} layers is weakened because of
the fully filled electron orbitals of Ga$^{3+}$(d$^{10}$) ions.[47] As a result, the super-exchange interactions between the GaO$_4$ and MnO$_6$ layers in the brownmillerite structure are suppressed perpendicular to the octahedral MnO$_6$ layers.[47]

**Fig. 4.** TGA curves of as-prepared samples (i.e. post-annealing Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$ at 1250 °C under Ar for 12 h): (a) Ca$_2$AlMnO$_5$, (b) Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_5$, (c) Ca$_2$(Al$_{0.5}$Ga$_{0.5}$)MnO$_5$, (d) Ca$_2$(Al$_{0.25}$Ga$_{0.75}$)MnO$_5$, (e) Ca$_2$GaMnO$_5$. The data were obtained under flowing O$_2$ with a scan rate of 1 °C/min between 20 and 900 °C.

**Table 2.** The maximum weight increase ($\Delta$W), the temperature, OSC and oxygen content obtained from the oxygen uptake/release curves in Fig. 4, assuming the valence of Mn in the as-prepared samples is Mn$^{3+}$.

<table>
<thead>
<tr>
<th>As-prepared sample</th>
<th>Maximum $\Delta$W</th>
<th>Temperature ($°$C)</th>
<th>OSC (µmol-O·g$^{-1}$)</th>
<th>Oxygen content $\delta$</th>
<th>Oxygenated sample</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ca$_2$AlMnO$_5$</td>
<td>2.97 wt%</td>
<td>392</td>
<td>1856</td>
<td>0.45</td>
<td>Ca$_2$AlMnO$_5.45$</td>
</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.75}$Ga$_{0.25}$)MnO$_5$</td>
<td>3.16 wt%</td>
<td>450</td>
<td>1975</td>
<td>0.5</td>
<td>Ca$<em>2$(Al$</em>{0.75}$Ga$_{0.25}$)MnO$_5.5$</td>
</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.5}$Ga$_{0.5}$)MnO$_5$</td>
<td>2.62 wt%</td>
<td>448</td>
<td>1637</td>
<td>0.43</td>
<td>Ca$<em>2$(Al$</em>{0.5}$Ga$_{0.5}$)MnO$_5.43$</td>
</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.25}$Ga$_{0.75}$)MnO$_5$</td>
<td>2.13 wt%</td>
<td>452</td>
<td>1331</td>
<td>0.36</td>
<td>Ca$<em>2$(Al$</em>{0.25}$Ga$_{0.75}$)MnO$_5.36$</td>
</tr>
<tr>
<td>Ca$_2$GaMnO$_5$</td>
<td>0.62 wt%</td>
<td>454</td>
<td>388</td>
<td>0.11</td>
<td>Ca$_2$GaMnO$_5.11$</td>
</tr>
</tbody>
</table>

Further increasing temperature would result in an abrupt weight loss at around 650 °C and plateau between 700 and 900 °C, and Ca$_2$AlMnO$_5$ shows the least weight difference with its initial value. Interestingly, during the subsequent cooling stage in flowing O$_2$, these samples could re-gain weight from ca. 600 °C and keep almost stable below 400 °C. In these five samples, Ca$_2$AlMnO$_5$ can almost re-gain the same weight (ca. 2.73 wt% corresponding to $\delta = 0.41$ and OSC of 1688 $\mu$mol-O·g$^{-1}$) as the weight lost during the heating process. However, for the Ga-substituted samples, they can't re-gain the same weight that is lost during the heating processes. With the increasing Ga-substituting amount, the re-gained weight during the cooling process decreased, which can be attributed to the suppressed oxygen-ion mobility and desorption by the increased intact GaO$_4$ layers in the structure. For Ca$_2$GaMnO$_5$ shown in Fig. 4c, there is almost no weight uptake during the cooling process, which is in good agreement with its XRD results. In addition, the starting temperature for oxygen uptake during the cooling process decreased from ca. 600 °C for Ca$_2$AlMnO$_5$ to 470 °C for Ca$_2$(Al$_{0.25}$Ga$_{0.75}$)MnO$_5$, which may be due to the different oxygen-ion mobility and distorted structure between (Al/Ga)O$_6$ and MnO$_6$ caused by the increased b-lattice parameter with the increasing of Ga-substituting amount. The thermal hysteresis and the differences in the TGA curves during the heating/cooling processes may be caused by their different oxygen uptake/release kinetics which is related with their composition.[27] Fig. 4 shows that the oxygen uptake/release in Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_{5+\delta}$ can be achieved under pure O$_2$ by adjusting temperature in a small range, which can be attributed to the redox equilibria of competing valence states (such as Mn$^{3+}$/Mn$^{4+}$) in the constituent transition metals,[27] similar to reported YBaCo$_4$O$_{7+\delta}$[48-50] and Dy$_{1-x}$Y$_x$MnO$_{3+\delta}$.[51]

Fig. 5 shows the oxygen uptake ability and OSCs of the as-prepared Ca$_2$(Al$_{1-
\( x \text{Ga}_x \text{MnO}_5 \) oxides at 450 °C under flowing O\(_2\) with a 25 mL/min flowing rate. These as-prepared samples were firstly treated from room temperature to 450 °C and maintained at 450 °C for 1 h under flowing argon with 25 mL/min to remove the possible adsorbed surface oxygen species. Then, the gas was switched from argon to O\(_2\) for the oxygen uptake test dependent on reaction time at 450 °C. For Ca\(_2\)AlMnO\(_5\) and Ca\(_2\)(Al\(_{0.75}\)Ga\(_{0.25}\))MnO\(_5\), they uptake oxygen immediately to the maximum amount (ca. 2.73 wt% corresponding to \( \delta = 0.41 \) and OSC of 1688 \( \mu \text{mol-O·g}^{-1} \) for Ca\(_2\)AlMnO\(_{5.41}\), and 2.41 wt% corresponding to \( \delta = 0.38 \) and OSC of 1500 \( \mu \text{mol-O·g}^{-1} \) for Ca\(_2\)(Al\(_{0.75}\)Ga\(_{0.25}\))MnO\(_{5.38}\)) in 10 min. However, further increasing the Ga-substitution amount to \( x \geq 0.5 \) in Ca\(_2\)(Al\(_{1-x}\)Ga\(_x\))MnO\(_5\) would result in less oxygen uptake in a longer time due to the lowered ionic diffusion related to the Al-O/Ga-O chemical bonds. For example, Ca\(_2\)(Al\(_{0.5}\)Ga\(_{0.5}\))MnO\(_5\) would uptake oxygen to the maximum amount (ca. 2.1 wt% corresponding to \( \delta = 0.34 \) and OSC of 1313 \( \mu \text{mol-O·g}^{-1} \) for Ca\(_2\)(Al\(_{0.5}\)Ga\(_{0.5}\))MnO\(_{5.34}\)) in about 50 min while Ca\(_2\)(Al\(_{0.25}\)Ga\(_{0.75}\))MnO\(_5\) needs even more time (ca. 220 min) to reach the maximum oxygen uptake amount (ca. 1.6 wt% corresponding to \( \delta = 0.27 \) and OSC of 1000 \( \mu \text{mol-O·g}^{-1} \) for Ca\(_2\)(Al\(_{0.25}\)Ga\(_{0.75}\))MnO\(_{5.27}\)). Ca\(_2\)GaMnO\(_5\) can uptake only a very small amount of oxygen (ca. 0.11 wt% corresponding to \( \delta = 0.02 \) and OSC of 13 \( \mu \text{mol-O·g}^{-1} \) for Ca\(_2\)GaMnO\(_{5.02}\)) with the increasing time to 220 min, in good agreement with their XRD patterns and lattice parameter analysis results.
Fig. 5. Isothermal TGA curves at 450 °C for as-prepared Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$ with various x values after switching the gas flow with 25 mL/min from Ar to O$_2$.

As shown by the results in Fig. 4, these as-prepared samples, especially for Ca$_2$AlMnO$_5$ and Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_5$, could change their oxygen content in a narrow temperature range (500-700 °C) under flowing O$_2$ atmosphere. Thus, it can be expected that remarkable oxygen uptake/release capacities can be obtained via adjusting the temperature in a small range even under pure O$_2$. As demonstrated in Fig. 6, a large amount of oxygen can be reversibly stored/released for Ca$_2$AlMnO$_5$ and Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_5$ by adjusting the temperature between 500 and 700 °C. Ca$_2$AlMnO$_5$ shows a reversible and stable weight change with 2.4 wt%, larger than that of Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_5$ with 1.9 wt%, and the results are consistent with the TGA results in Fig. 4.
Fig. 6. TGA curves for Ca$_2$AlMnO$_5$ (a) and Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_5$ (b) in flowing O$_2$ upon temperature swing between 700 and 500 °C. The sample temperature is also shown with a blue curve.

Fig. 7 shows the oxygen uptake/release cycles of as-prepared Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$ oxides at 500 °C under alternating Ar and O$_2$ for three cycles. The results indicate that all these five samples exhibit excellent reversibility in oxygen uptake/release via switching the gas flow between argon and O$_2$ at 500 °C. With the increase of Ga substituting amount from $x = 0$ to $x = 1.0$ in Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$, the total oxygen uptake/release content decreased from about 2.4 wt% for Ca$_2$AlMnO$_5$ to 0.05 wt% for Ca$_2$GaMnO$_5$. Even though the OSCs of these as-prepared samples are different, they all have good reversibility for three cycling times under inert (Ar) and oxidative gas (O$_2$) at 500 °C, which is important for long-term usage in practical applications.
Fig. 7. TGA curves of as-prepared samples (i.e., post-annealing Ca(Al_{1-x}Ga_x)MnO_5 at 1250 °C under Ar for 12 h) obtained at 500 °C under alternating Ar and O_2 for three cycles: (a) Ca_2AlMnO_5, (b) Ca_2(Al_{0.75}Ga_{0.25})MnO_5, (c) Ca_2(Al_{0.5}Ga_{0.5})MnO_5, (d) Ca_2(Al_{0.25}Ga_{0.75})MnO_5, (e) Ca_2GaMnO_5.

The thermal stability of the oxygenated samples under pure argon with temperature is shown in Fig. 8. The results show that these samples are almost stable between room temperature and 400 °C under flowing argon. Further increasing temperature would result in sharp weight loss from 400 to 600 °C, and then keep stable between 600 and 900 °C. With the increasing amount of Ga, the weight loss amount between 400 and 650 °C decreased from about 2.9 wt% for Ca_2AlMnO_{5+δ} to 0.25 wt% for Ca_2GaMnO_{5+δ}. The results also indicate that substituting Ga into the Al site would reduce the OSC under flowing argon, which is in good agreement with the TGA cycle test results shown in Fig. 7. As discussed in the aforementioned paragraphs, in the Ga-substituting samples, Ga would occupy the Al-site of tetrahedral AlO_4 to form GaO_4 blocks. During the oxygenation process, tetrahedral AlO_4 would transfer to octahedral AlO_6 while tetrahedral GaO_4 is very difficult to be oxygenated to octahedral GaO_6 under 1 atm O_2 conditions, therefore Ga-substituted samples would result in less OSC.
Fig. 8. TGA curves under flowing Ar of as-prepared samples after treating at 450 °C in air for 12 h: (a) Ca$_2$AlMnO$_{5+δ}$, (b) Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_{5+δ}$, (c) Ca$_2$(Al$_{0.5}$Ga$_{0.5}$)MnO$_{5+δ}$, (d) Ca$_2$(Al$_{0.25}$Ga$_{0.75}$)MnO$_{5+δ}$, (e) Ca$_2$GaMnO$_{5+δ}$. The data were measured up to 900 °C under flowing pure Ar with 10 °C/min increasing rate.

Based on the TGA results, the Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_5$ could uptake as much as 1975 µmol-O·g$^{-1}$ during the heating process under flowing O$_2$ even though its OSC in reversible test is a bit lower than that of Ca$_2$AlMnO$_5$. In addition, Ca$_2$(Al$_{1-x}$Ga$_x$)MnO$_5$ (x < 1) can store/release oxygen reversibly via controlling the temperature or alternating Ar and O$_2$. The OSCs of Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_5$ and Ca$_2$AlMnO$_5$ are higher or comparable with the OSCs in reported results, as show in Table 3. Even though the OSCs of YBaCo$_4$O$_7$ and BaYMn$_2$O$_5$ are higher than those of Ca$_2$(Al$_{0.75}$Ga$_{0.25}$)MnO$_5$ and Ca$_2$AlMnO$_5$ in this work, the much lower abundance and higher price of Y than those of Ca and Al would increase the cost and limit their wide applications.

Table 3. Comparison of the OSC values in this work with the reported results.

<table>
<thead>
<tr>
<th>OSMs</th>
<th>OSC ($\mu$mol-O·g$^{-1}$)</th>
<th>Temperature (°C)</th>
<th>Gas atmosphere</th>
<th>Rate (°C·min$^{-1}$)</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ca$_2$AlMnO$_5$</td>
<td></td>
<td>450</td>
<td>Ar</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.75}$Ga$_{0.25}$)MnO$_5$</td>
<td>1975</td>
<td>900</td>
<td>Pure Ar</td>
<td>10</td>
<td></td>
</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.5}$Ga$_{0.5}$)MnO$_5$</td>
<td></td>
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</tr>
<tr>
<td>Ca$<em>2$(Al$</em>{0.25}$Ga$_{0.75}$)MnO$_5$</td>
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</tr>
<tr>
<td>Ca$_2$GaMnO$_5$</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
This work

\[
\text{Ca}_2\text{(Al}_{1-x}\text{Ga}_x)\text{MnO}_5 \quad 1975 \quad 450 \quad \text{O}_2 \quad 1.0 \quad \text{This work}
\]

\[
\text{Ca}_2\text{AlMnO}_5 \quad 1856 \quad 392 \quad \text{O}_2 \quad 1.0 \quad \text{This work}
\]

\[
\text{Y}_{0.7}\text{Tb}_{0.3}\text{MnO}_{3+\delta} \quad \sim 770 \quad \sim 320 \quad \text{O}_2 \quad 1.0 \quad [23]
\]

\[
\text{DyMnO}_3 \quad \sim 810 \quad \sim 400 \quad \text{O}_2 \quad 1.0 \quad [51]
\]

\[
\text{YBaMn}_2\text{O}_{5+\delta} \quad \sim 2240 \quad \sim 400 \quad \text{H}_2^* \quad 1.0 \quad [52]
\]

\[
\text{YBaCo}_4\text{O}_{7+\delta} \quad \sim 2180 \quad \sim 400 \quad \text{O}_2 \quad 1.0 \quad [53]
\]

\[
\text{Ce}_{1-x}\text{Zr}_x\text{O}_{2+\delta} \quad 360-750 \quad \sim 500 \quad \text{H}_2^* \quad - \quad [54]
\]

\*H\(_2\) is used for reduction to calculate the OSC.

4. Conclusions

Several pure-phased brownmillerite-type oxides with the formula of \(\text{Ca}_2\text{(Al}_{1-x}\text{Ga}_x)\text{MnO}_5\) were prepared via a solid state method. The substitution of Ga on the Al-site would result in the change of space group from \(\text{Ibm}2\) for \(\text{Ca}_2\text{AlMnO}_5\) to \(\text{Pnma}\) for \(\text{Ca}_2\text{(Al}_{1-x}\text{Ga}_x)\text{MnO}_5\) \((x > 0)\). Substituting Al with 0.25 atomic Ga (i.e., \(\text{Ca}_2\text{(Al}_{0.75}\text{Ga}_{0.25})\text{MnO}_5\)) would result in improved oxygen uptake content from 1856 \(\mu\text{mol-O}\cdot\text{g}^{-1}\) for \(\text{Ca}_2\text{AlMnO}_5\) to 1975 \(\mu\text{mol-O}\cdot\text{g}^{-1}\) for \(\text{Ca}_2\text{(Al}_{0.75}\text{Ga}_{0.25})\text{MnO}_5\) during the heating process under flowing \(\text{O}_2\) but less oxygen release amount during the cooling process. The \(\text{Ca}_2\text{(Al}_{1-x}\text{Ga}_x)\text{MnO}_5\) \((x < 1)\) can store/release oxygen reversibly depending on temperature between 500 and 700 °C, or alternating \(\text{Ar}\) and \(\text{O}_2\) at 500 °C, in which \(\text{Ca}_2\text{AlMnO}_5\) shows the highest OSC of about 2.9 wt% depending on temperature, and 2.4 wt% depending on alternative \(\text{Ar}\) and \(\text{O}_2\) at 500 °C, as well as excellent reversibility. With the increasing of Ga-substituting amount on the Al-site, their OSC decreased to only 0.11 wt% for \(\text{Ca}_2\text{GaMnO}_5\) which are attributed to the difficult oxidation of tetrahedral \(\text{GaO}_4\) into octahedral \(\text{GaO}_6\) blocks under 1 atm \(\text{O}_2\) condition. In brief, the ability of \(\text{Ca}_2\text{(Al}_{1-x}\text{Ga}_x)\text{MnO}_5\) OSMs to work both in pressure and temperature-controlled modes enables them great potentials in many energy-related applications.
Acknowledgements

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