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# Implementing vanadium peroxides as direct air carbon capture materials†

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Direct air capture (DAC) removal of anthropogenic CO<sub>2</sub> from the atmosphere is imperative to slow the catastrophic effects of global climate change. Numerous materials are being investigated, including various alkaline inorganic metal oxides that form carbonates *via* DAC. Here we explore metastable early d<sup>0</sup> transition metal peroxide molecules that undergo stabilization *via* multiple routes, including DAC. Specifically here, we describe *via* experiment and computation the mechanistic conversion of A<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> (tetraperoxovanadate, A = K, Rb, Cs) to first a monocarbonate VO(O<sub>2</sub>)<sub>2</sub>(CO<sub>3</sub>)<sup>3-</sup>, and ultimately HKCO<sub>3</sub> plus KVO<sub>4</sub>. Single crystal X-ray structures of rubidium and cesium tetraperoxovanadate are reported here for the first time, likely prior-challenged by instability. Infrared spectroscopy (FTIR), powder X-ray diffraction (PXRD), <sup>51</sup>V solid state NMR (nuclear magnetic resonance), tandem thermogravimetry-mass spectrometry (TGA-MS) along with calculations (DFT, density functional theory) all converge on mechanisms of CO<sub>2</sub> capture and release that involve the vanadium centre, despite the end product of a 300 days study being bicarbonate and metavanadate. Electron Paramagnetic Resonance (EPR) Spectroscopy along with a wet chemical assay and computational studies evidence the presence of ~5% adventitious superoxide, likely formed by peroxide reduction of vanadium, which also stabilizes *via* the reaction with CO<sub>2</sub>. The alkalis have a profound effect on the stability of the peroxovanadate compounds, stability trending K > Rb > Cs. While this translates to more rapid CO<sub>2</sub> capture with heavier alkalis, it does not necessarily lead to capture of more CO<sub>2</sub>. All compounds capture approximately two equivalents CO<sub>2</sub> per vanadium centre. We cannot yet explain the reactivity trend of the alkali peroxovanadates, because any change in speciation of the alkalis from reactions to product is not quantifiable. This study sets the stage for understanding and implementing transition metal peroxide species, including peroxide-functionalized metal oxides, for DAC.

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## Introduction

Burning fossil fuels produces high levels of greenhouse gases including CO<sub>2</sub>, which has led to alarming changes in the Earth's climate, ecosystems and geography.<sup>1,2</sup> To mitigate the negative impacts of climate change, it is essential to transition to low emissions energy technologies, and to also decrease the current high levels of CO<sub>2</sub> in the atmosphere. A number of technologies

have been developed and deployed to decrease legacy and future CO<sub>2</sub> levels, respectively by direct air capture (DAC), and at-the-source capture.<sup>3</sup> Experts agree, however, that no single technology can meet all the demands of rapid and high capacity capture, energy-efficient regeneration and cyclability, and conversion of CO<sub>2</sub> to useful products. Moreover, global, local and seasonal atmospheric differences including temperature, temperature swings, relative humidity, atmospheric pressure, and presence of competing gas phase molecules can either promote or inhibit carbon capture chemistries. Therefore, expanding our chemistry toolbox for carbon dioxide removal can supplement, enhance and diversify DAC technologies.<sup>4-7</sup> Perhaps one of the most intriguing feature of solid phase materials for DAC is their potential to be integrated into existing infrastructure, buildings or vehicles, becoming a more decentralized and flexible solution for carbon capture.<sup>8</sup>

The simplest and most common carbon capture chemistries are amine solutions and basic oxides (both synthetic and earth materials). Aqueous amines is one of the most mature chemical

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technologies, implemented at the source.<sup>9</sup> Amines are inexpensive and effective, but they undergo degradation with reuse, can be corrosive, and require high temperatures and pressures to regenerate *via* CO<sub>2</sub> release.<sup>10</sup> Basic oxides, *i.e.* MgO and CaO are inexpensive and lightweight, and they present similar strengths and limitations as amine solutions, with surface passivation as an additional challenge.<sup>11–14</sup> Other effective solid-phase materials are most often basic oxides or basic hybrid materials, with or without porosity, including zeolites,<sup>15–17</sup> hydrotalcites (synthetic or mine tailings),<sup>18,19</sup> amine-functionalized silica,<sup>20,21</sup> and metal organic frameworks (MOFs).<sup>22–24</sup>

The implementation of peroxide or superoxide ligation of metals (*i.e.* alkalis, alkaline earths) is an interesting approach to solid DAC materials. For example, potassium superoxide has been used since the dawn of space travel to remove CO<sub>2</sub> and replenish O<sub>2</sub> within enclosed space vehicles.<sup>25</sup> These compounds are simple, lightweight, inexpensive and very reactive. However, their reactivity with CO<sub>2</sub> and the resultant product is very sensitive to lattice water and atmospheric water. For example, either carbonate or bicarbonate can form, where the former has a decomposition temperature greater than 600 °C, and the latter has a decomposition temperature less than 150 °C; a critical difference for regeneration.

Peroxometalates of the early d<sup>0</sup> transition metals include titanium,<sup>26</sup> molybdenum,<sup>27</sup> tungsten,<sup>28</sup> niobium,<sup>29</sup> tantalum,<sup>30</sup> chromium<sup>31</sup> and vanadium.<sup>32</sup> These have been known for decades as either homoleptic anions, or more often mixed organic-inorganic hybrid complexes, and heteroligation is one strategy to tune stability/reactivity. Due to their high oxidative reactivity (especially Mo<sup>VI</sup> and V<sup>V</sup>), they have been studied for biological and catalytic applications.<sup>33,34</sup> Notable to the periodic table relationship between early d<sup>0</sup> and f<sup>0</sup> metals, U<sup>VI</sup> as the linear uranyl species (UO<sub>2</sub><sup>2+</sup>) also forms stable peroxide complexes. Transition metal peroxide complexes are typically pseudo-tetrahedral with four bonded peroxides (M(O<sub>2</sub>)<sub>4</sub><sup>x-</sup>), while uranyl binds three peroxides, equatorial to the linear *yl*-dioxo complex (Fig. 1). Though rarely mentioned in the literature because it is deemed a ‘failed experiment’, uranyl triperoxides, as both solids and solutions, will often form uranyl carbonates, *via* reaction with atmospheric CO<sub>2</sub>. However, it is uncertain if this is related to the basicity of the solution, the peroxide reactivity, or both. Forbes *et al.*<sup>35,36</sup> described synthesis of uranyl diperoxide-monosuperoxide, where superoxide is formed from atmospheric O<sub>2</sub> *via* benzyl alcohol oxidation; and this reactive species converts to the uranyl tricarbonate. While the focus of this prior paper is on the superoxide, the end product is uranyl tricarbonate *via* direct air capture, meaning both the peroxide and superoxide react with atmospheric CO<sub>2</sub>, when bonded to uranyl. However, the mechanism of conversion of uranyl peroxide/uranyl peroxide-superoxide to uranyl carbonate *via* DAC is not well-understood, since it happens spontaneously without intent (with both peroxide and superoxide ligands), under a variety of conditions (in solution and as solids). The reactivity of the related transition metal peroxometalates has been largely unexplored. This possibility was mentioned in a review paper<sup>37</sup> concerning the reactivity of

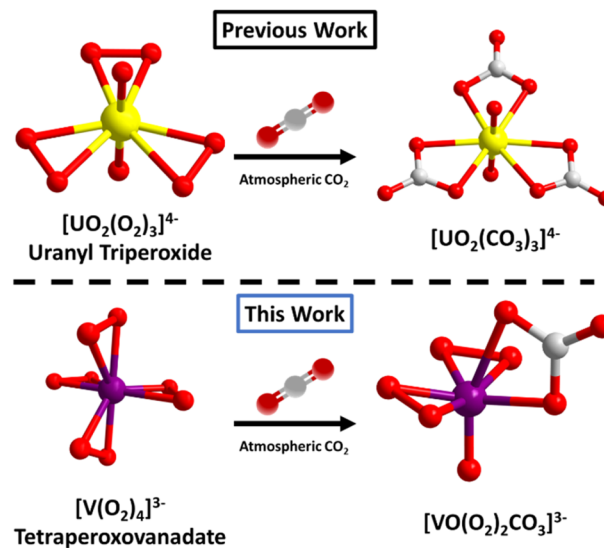


Fig. 1 Comparing reaction of f<sup>0</sup>-actinide peroxide and d<sup>0</sup> transition metal peroxide (superoxide) with atmospheric CO<sub>2</sub>. Under some conditions, uranyl triperoxide (and similar compounds) converts to a uranyl tricarbonate (top).<sup>35</sup> In this work we report a related reaction pathway with tetraperoxovanadates (bottom).

K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> with atmospheric CO<sub>2</sub>, which in turn cites a reference that reports the reactivity of dimeric [V<sub>2</sub>O<sub>3</sub>(O<sub>2</sub>)<sub>4</sub>]<sup>4-</sup> with CO<sub>2</sub>.<sup>38</sup>

Considering transition metal peroxometalates as candidates for DAC, it is hypothetically desirable to have the maximum number of peroxide ligands bound to the metal centre in order to maximize efficiency, so these M(O<sub>2</sub>)<sub>4</sub><sup>x-</sup> species are intriguing candidates. Mixed ligand peroxocarbonato metalates have been synthesized and studied, *i.e.* for second harmonic generation materials.<sup>39,40</sup> However, the transition from peroxo complexes to carbonates by reaction with atmospheric CO<sub>2</sub> has not been extensively investigated. In a related study, Cummins demonstrated the affinity of Mo for carbonate ligands *via* a reaction between orthomolybdate (MoO<sub>4</sub><sup>2-</sup>) and two equivalents of CO<sub>2</sub> in organic solvent.<sup>41</sup> Moreover, reduction of CO<sub>3</sub><sup>2-</sup> to formate was achieved, demonstrating the potential of early d<sup>0</sup> transition metals for not only capture, but conversion.

Here we present a detailed study of the reaction of tetraperoxovanadate with atmospheric CO<sub>2</sub> in ambient lab conditions, using K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> (ref. 42) (compound 1) and synthesized and structurally described here for the first time, Rb<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>·2H<sub>2</sub>O·H<sub>2</sub>O (compound 2) and Cs<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>·3H<sub>2</sub>O (compound 3). A combination of thermogravimetric analysis – mass spectrometry (TGA-MS), <sup>51</sup>V MAS NMR (magic angle spinning nuclear magnetic resonance spectrometry), quantitative carbon analysis, infrared spectroscopy (FTIR) and powder X-ray diffraction (PXRD) benchmarks the conversion of K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> immediately to K<sub>3</sub>VO(O<sub>2</sub>)<sub>2</sub>(CO<sub>3</sub>), and ultimately a mixture containing mostly KVO<sub>3</sub> plus KHCO<sub>3</sub> over several months. Parallel studies of the Rb and Cs analogues show the same reaction pathway, higher reactivity rates, but also a possible competing reaction of autodegradation, or water-promoted degradation. The vanadyl-bound CO<sub>2</sub> is thermally removed below 300 °C for



all three alkali compounds simultaneously with the bound peroxide, indicating a concerted mechanism which is supported by computational studies. While the alkali affects reactivity of the peroxovanadate complex, the vanadium plays a clear role in both CO<sub>2</sub> capture, and its subsequent release. Finally, we also consider the role of superoxides both experimentally and computationally. These studies suggest that superoxide is present in small concentrations, and are likewise reactive towards DAC.

## Results and discussion

### Synthesis

**Warning:** These compounds can be shock sensitive in the dry form, especially with heavier alkali counterions. K, Rb and Cs salts of V(O<sub>2</sub>)<sub>4</sub><sup>3-</sup> were synthesized adapting reported procedures.<sup>42</sup> Details for each compound are included in the ESI.† Summarily, V<sub>2</sub>O<sub>5</sub> was suspended in a basic solution of AOH (A = K, Rb, Cs) and introduced into an ice bath. Then cold 30% H<sub>2</sub>O<sub>2</sub> was added, causing a colour change from brown to green and ultimately purple. The intermediate colour could be due to transient reduction of vanadium, but the final obtained product is pentavalent, based on alkalis for charge-balance as well as experimental precedence (discussed below). After a short time, the solid A<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> is obtained by precipitation with cold ethanol. Crystals suitable for single crystal X-ray diffraction (SCXRD) were obtained by ethanol vapour diffusion at low temperature. The ratios of metal to alkali to peroxide in reaction solutions were commonly 1:5:10 but, a number of other combinations were also attempted. For instance, when the amount of base is increased, the isolation of the material became challenging due to the solid being very effervescent and decomposing quickly. Similarly, when the peroxide amount was lowered to stoichiometric ratios, the isolated crystalline material underwent rapid colour change, from deep purple to yellow. In this case, the small amount of peroxide present is not enough to coordinate to the vanadium and form the tetraperoxovanadate. Despite these challenges, the syntheses of these compounds are reproducible with high yields. Considering all three compounds are obtained from the same procedure, the prior paucity of structures of the Rb and Cs analogues was somewhat surprising, perhaps owed to their high reactivity, similar to the alkali uranyl triperoxides.<sup>43</sup> Single crystals of the Cs-analogue were only obtained when the samples were flushed with argon gas. This was not a necessary for the Rb-analogue, but it helped obtain better quality crystals. This reactivity hints at a clear trend of the alkalis influencing reactivity of the peroxovanadates.

### Structure description

K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> (1) was described prior,<sup>42</sup> but the Rb and Cs analogues are described here for the first time, below (full details on lattice packing and models used for DFT calculations shown in Fig. S1–S4†). All crystallographic information is compiled in Table S1.† Here we briefly note that the simulated PXRD from the single-crystal structures for 2 and 3 do not match that of the

experimental (Fig. S5 and S6).† We attribute this to the ease of partial to complete removal of the free peroxide molecules in the lattice upon isolating the dry powder for PXRD. While compounds 2 and 3 contain extra-framework species according to the single crystal X-ray data (2 peroxide molecules and one water molecule per formula unit for 2; 3 lattice waters per formula unit for 3), the bulk characterization suggests otherwise. Thermogravimetry (Table S2†) of the pristine materials showed weight loss consistent with A<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> for all three compounds. Only the Rb analogue has ~2% excess weight loss, the equivalent of 8.9 mass units, or approximately half a water molecule or a quarter peroxide molecule. Experiments discussed later suggest a small amount of residual reactive oxygen species (peroxide or superoxide) remains in the lattice in the bulk material. The PXRD for K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> also does not match the simulated PXRD from single crystal-data, despite not having extra-framework molecules. In this case (and certainly also relevant for the Rb and Cs analogues); we attribute this lattice mismatch to the observed evolution of oxygen from the solid, disrupting long-range order. Nonetheless, bulk characterization shows the peroxyanions charge-balanced by the appropriate number of alkalis with a starting composition of A<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> comprises the bulk solids. In fact, a starting bulk composition of A<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> for all three compounds means their reactivity is more readily compared, with the alkali being the main difference.

Despite the lattices characterized by SCXRD being different that the studied material, they are worth describing here because; (1) they are new structures, and (2) this provides opportunity to compare the influence of the different alkalis on O–O and V–O bond distance. Finally, for the sake of accuracy in naming, we refer to the bulk material as K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>, Rb<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>, and Cs<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>; or K-analogue, Rb-analogue and Cs-analogue.

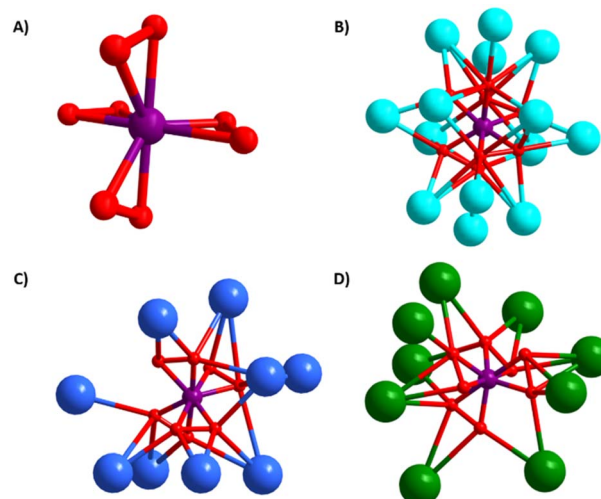


Fig. 2 (A) Ball-and-stick representation of the tetraperoxovanadate core common to all synthesized structures and illustrating the alkali-[V(O<sub>2</sub>)<sub>4</sub>]<sup>3-</sup> bonding for: (B) the K-analogue (1), (C) the Rb-analogue (2), and (D) the Cs-analogue (3). Vanadium is purple, potassium is light blue, rubidium is dark blue, caesium is green and oxygen is red.



All three compounds feature the  $V(O_2)_4^{3-}$  anion (Fig. 2a). The peroxide ligands are in a pseudo-tetrahedral arrangement around  $V^V$  with  $O_p-O_p$  ( $O_p$  = peroxide oxygen) bond lengths ranging from 1.38–1.52 Å, and  $V-O_p$  bond lengths ranging from 1.86–1.98 Å. Notably, the bond lengths found in all the crystal structure are consistent with peroxide; superoxide O–O bond lengths are shorter ( $\sim 1.2$ – $1.3$  Å).<sup>44</sup> In addition, superoxide-peroxide disorder in the electron density maps was not detected. Therefore, the  $\sim 5\%$  adventitious superoxide discussed later is not stoichiometric. While the  $V-O_p$  bond lengths increase slightly with increasing alkali size (Table 1), the  $O_p-O_p$  bond lengths are notably shorter for **1** than for **2** and **3**. Both the total number of bonded alkalis per peroxovanadate and the A–O bond lengths are nearly identical for **2** and **3**, suggesting it is merely packing effects. We also note there is considerably more A– $O_p$  bonding for **1** than for **2** and **3**; either because (1)  $K^+$  is smaller and more can pack around the anion, or (2) while **1** crystallizes with no additional lattice molecules, **2** contains free peroxide and lattice water, and **3** contains lattice water, meaning there are more ligands besides the vanadium-bound peroxide to which the alkalis can bond. Bond valence sum (BVS) calculations are not feasible since parameters for a  $V-O^{1-}$  bond are not available in standard databases, but number of charge-balancing cations confirms pentavalent vanadium, despite the deep purple colour of these compounds (Fig. S7a†).

The Rb-analogue (**2**) crystallizes in the monoclinic system with the space group  $P12_1/n1$  ( $V = 1153.65(6)$  Å<sup>3</sup>) and is fully formulated  $Rb_3[V(O_2)_4] \cdot 2H_2O_2 \cdot H_2O$ . The  $V-O_p$  bond lengths range between 1.876(6)–1.985(5) Å, and  $O_p-O_p$  distances are 1.461(7)–1.470(9) Å. The  $O_p-V-O_p$  angle is  $\sim 45^\circ$  ( $\pm 2^\circ$ ) for all four peroxides, but the  $V-O_p-O_p$  angles vary between 60 and  $70^\circ$ , consistent with the wide range of  $V-O_p$  bond distances.

There are two additional hydrogen peroxide molecules in the lattice. The hydrogen atoms were not located in the electron density map, but they are necessary for charge-balance, and when included in the solid-state calculations, result in a stable structure with an H-bonding network (Fig. S1 and S2†). Protonation is expected, since deprotonation of peroxide is the first step to decomposition, and we would expect a free peroxide anion to be unstable. In addition, the free peroxide oxygen atoms are located 2.7 Å from the vanadium-bound peroxide ligands, consistent with hydrogen-bonding. A prior structure reported for the Na-analogue of  $V(O_2)_4^{3-}$  also contained free peroxide molecules.<sup>32</sup> The hydrogen peroxide molecules are bonded terminally, bridging and side-on to Rb with bond lengths ranging from 2.98 to 3.43 Å (Fig. S1a†). Finally, a water molecule coordinates between Rb2 and Rb3, a lattice arrangement reported in prior structures.<sup>28,32</sup> The  $[V(O_2)_4]^{3-}$  anions are aligned along the  $a$ -axis in the lattice. The Rb1 and Rb3 each

form four-atom parallel rows that are broken by a zigzag column of Rb2 cations (Fig. S1b†). This bears resemblance to the prior-reported sodium analogue structure, where the cations interconnect with one another through bound water in octahedral configuration.<sup>32</sup>

Compound **3** crystallizes in the monoclinic space group  $P12_1/n1$  ( $V = 1162.57(8)$  Å<sup>3</sup>) and is fully formulated  $Cs_3[V(O_2)_4] \cdot 3H_2O$ . The  $[V(O_2)_4]^{3-}$  of compound **3** is more symmetric than that of compound **2**; with deviation between the bonds significantly lower in compound **3**; almost three times lower for  $V-O$  bonds and over five times lower for  $O-O$  bonds. The same is true for the  $O-V-O$  and  $V-O-O$  angles, which also have an average of  $45^\circ$  and  $67^\circ$  respectively, but with a smaller range in both cases. This suggests the distortions in **2** arise from H-bonding to the free peroxide.

Compound **3** has three lattice water molecules per formula unit, each water molecule sits between four  $Cs^+$  and one tetraperoxovanadate. The lack of free hydrogen peroxide molecules in the lattice is a notable difference between compounds **2** and **3**, but is also not unprecedented for this family of compounds and the DFT calculated unit cell of  $Cs_3[V(O_2)_4] \cdot 3H_2O$  with waters explicitly included converged to a stable geometry (Fig. S3 and S4†). The most-studied  $Na_3V(O_2)_4$  compound has been crystallized with a range of  $H_2O_2$  and  $H_2O$  in its lattice,<sup>32</sup> and it is likely that other ordered lattices for K, Rb and Cs analogues could be isolated, with or without the inclusion of free  $H_2O$  and  $H_2O_2$ .

The lattice of **3** features alternating layers of alkalis and tetraperoxovanadates, stacked in the  $b$ -direction (Fig. S3c†). Along the  $a$ -axis (Fig. S3d†), parallel layers of  $Cs_2$  and  $Cs_3$  alternate with double layers of  $Cs_1$ . Viewed along the  $c$ -direction, the  $Cs^+$  cations appear as triads of all three independent Cs atoms (Fig. S3b†), bearing resemblance to the  $Na_3[V(O_2)_4]$  lattice.<sup>32</sup> The main difference, as discussed above, is that the Na-triads are interconnected through lattice water for the Na-analogue.

### Reaction with atmospheric CO<sub>2</sub>

All analogues of  $A_3V(O_2)_4$  react immediately upon air exposure, if not stored cold and preferably (esp. Rb and Cs analogues) under an inert atmosphere, best depicted by a colour change, from purple to yellow (Fig. S7†). This colour change is commensurate with reaction of the unstable oxygen ligands (peroxide and adventitious superoxide), either *via* auto-decomposition or capture of atmospheric CO<sub>2</sub>, discussed below. The macroscopic colour change occurs more rapidly for  $Rb_3V(O_2)_4$  and  $Cs_3V(O_2)_4$ ; no purple is visible after just 5 days, while  $K_3V(O_2)_4$  retains a purple hue two weeks to a month. This

Table 1 Summary of alkali- $[V(O_2)_4]^-$  bonding (Å) for compounds **1**, **2** and **3**

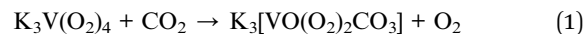
Alkali cation	$V-O_p$ , avg	$O_p-O_p$ , avg	$A^+-O_p$ , avg	# $A^+$ bonded to each $[V(O_2)_4]^-$	# $A^+-O_p$ bonds per $[V(O_2)_4]^-$
K ( <b>1</b> )	1.9181	1.5508	2.8822	14	31
Rb ( <b>2</b> )	1.9198	1.4654	3.0982	9	21
Cs ( <b>3</b> )	1.9276	1.4762	3.0985	9	20



is in agreement with previous reports that describe larger alkali tetraperoxovanadates (and also Mo and W counterparts) as more reactive, or even explosive.<sup>45</sup> That said, this faster reactivity does not necessarily translate to higher carbon capture, as revealed by characterization of the degradation products *via* Fourier transform infrared (FTIR), thermogravimetric analysis/mass spectrometry (TGA-MS), powder X-ray diffraction (PXRD), <sup>51</sup>V solid state MAS NMR (magic angle spinning nuclear magnetic resonance) and elemental analysis, described below.

To investigate the changes of A<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>, the K-analogue was left on a lab bench under ambient conditions (temperature ~21 °C). FT-IR spectra were recorded over time as a first indication of moieties present in the decomposition products (Fig. 3, S8 and S9 for K, Rb and Cs analogues, respectively). Here we describe the spectral bands citing K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>, and bands for Rb<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> and Cs<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> are summarized in Table S3.† It is apparent that these compounds react with atmospheric CO<sub>2</sub> to form carbonates due to the presence of characteristic bidentate carbonate peaks at 1570 cm<sup>-1</sup>, 1340 cm<sup>-1</sup>, and 1050 cm<sup>-1</sup>.<sup>46</sup> The increasing absorbance at 935 cm<sup>-1</sup> and 735 cm<sup>-1</sup> were assigned to V=O stretching and V-O vibration respectively, both

indicating a change in the vanadium-oxygen bonding mode. This was further confirmed by the peak growth at 740 cm<sup>-1</sup>, which matches the V-O-C vibrational mode and implies that the carbonate is bonded to the vanadium centre. The absence of these peaks in the fresh K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> evidences DAC is the source. Other peaks were assigned to O-O stretching (peroxide, 850 cm<sup>-1</sup>) and V-(O<sub>2</sub>) vibrations (625 cm<sup>-1</sup>, 590 cm<sup>-1</sup>, and 560 cm<sup>-1</sup>). Overall, the new spectrum resembles that of synthesized K<sub>3</sub>[VO(O<sub>2</sub>)<sub>2</sub>CO<sub>3</sub>].<sup>39</sup> We can describe the reaction as:



Solid-state density-functional theory (DFT) calculations were carried out in CASTEP<sup>47</sup> (see ESI† for details) to assist in the interpretation of the experimental IR spectra. Phonon calculations of the starting material, K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>, the monocarbonate (K<sub>3</sub>VO(O<sub>2</sub>)<sub>2</sub>CO<sub>3</sub>), and other species thought to be involved in decomposition/side reactions were simulated in this manner, namely KOH, K<sub>2</sub>O, K<sub>3</sub>VO<sub>4</sub>, and K<sub>2</sub>CO<sub>3</sub>. Comparison between calculated and observed peaks in solid-state IR is complicated

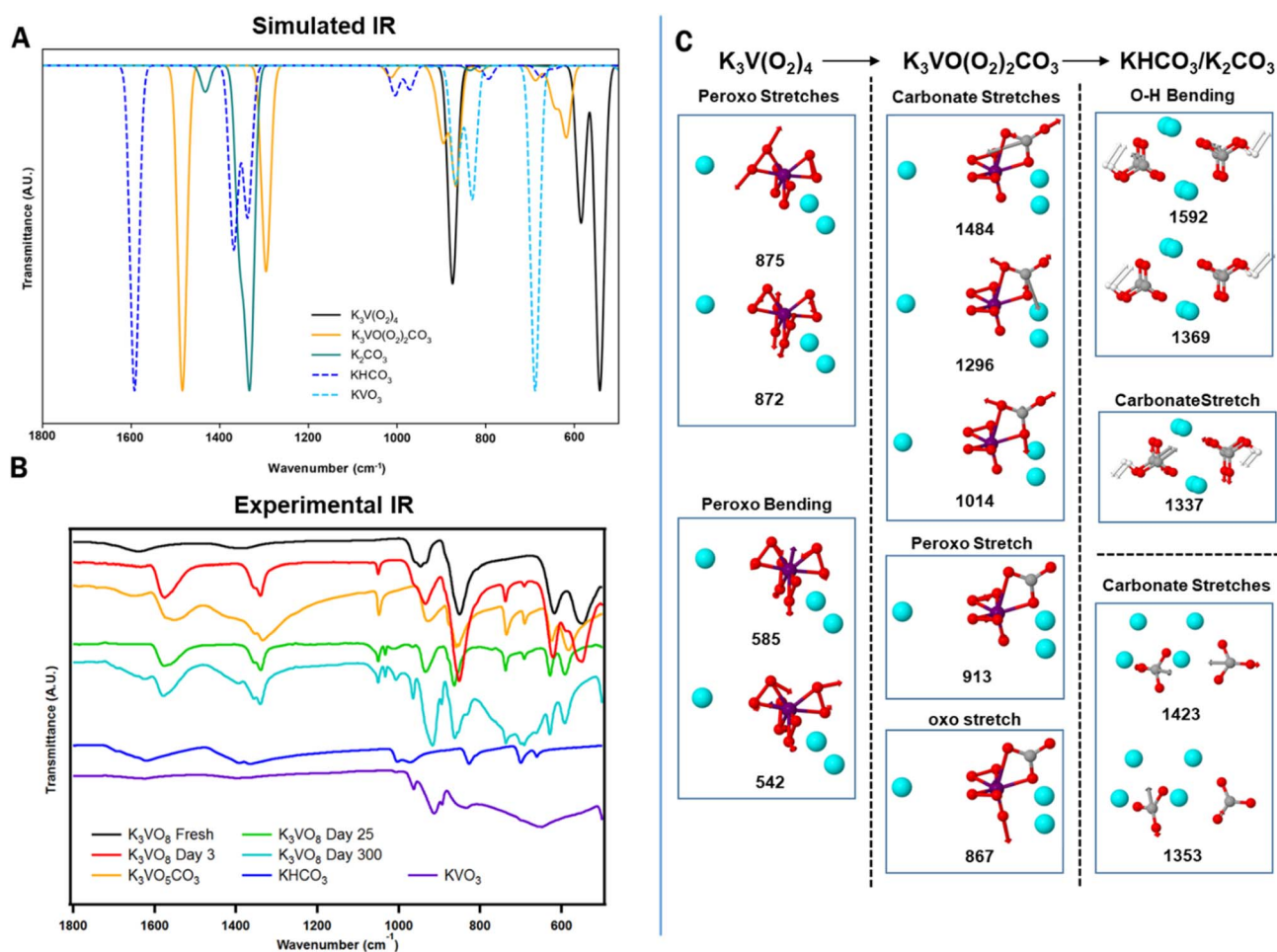
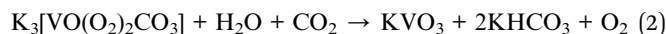


Fig. 3 Comparison of the calculated IR spectrum of K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> and its possible decomposition products (A) with the experimental data (B). Representative carbonate peaks for K<sub>3</sub>[VO(O<sub>2</sub>)<sub>2</sub>(CO<sub>3</sub>)] (1570 cm<sup>-1</sup>, 1340 cm<sup>-1</sup>, and 1050 cm<sup>-1</sup>) are observed in Day 3, Day 25, and Day 300 samples. The spectrum of Day 300 also contains IR peaks representative of KHCO<sub>3</sub> (1630 cm<sup>-1</sup>, 1390 cm<sup>-1</sup>, and 1010 cm<sup>-1</sup>) and KVO<sub>3</sub> (965 cm<sup>-1</sup>, and 890 cm<sup>-1</sup>). The modes of vibration to which they are assigned are shown in (C).



by the experimental inevitability of there being a superposition of reactant, product, and other species. Other discrepancies can be attributed to inaccuracies in the unit cells, lack of surface terminations in the calculations and the method's own limitations. Nonetheless, a few supported comments can be made here. If only considering motions above  $500\text{ cm}^{-1}$ ,  $\text{K}_3\text{V}(\text{O}_2)_4$  has effectively four signals: V–O–O bending motions at  $542$  and  $585\text{ cm}^{-1}$ , and O–O stretching motions at  $871$  and  $876\text{ cm}^{-1}$ , as would be expected for a homoleptic metal-peroxide compound.<sup>48</sup>  $\text{K}_3\text{VO}(\text{O}_2)_2\text{CO}_3$  also has signals in this region including a V=O stretch at  $867\text{ cm}^{-1}$ , an asymmetric peroxy stretch at  $895\text{ cm}^{-1}$ , and a symmetric peroxy stretch at  $914\text{ cm}^{-1}$ . Distinct from  $\text{K}_3\text{V}(\text{O}_2)_4$ , however,  $\text{K}_3\text{VO}(\text{O}_2)_2\text{CO}_3$  is predicted to have three intense peaks above  $1000\text{ cm}^{-1}$  attributed to the carbonate group. These include symmetric  $[\text{CO}_3]^{2-}$  stretch at  $1014\text{ cm}^{-1}$ , and asymmetric  $[\text{CO}_3]^{2-}$  stretches at  $1296\text{ cm}^{-1}$  and  $1484\text{ cm}^{-1}$ . This collection of peaks are found experimentally only after the  $\text{A}_3\text{V}(\text{O}_2)_4$  sample has been exposed to the air. Further confirmation of this transition comes from PXRD patterns. Fig. 4 displays a structural transition from  $\text{K}_3\text{V}(\text{O}_2)_4$  to  $\text{K}_3[\text{VO}(\text{O}_2)_2\text{CO}_3]$  by ingrowth of peaks that are consistent with the latter compound. Longer ambient exposure times bring a new set of peaks at  $2\theta$  ( $^\circ$ ) =  $17.2$ ,  $24.3$ ,  $24.7$ ,  $34.2$ ,  $34.6$ , and  $36.9$ ; indicating a mixture of  $\text{KHCO}_3$  and  $\text{KVO}_3$ . Given the order of peak appearance, we deduce that these simple salts form *via* capture of additional  $\text{CO}_2$ , thus marking a possible end-product of the reaction, which can be formulated as:



It is worth noting, however, that peaks for both  $\text{K}_3[\text{VO}(\text{O}_2)_2\text{CO}_3]$  and  $\text{K}_3[\text{V}(\text{O}_2)_4]$  persist, despite the ingrowth of peaks for the secondary decomposition products, which is further verified by TGA-MS experiments, discussed later.

We synthesized prior-reported  $\text{K}_3[\text{VO}(\text{O}_2)_2\text{CO}_3]$  to use as a standard for various spectroscopic characterizations (see ESI† for synthesis details).<sup>42</sup> The  $^{51}\text{V}$  MAS NMR of synthesized

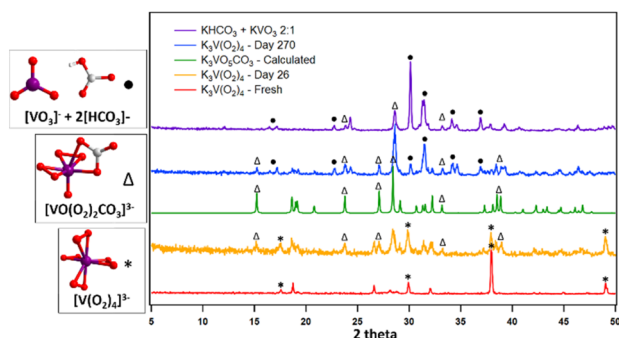


Fig. 4 PXRD patterns of  $\text{K}_3\text{V}(\text{O}_2)_4$  at different time stamps compared to the synthesized  $\text{K}_3\text{VO}(\text{O}_2)_2\text{CO}_3$  as an intermediate and  $\text{KVO}_3$  and  $\text{KHCO}_3$  as end products. Peaks corresponding to compounds containing the key molecular species are shown on the left and marked accordingly.

$\text{K}_3[\text{VO}(\text{O}_2)_2\text{CO}_3]$  and **1** aged for nine months is shown in Fig. 5.  $\text{K}_3[\text{VO}(\text{O}_2)_2\text{CO}_3]$  has a single  $^{51}\text{V}$  peak at  $789\text{ ppm}$ . The central peak was identified as the non-shifting peak upon changing spin speeds (Fig. S10†), and the remainder are spinning side bands. The single  $^{13}\text{C}$  peak for this compound is located at  $161\text{ ppm}$ . Aging  $\text{K}_3\text{V}(\text{O}_2)_4$  for 9 months yields a  $^{51}\text{V}$  MAS NMR spectrum with two peak envelopes (Fig. 5b). The minor envelope (spinning sidebands marked by red stars) is  $\text{K}_3[\text{VO}(\text{O}_2)_2\text{CO}_3]$ , and the major envelope (spinning sidebands marked by black stars) is  $\text{KVO}_3$ ,<sup>49</sup> consistent with both IR and PXRD. Notably, we were not able to collect data on any fresh  $\text{V}(\text{O}_2)_4^{3-}$ . A single attempt led to ‘explosion’ of the sample rotor, because the excessive heat generated by spinning rapidly decomposed peroxide ligands.

To gain insight into the possible mechanistic steps associated with the conversion of gaseous  $\text{CO}_2$  into complexed  $[\text{CO}_3]^{2-}$ , cluster-model (Fig. 6A) DFT calculations in Gaussian<sup>50</sup> were performed on one unit of the tetraperoxovanadate complex with three potassium ions explicitly included (see ESI† for details). The resulting proposed mechanism (summarized in Fig. 6B; see Fig. S11† for complete diagram) consists of five steps for the  $\text{CO}_2$  addition: (1) barrierless addition of  $\text{CO}_2$  to  $\text{K}_3\text{V}(\text{O}_2)_4$ , forming  $\text{K}_3\text{V}(\text{O}_2)_3(\kappa^1\text{-CO}_4)$ ; (2) isomerization of

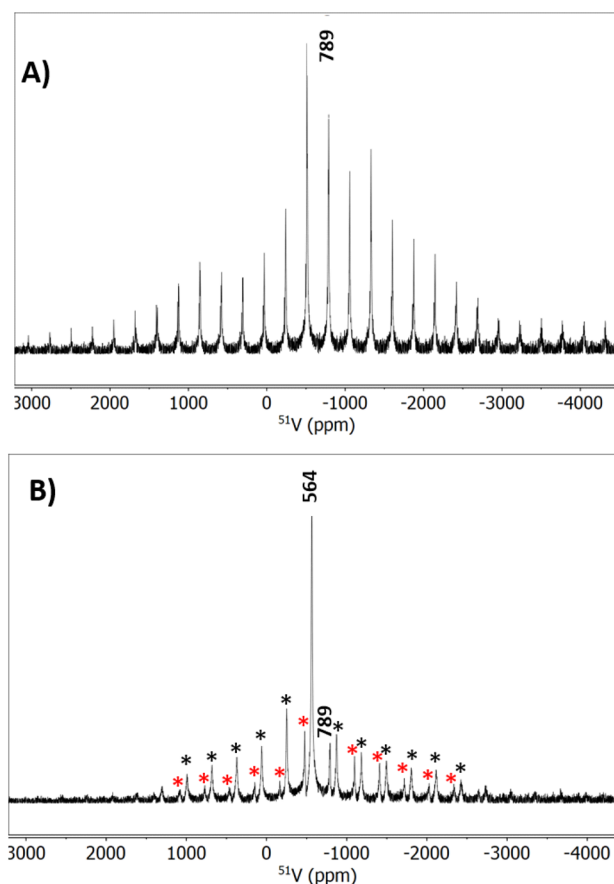


Fig. 5  $^{51}\text{V}$  MAS NMR for (A)  $\text{K}_3\text{VO}(\text{O}_2)_2\text{CO}_3$  and (B)  $\text{K}_3\text{V}(\text{O}_2)_4$  aged 9 months. The red stars indicate spinning sidebands for the peak at  $789\text{ ppm}$  ( $\text{K}_3\text{VO}(\text{O}_2)_2\text{CO}_3$ ) and the black stars indicate spinning sidebands for  $\text{KVO}_3$  ( $564\text{ ppm}$ ).



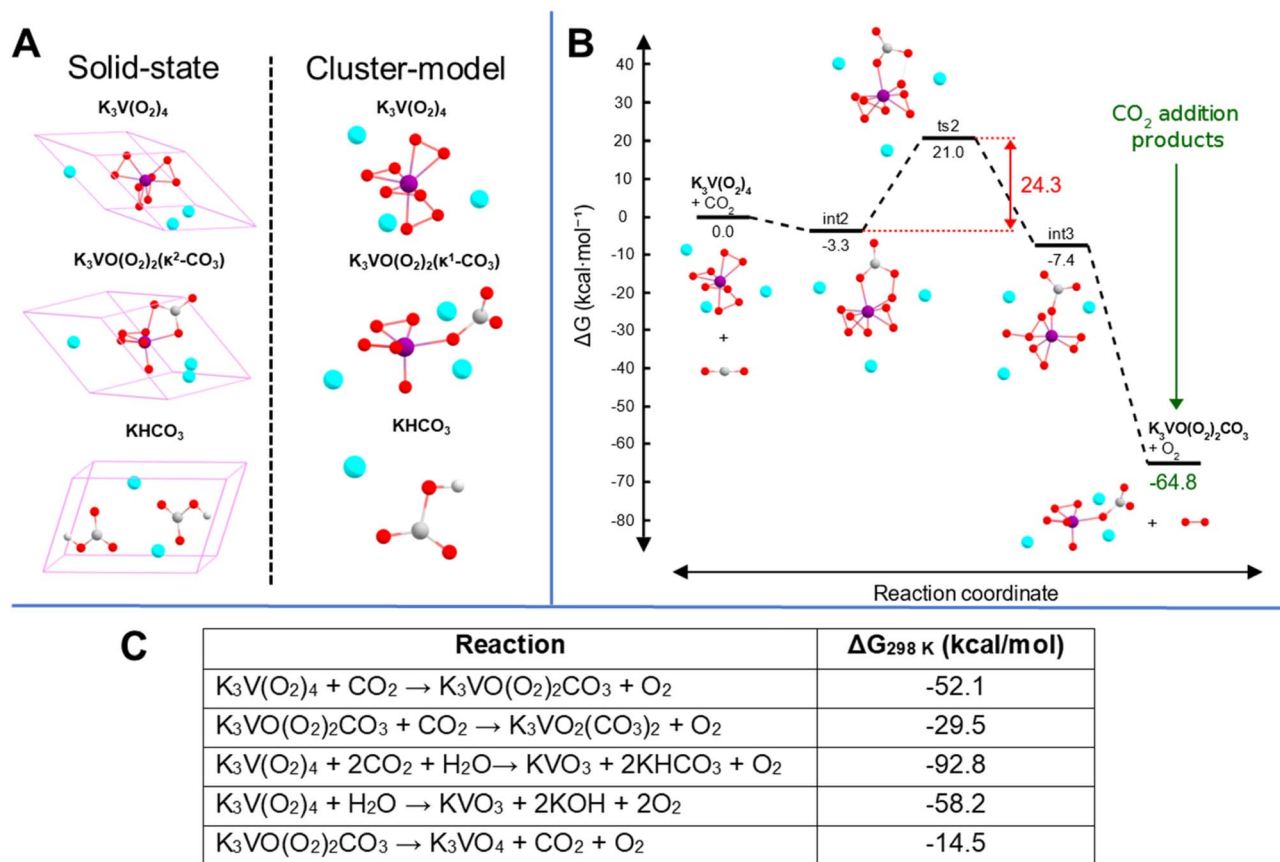


Fig. 6 (A) Two computational modelling approaches used: solid-state DFT with periodic boundary conditions to simulate IR spectra and gas-phase cluster-model DFT to describe CO<sub>2</sub> capture by K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>. (B) Simplified reaction mechanism, displaying only the thermodynamics and rate-limiting steps associated with the addition of CO<sub>2</sub> to K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>. (C) Room-temperature solid-state thermodynamics of some selected reactions.

K<sub>3</sub>V(O<sub>2</sub>)<sub>3</sub>(κ<sup>1</sup>-CO<sub>4</sub>) to K<sub>3</sub>V(O<sub>2</sub>)<sub>3</sub>(κ<sup>2</sup>-CO<sub>4</sub>), (3) transfer of an oxygen atom from the peroxy carbonate to one of the peroxo ligands to yield (η<sup>2</sup>-O<sub>3</sub>)(κ<sup>1</sup>-CO<sub>3</sub>), (4) detachment of one of the coordinating O atoms in η<sup>2</sup>-O<sub>3</sub>, meaning η<sup>2</sup>-O<sub>3</sub> becomes η<sup>1</sup>-O<sub>3</sub>; (5) cleavage of the dangling O<sub>2</sub> from η<sup>1</sup>-O<sub>3</sub>, producing K<sub>3</sub>VO(O<sub>2</sub>)<sub>2</sub>(κ<sup>1</sup>-CO<sub>3</sub>). The formation of a peroxy carbonate ligand is the natural starting point from chemical intuition and is consistent with previous computational and experimental studies that demonstrate the ability of peroxo-containing metal species to form stable peroxy carbonate complexes.<sup>51,52</sup> Additionally, the transfer of an oxygen atom to form an ozone intermediate in step 3 (see Fig. S11†) has some literature precedent in zirconium structures featuring η<sup>2</sup>-O<sub>3</sub> involved in the epoxidation of propene.<sup>53</sup> Although the cluster-model product geometry, K<sub>3</sub>VO(O<sub>2</sub>)<sub>2</sub>(κ<sup>1</sup>-CO<sub>3</sub>), does feature the carbonate ligand bound in a κ<sup>1</sup>-fashion that is different from both the experimental and solid-state computational structures, this binding mode can be rationalized on the basis of the lack of crystal packing in the cluster (vacuum) calculation and the small atomic radius of V<sup>5+</sup>. We performed analogous simulations with the heavier group congeners Nb and Ta, and found that only Ta exhibits bidentate bonding of the carbonate to the metal centre (see Fig. S12 and S13†). As shown in Fig. 6B, the overall net change in Gibbs free energy for the CO<sub>2</sub> addition is -64.8 kcal mol<sup>-1</sup> with a rate

limiting barrier of 24.3 kcal mol<sup>-1</sup>. Given the predictions of highly exergonic free energy changes, surmountable kinetic barriers for a room-temperature reaction, and qualitatively matching IR simulations, we believe the reaction described here is a plausible pathway for the initial carbon capture process observed, invoking conversion of K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> to K<sub>3</sub>VO(O<sub>2</sub>)<sub>2</sub>CO<sub>3</sub>.

We attempted to explain the relative reactivity dependence of V(O<sub>2</sub>)<sub>4</sub><sup>3-</sup> on the alkalis (Cs > Rb > K) using computation. If an idealized K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> structure is assumed for the Rb and Cs analogues, the calculated solid-state relative electronic energy of Rb<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> is destabilized by 25.0 kcal mol<sup>-1</sup> relative to K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>, while Cs<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> is destabilized by 48.9 kcal mol<sup>-1</sup> compared to the K-analogue. Interestingly, despite the notable difference in stability between all three compounds, the  $\Delta G$  of the decomposition reactions (described in eqn (1) and (2)) are rather similar. We had expected this, due to the higher intrinsic reactivity of the Rb and Cs analogues. However, the presence of a larger alkali destabilizes both the starting materials and the products of the reaction, meaning that the net effect is comparable between all three reactions, so the experimentally observed difference in reactivity cannot be attributed to simple electrostatic differences in the counterions. Our computational method is limited by the potential role of reagents that cannot be accounted for in a balanced chemical reaction, e.g. water,



free hydrogen peroxide, hydroxide, and even the chemical role of the alkalis.

The PXRD patterns of  $\text{Rb}_3\text{V}(\text{O}_2)_4$  and  $\text{Cs}_3\text{V}(\text{O}_2)_4$ , while displaying similar behaviour to those of  $\text{K}_3\text{V}(\text{O}_2)_4$  initially, show no evidence for the  $\text{A}_3[\text{VO}(\text{O}_2)_2\text{CO}_3]$  intermediate beyond the 90 days mark (Fig. S5 and S6†). This suggests that after a 3 months period, all the  $\text{CO}_2$  captured is transformed to  $[\text{HCO}_3^-]$  and the vanadium complex has fully decomposed to its simplest metavanadate form, with no remaining peroxide. While this information can be extracted from the PXRD data; as the material decomposes over time it also loses crystallinity, making peak assignment somewhat more challenging. Therefore, in order to confirm this observation as well as determine temperature and mechanism of  $\text{CO}_2$  release and monitor the thermal degradation pathway, thermogravimetric analysis-mass spectrometry (TGA-MS) was employed.

Fig. 7 displays the simultaneous thermogravimetric analysis and mass spectrometry (TGA-MS) analysis of  $\text{K}_3\text{V}(\text{O}_2)_4$  with aging (up to 9 months) in the ambient lab atmosphere, and the data discussed below is summarized in Table S4.† Mass spectrometry identifies volatile species evolved from  $\text{K}_3\text{V}(\text{O}_2)_4$ , and observed changes with time are consistent with PXRD, IR and  $^{51}\text{V}$  NMR analyses. The fresh  $\text{K}_3\text{V}(\text{O}_2)_4$  served as a reference and a starting point of the carbon capture reaction (Fig. 7A). The loss of molecular oxygen originated from the four peroxide ligands

bound to the metal centre occurs at 120 °C. The absence of any  $\text{CO}_2$  signals in the mass spectrum indicates the fresh product is pure, prior to initiating carbon capture.

After 5 days on the lab benchtop, the oxygen peak from unreacted tetraperoxovanadates remains, but a new simultaneous release of oxygen and carbon dioxide appears at 240 °C (Fig. 7B). This simultaneous release is assigned to peroxides and carbonates bonded to vanadium, namely the newly formed  $\text{K}_3\text{VO}(\text{O}_2)_2\text{CO}_3$ , also observed by PXRD (Fig. S15†). This was confirmed by performing the TGA-MS experiment on the standard (synthesized)  $\text{K}_3\text{VO}(\text{O}_2)_2\text{CO}_3$ , which displayed a comparable simultaneous weight loss of  $\text{O}_2$  and  $\text{CO}_2$  (Fig. S16†). We used computation to understand the mechanism of simultaneous release. These studies revealed favourable thermodynamics for the two-step sequential removal of  $\text{CO}_2$  and  $\text{O}_2$  from  $\text{K}_3\text{VO}(\text{O}_2)_2\text{CO}_3$  to form  $\text{K}_3\text{VO}_4$  at room and elevated temperatures (Fig. S14†). A new peak at 370 °C that broadens in later spectra also appears after 10 days of ambient air exposure. This  $\text{CO}_2$  release appears to originate from decomposition of the formed alkali carbonate, catalysed by the vanadium present in the mixture (discussed below). In addition, higher temperature  $\text{CO}_2$  loss (670 °C) originating from carbonates bonded to alkali metals is observed. The  $\text{CO}_2$  has a weight percent of 8.29% and the  $\text{O}_2$  is 13.41%, where the water only accounts for 2.26%. Similar signals can be found after 10 days (Fig. 7C), with an

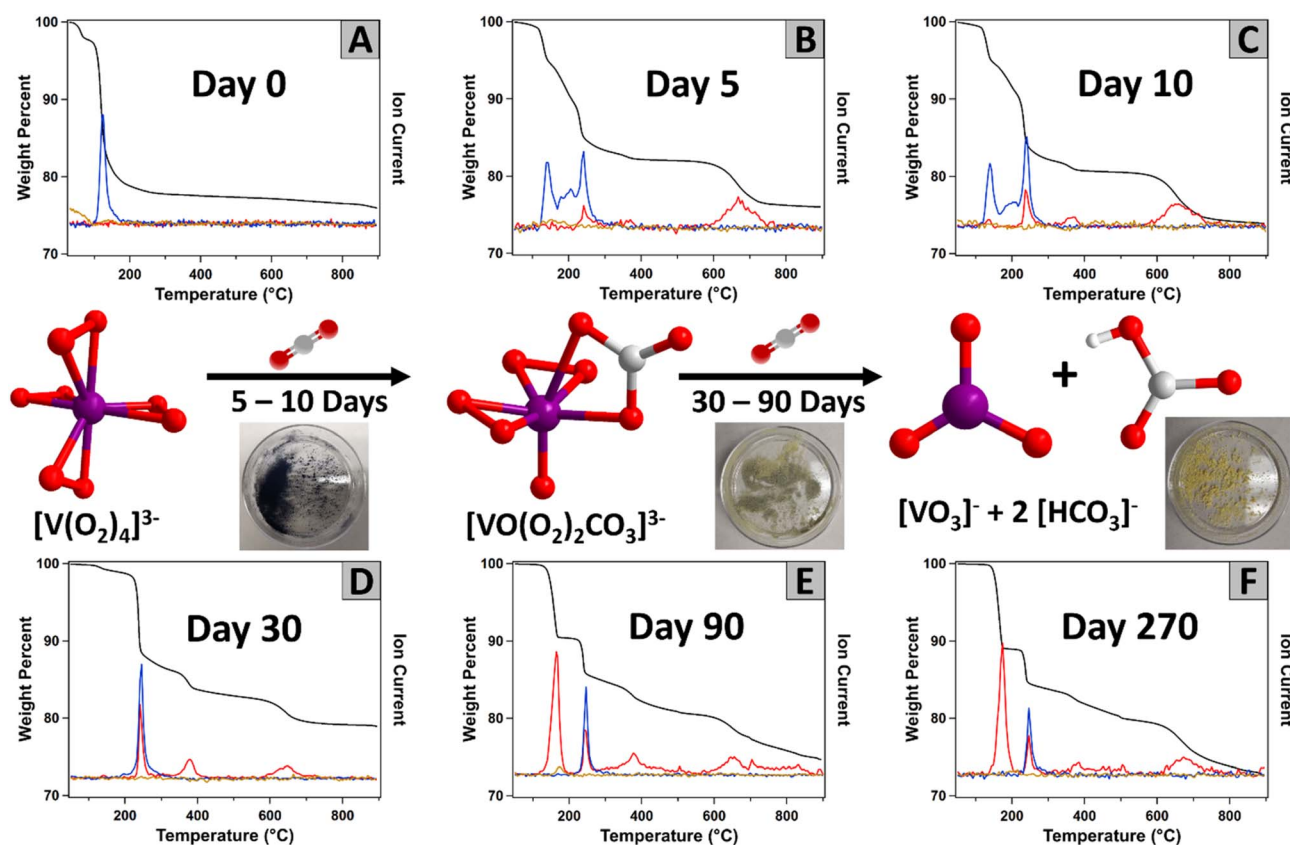


Fig. 7 Evolution of TGA-MS of  $\text{K}_3\text{V}(\text{O}_2)_4$  following exposure to ambient air with time (increasing time from (A) to (F)). The thermogravimetric signal is black. The red, blue, and light orange traces are the signals of  $\text{CO}_2$ ,  $\text{O}_2$  and  $\text{H}_2\text{O}$  by tandem mass spectrometry, respectively. Photographs show the colour evolution of the solid materials with aging in ambient conditions.



increase in the carbon dioxide signal (weight percent of CO<sub>2</sub>, O<sub>2</sub>, and H<sub>2</sub>O were 10.37%, 13.54%, and 2.16%, respectively).

With continued ambient exposure, the oxygen signal at 140 °C is completely absent after 30 days (Fig. 7D), indicating that any remaining K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> decomposed to other materials. At this point, the multiple CO<sub>2</sub> signals have increased (total weight loss 12.36%), and less O<sub>2</sub> (weight loss 8.48%) was released. The CO<sub>2</sub> peak at 165 °C, dominates after 90 days of exposure (Fig. 7E). This peak is assigned to the decomposition of bicarbonate and suggests that this new carbon environment is the most prevalent form in the material. We confirmed this by preparing a mixture of potassium bicarbonate and KVO<sub>3</sub> (as observed by PXRD) and performing the TGA-MS experiment on this mixture (Fig. S17†). In this reference spectrum, the peak appearing at 300–500 °C can also be observed, confirming it belongs to a vanadium catalysed release of CO<sub>2</sub>. Between day 90 and day 270, the TGA-MS data exhibited little change. The weight loss is dominated by CO<sub>2</sub> (weight loss 21.16%) and some remaining O<sub>2</sub> from the [VO(O<sub>2</sub>)<sub>2</sub>CO<sub>3</sub>]<sup>3-</sup> intermediate (weight loss 2.88%). A series of IR spectra were run of this sample after being heated at different temperatures. As expected, there is a decrease in the intensity of the carbonate bands after the sample is heated to temperatures higher than 300 °C in favour of bands assigned to V=O stretches (Fig. S18†). In summary, the TGA-MS and PXRD data both show an evolution of V(O<sub>2</sub>)<sub>4</sub><sup>3-</sup> to VO(CO<sub>3</sub>)(O<sub>2</sub>)<sub>2</sub><sup>3-</sup> to VO<sub>3</sub><sup>-</sup> plus bicarbonate, with all anions charge-balanced by potassium. At each step of air exposure, VO(CO<sub>3</sub>)(O<sub>2</sub>)<sub>2</sub><sup>3-</sup> is always present.

The thermal behaviour of the Rb and Cs analogues following ambient air exposure was investigated in the same way as K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>. Details for these results for Rb<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> are in Fig. S19–S24, summarized in Table S5, and in Fig. S25–S30, summarized in Table S6,† for Cs<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>. There are distinct differences between the three analogues. While the oxygen release occurs in a single step at 120 °C for 1, the oxygen loss for both Rb<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> and Cs<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> are represented by two temperature maxima, 130 °C and 180 °C. The shifts observed in the CO<sub>2</sub> peak positions at 300–400 °C and 550–750 °C can be explained by the varying thermal stabilities of the alkali carbonates, to which these peaks are assigned (Fig. S31†). In turn, the minimal shift for the CO<sub>2</sub> released at 250 °C confirms this peak to correspond to vanadium-bonded carbonate. However, the biggest difference between all three compounds appears after 90 days when the simultaneous release of CO<sub>2</sub> and O<sub>2</sub> at 250 °C, characteristic of the [VO(O<sub>2</sub>)<sub>2</sub>CO<sub>3</sub>]<sup>3-</sup> complex, disappears for Rb<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> and Cs<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>. This is confirmed by the PXRD data that shows after 90 days, the [VO(O<sub>2</sub>)<sub>2</sub>CO<sub>3</sub>]<sup>3-</sup> complex is fully decomposed to bicarbonate and metavanadate (Fig. S5 and S6†). This key difference between the K-analogue and Rb<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> and Cs<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> can, once again, be explained based on the relative stability of the different compounds.

To quantify CO<sub>2</sub> capture per tetraperoxovanadate molecule, we used CHN microanalysis at similar time stamps as the above-described analyses. In addition to this, the weight loss and the integration of the peak areas from the TGA-MS spectra was also used to validate the results. Fig. 8 shows the capture of CO<sub>2</sub> occurs most rapidly the first few days of atmospheric exposure,

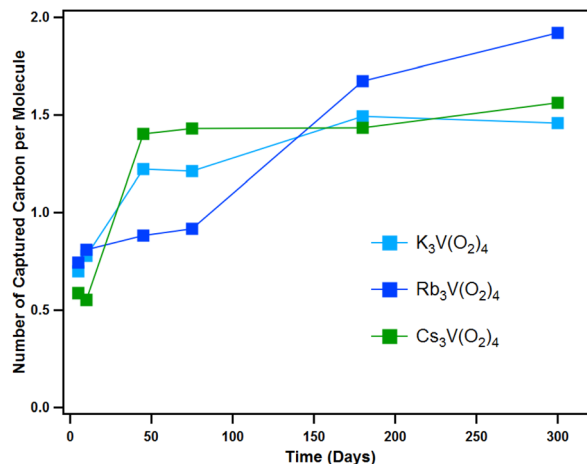


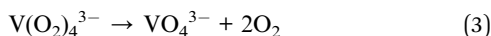
Fig. 8 Carbon capture performance for the three A<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> analogues under ambient conditions evaluated by CHN analysis.

at which time the total carbon captured changes notably on a daily basis (full details of CHN analysis are shown in Tables S7–S9 and Fig. S32–S35†). This is consistent with the energetically favourable conversion of V(O<sub>2</sub>)<sub>4</sub><sup>3-</sup> to [VO(O<sub>2</sub>)<sub>2</sub>CO<sub>3</sub>]<sup>3-</sup> ( $\Delta G = -51.6$  kcal mol<sup>-1</sup>,  $-53.2$  kcal mol<sup>-1</sup> and  $-51.4$  kcal mol<sup>-1</sup> for K, Rb and Cs analogues, respectively, in the solid-state). As discussed above, this intermediate continues to degrade into even more stable products, namely metavanadate and bicarbonate ( $\Delta G = -41.2$  kcal mol<sup>-1</sup>,  $-41.5$  kcal mol<sup>-1</sup> and  $-41.1$  kcal mol<sup>-1</sup> for K, Rb and Cs analogues, respectively). The possibility of binding the second CO<sub>2</sub> molecule to the peroxyvanadate complex was hypothesized briefly. Computational calculations show a favoured process, with a net free energy of  $-50.4$  kcal mol<sup>-1</sup> as a cluster-model ( $-29.5$  kcal mol<sup>-1</sup> in the solid-state) with a rate limiting barrier of 24.5 kcal mol<sup>-1</sup>. However, this product was never observed. After the initial transformation, the capture rate seems to slow down and plateau at 1.5–2.0 CO<sub>2</sub> per molecule. The fact that all three compounds reach a similar endpoint is expected since the reaction path is the same for all three, and they have similar overall change in free energy ( $\Delta G$ ). Therefore, even though their thermodynamic behaviour and reaction pathway is similar, Cs<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub> reacts faster. Furthermore, the prospect of binding a third CO<sub>2</sub> molecule through a reaction with the remaining peroxy ligand in hypothetical K<sub>3</sub>VO<sub>2</sub>(CO<sub>3</sub>)<sub>2</sub> to form K<sub>3</sub>VO( $\kappa^1$ -CO<sub>3</sub>)<sub>3</sub> was considered, but was found to be uphill in relative free energy by 3.2 kcal mol<sup>-1</sup>, and the fourth addition product, K<sub>3</sub>V( $\kappa^1$ -CO<sub>3</sub>)<sub>4</sub>, did not result in a stable complex with cluster calculations.

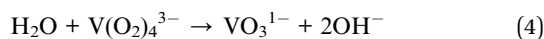
The alkali counteranions play a role in the reactivity of V(O<sub>2</sub>)<sub>4</sub><sup>3-</sup>, with a clear trend in increasing reactivity with increasing alkali size. However, the exact mechanism, and therefore the role of the alkalis is still poorly understood, despite the observed consequences. On the other hand, the number of CO<sub>2</sub> molecules per metal centre (1.5 to 2) is similar for K<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>, Rb<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>, and Cs<sub>3</sub>V(O<sub>2</sub>)<sub>4</sub>; suggesting the effect of the alkalis is largely kinetic. At the end of the experiment time (300 days), the K-analogue exhibits carbonate directly bound to



the vanadium as well as carbonate bound to the alkalis, based on release temperature in TGA-MS and PXRD. On the other hand, the Rb and Cs analogues only exhibit carbonate bound to the alkalis after 300 days of ambient exposure. Competing decomposition pathways of the peroxometalates that complicate mechanistic understanding is auto-degradation (eqn (3)) or water-promoted degradation (eqn (4)):



or



The role of the alkalis is also not readily probed by computation since we have no evidence for its change in speciation. Finally, the role of water, whose presence is coupled to alkali charge-density, cannot be dismissed in any of the solid-state reactions; DAC, auto-degradation, or water-promoted degradation.

The vanadium-bound carbonate for  $\text{K}_3\text{V}(\text{O}_2)_4$  (observed for up to 300 days of monitoring), coupled with the computational studies indicating monodentate-bound carbonate is more stable than chelating carbonate, leads us to some hypotheses about the progression of carbon capture for  $\text{K}_3\text{V}(\text{O}_2)_4$ . It is possible that the vanadium is the steady-state point of  $\text{CO}_2$  capture and conversion to  $\text{CO}_3^{2-}$  via reactivity of the peroxides, and  $\text{V-OCO}_2$  binding is followed by migration of  $\text{CO}_3^{2-}$  to the alkalis, relieving steric hindrance and strain at the vanadium centre (Fig. S12†).

A final hypothesis considered for this DAC process was the presence of superoxide species in the material. Superoxide,  $\text{O}_2^-$ , is a potential decomposition product of peroxide, especially in the presence of a redox active metal such as vanadium. As discussed in the introduction; (1) a uranyl peroxide-superoxide anion was shown to evolve to a uranyl tricarbonatate via direct air capture,<sup>35</sup> and (2) potassium superoxide has been used for air purification for space travel, given its reaction with  $\text{CO}_2$  to release  $\text{O}_2$ . To identify and quantify superoxides, two approaches were employed: an experimental approach using a superoxide scavenger and Electron Paramagnetic Resonance (EPR) spectroscopy to detect unpaired electrons. First, following a reported procedure,<sup>35</sup>  $\text{K}_3\text{V}(\text{O}_2)_4$ ,  $\text{Rb}_3\text{V}(\text{O}_2)_4$ , and  $\text{Cs}_3\text{V}(\text{O}_2)_4$  were suspended in DMSO and mixed with nitro blue tetrazolium chloride (NBT). A colour change was observed immediately in the solution, suggesting the presence of superoxide bound to the vanadium or in the lattice. While there was no appreciable difference in the rate at which the reagent solutions containing the three compounds changed colour, we saw potential to finally establish a reactivity trend for the alkalis-analogues (discussed later). It is known that larger alkalis stabilize superoxide better due to poorer polarization of O–O bonds. Hence if the formation of superoxides is necessary for the  $\text{CO}_2$  capture to proceed, it could explain the increased reactivity with alkali size. To supplement these results, we also analysed both hydroxide and hydroxide-peroxide solutions for the presence of superoxide, where the latter has been reported.<sup>54</sup> The mixture of

hydroxides and NBT in DMSO without peroxide also gave a colour change indicating superoxide, suggesting this technique must be used with caution. However, water did not give a false positive, so further tests were performed in water (see ESI† for details). In the presence of superoxide, NBT is reduced to form formazan, which produces a colour change that can be monitored via UV-vis.<sup>55</sup> The aqueous NBT solutions with added KOH-peroxide, RbOH-peroxide, and CsOH-peroxide, all showed linear increase in the formazan absorption peak, and therefore increase in formation of superoxides with time (Fig. S36†). The concentration of superoxide trended  $\text{CsOH} > \text{RbOH} > \text{KOH}$ , suggesting increasing basicity increases peroxide decomposition in water, with superoxide as one of the decomposition products. We also measured TMAOH-peroxide and  $\text{NH}_4\text{OH}$ -peroxide (TMA = tetramethylammonium) because (1)  $\text{NH}_4\text{OH}$  is a weak base, and (2) TMAOH is generally considered a strong base, and (3) TMA exhibits minimal bonding/association interactions that we expect from the alkalis and provides competing stabilization/destabilization effects. Indeed, TMAOH exhibited similar superoxide presence as KOH according to the NBT test, while minimal superoxide was detected for the  $\text{NH}_4\text{OH}$  solution (Fig. S36†). UV-vis of  $\text{K}_3\text{V}(\text{O}_2)_4$ ,  $\text{Rb}_3\text{V}(\text{O}_2)_4$ , and  $\text{Cs}_3\text{V}(\text{O}_2)_4$  measured in aqueous NBT solutions also showed an increase in superoxide concentration, which plateaued over time (Fig. S37–S39† and 9A).  $\text{Rb}_3\text{V}(\text{O}_2)_4$  showed the maximum absorbance, perhaps due to residual free peroxide (Table S2†). This apparent increase in superoxide concentration with time may also be an artifact of the NBT test, and the results should be considered with caution. Additionally, dissolution of  $\text{V}(\text{O}_2)_4^{3-}$  in water leads to dissociation of the peroxide ligands and formation of orthovanadate. Nonetheless, similar results were obtained from EPR and the aqueous NBT test, discussed below. Similar tests performed on samples

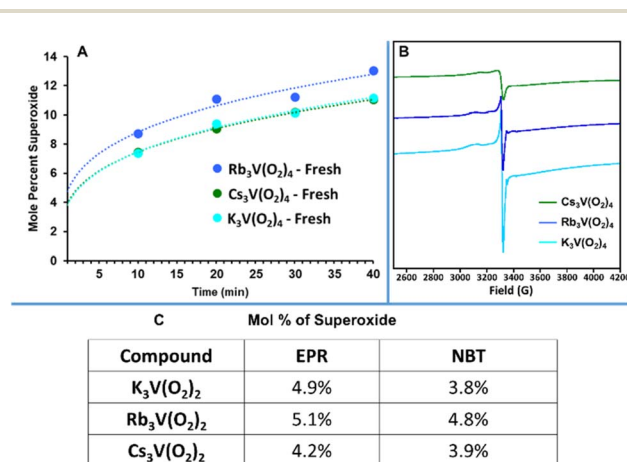


Fig. 9 Quantification of superoxide present in  $\text{K}_3\text{V}(\text{O}_2)_4$ ,  $\text{Rb}_3\text{V}(\text{O}_2)_4$ , and  $\text{Cs}_3\text{V}(\text{O}_2)_4$  by: (A) UV-vis data from NBT tests, based on a  $\text{KO}_2$  standard (Fig. S41†). The trendline of the data points was extrapolated to time  $\approx 0$  to estimate the starting concentration of superoxide in the solid material (error on data points  $\sim 5\%$ , based on triplicate measurements). (B) EPR spectra of pristine  $\text{K}_3\text{V}(\text{O}_2)_4$ ,  $\text{Rb}_3\text{V}(\text{O}_2)_4$ , and  $\text{Cs}_3\text{V}(\text{O}_2)_4$  recorded at 100 K, using a copper imidazolite standard. (C) Comparison of superoxide quantification present in pristine  $\text{K}_3\text{V}(\text{O}_2)_4$ ,  $\text{Rb}_3\text{V}(\text{O}_2)_4$ , and  $\text{Cs}_3\text{V}(\text{O}_2)_4$  from the EPR tests and NBT tests.



exposed to ambient air for three days showed little to no absorbance for  $\text{Rb}_3\text{V}(\text{O}_2)_4$ , and  $\text{Cs}_3\text{V}(\text{O}_2)_4$ , and absorbance similar to that of the fresh sample of  $\text{K}_3\text{V}(\text{O}_2)_4$  (Fig. S40†). Unlike  $\text{Rb}_3\text{V}(\text{O}_2)_4$ , and  $\text{Cs}_3\text{V}(\text{O}_2)_4$ ,  $\text{K}_3\text{V}(\text{O}_2)_4$  had retained much of its purple colour, characteristic of less decomposition. This indicates that the peroxide (and superoxide) present in the fresh  $\text{Rb}_3\text{V}(\text{O}_2)_4$ , and  $\text{Cs}_3\text{V}(\text{O}_2)_4$  materials soon decomposes upon atmospheric exposure, by reaction with  $\text{CO}_2$  and/or water. Infrared analysis supports this data (Fig. S8 and S9†). The peaks at  $2500\text{--}3200\text{ cm}^{-1}$  present in fresh  $\text{Rb}_3\text{V}(\text{O}_2)_4$  and  $\text{Cs}_3\text{V}(\text{O}_2)_4$ , consistent with either water, peroxide, or other reactive oxygen species such as superoxide,<sup>56</sup> disappear after one day. We used aqueous solutions of  $\text{KO}_2\text{-NBT}$  to create a calibration curve and quantify superoxide in the peroxovanadates (Fig. S41†). Concentration of superoxide *vs.* time in all three compounds best fits a power trend line (details in the ESI, Table S10†). Measuring the beginning of the reaction proved impossible since upon dissolution the three compounds retained their natural purple colour before the peroxide dissociates from the vanadium-centre, which interfered with the absorbance caused by the reaction with NBT. Extrapolation to the initiation of the reaction, it appears that the solids contained an initial 4–5% of superoxide (Fig. 9C). Because the NBT test are done a liquid medium (DMSO or water) that seems to promote superoxide formation (from peroxide in water and from an unknown source in DMSO), and the  $\text{CO}_2$  capture studies are performed in the solid-state, additional evidence and quantification of superoxide in the solid-state was necessary.

Additional evidence was provided by EPR of  $\text{K}_3\text{V}(\text{O}_2)_4$ ,  $\text{Rb}_3\text{V}(\text{O}_2)_4$ , and  $\text{Cs}_3\text{V}(\text{O}_2)_4$  recorded at 100 K (Fig. 9B). These spectra all show an axial lineshape associated with  $\text{V}^{5+}$ /superoxide over a broad background peak,<sup>57,58</sup> with the *g*-values tabulated in Table S11.† The *g*-values show a clear correlation with the cation. Variable temperature spectra were collected from 100 K up to 260 K for  $\text{K}_3\text{V}(\text{O}_2)_4$ ,  $\text{Rb}_3\text{V}(\text{O}_2)_4$ , and  $\text{Cs}_3\text{V}(\text{O}_2)_4$  (Fig. S42†). With increased temperature, the spectra coalesced into a symmetric peak, likely due to increased motion. If the superoxide is bonded to vanadium in the solid-state, we expect hyperfine splitting of the  $\text{O}_2^-$  signal into eight peaks from the quadrupolar vanadium nucleus ( $^{51}\text{V}$ , 99.75% natural abundance, spin 7/2).<sup>57</sup> However, not observing this in the solid-state could be due to coupling of proximal spins, typical solid-state broadening, or evidence that the superoxide is not bound to the vanadium-centres. Dilute solution measurements to determine if V-superoxide bonds are present were not possible because the compounds are only soluble in water, and the peroxide ligands dissociate in water. Attempted dissolution in DMSO did not evidence any purple coloration of the solution. However, when spun down to remove the solids, the resulting supernatant exhibited a frozen solution spectrum similar to “free” superoxide in DMSO ( $\text{KO}_2$  in DMSO is shown for comparison, Fig. S43†). This indicates that the superoxide detected was released from the solid lattice. We determined number of spins (100 K) in the peroxovanadates by double integration of the spectra and comparison with a spin standard (see ESI and Table S10† for details). This revealed that less than 5% (mole percent) superoxide is present per vanadium (Fig. 9C).

In summary, both NBT and EPR measurements confirmed that peroxide, not superoxide is the most prevalent reactive oxygen species present in the bulk material. The ~5% superoxide detected may be produced at steady-state as a decomposition product of peroxide, and then further reacts with water or with  $\text{CO}_2$ , also at steady-state.

To complement the experimental determination of the role of superoxide in DAC, we computationally explored the reaction of hypothetical triperoxosuperoxovanadate,  $[\text{V}(\text{O}_2^{2-})_3(\text{O}_2^-)]^{2/3-}$  with  $\text{CO}_2$ . The salient points are summarized here, while the details are presented in the ESI.† First,  $[\text{V}^{\text{IV}}(\text{O}_2^{2-})_3(\text{O}_2^-)]^{3-}$  instead of  $[\text{V}^{\text{V}}(\text{O}_2^{2-})_3(\text{O}_2^-)]^{2-}$  was chosen as the superoxide model compound because; (1) it is only 1–3 kcal mol<sup>-1</sup> less stable than the parent  $[\text{V}^{\text{V}}(\text{O}_2^{2-})_4]^{3-}$  (Table S12†) and therefore accessible, (2) the formation of  $[\text{V}^{\text{IV}}(\text{O}_2^{2-})_3(\text{O}_2^-)]^{3-}$  is a likely mechanism of superoxide formation from peroxide, and (3) the total charge on the anion is constant between  $[\text{V}^{\text{IV}}(\text{O}_2^{2-})_3(\text{O}_2^-)]^{3-}$  and  $[\text{V}^{\text{V}}(\text{O}_2^{2-})_4]^{3-}$ , allowing for easier direct comparison. Notably, the  $\text{CO}_2$  favourably binds the peroxide ligand, not the superoxide ligand (likely due to the longer O–O bond with more polarizable nature for peroxide), and the vanadium reverts to  $\text{V}^{\text{V}}$  in the end-product (Fig. S44†). The superoxide lowers the barrier of the rate-determining step by ~3 kcal mol<sup>-1</sup>, and the energy difference of the complete reaction of  $\text{K}_3\text{V}(\text{O}_2)_4 + \text{CO}_2 \rightarrow \text{K}_3\text{VO}(\text{CO}_3)(\text{O}_2)_2$  for the complex with superoxide *vs.* without superoxide is likewise <3 kcal mol<sup>-1</sup>. In summary, computation suggests that vanadium-bound peroxide and superoxide could both be important in the observed  $\text{CO}_2$  DAC capture reaction.

Practical Aspects of  $\text{V}(\text{O}_2)_4^{3-}$  DAC molecules. Compiled in Table S13† is performance of a variety of leading materials for DAC (including the peroxovanadates reported here) or carbon capture in  $\text{CO}_2$ -rich environments, selected and summarized from two recent reviews.<sup>59,60</sup> The range is 0.4 to 17 mmol g<sup>-1</sup>; and  $\text{K}_3\text{V}(\text{O}_2)_4$  has similar  $\text{CO}_2$  capture capacity (~4.9 mmol g<sup>-1</sup>) as carbon-based adsorbents.

To further put the peroxovanadate performance into context, the DAC behaviour of sodium peroxide ( $\text{Na}_2\text{O}_2$ ) was demonstrated. Total DAC capacity reached a maximum of 9.6 carbon weight% after 3 days (Fig. S45†). Both rate and capacity are greater than that of  $\text{A}_3\text{V}(\text{O}_2)_4$ . On the other hand, TGA-MS analyses shows no  $\text{CO}_2$  release up to 900 °C (Fig. S46 and S47†). This is due to the formation of sodium carbonate, which has a high decomposition temperature. Both  $\text{Na}_2\text{O}_2$  and its decomposition products are hygroscopic, evidenced by the increased hydrogen weight% and higher weight loss at lower temperatures, due to water present in the solid (Fig. S47†).

Finally, recovering the starting  $\text{Na}_2\text{O}_2$  is not trivial since its commercial synthesis is based on the oxygenation of sodium oxide,<sup>61</sup> which would be hard to obtain from sodium carbonate due its aforementioned high thermal stability. Concerning reusability of the peroxovanadate materials for DAC; the peroxovanadate salts are easily dissolved in water at any stage of the process from  $(\text{CO}_3)\text{VO}(\text{O}_2)_2^{3-}$  to the end product of  $\text{VO}_3$ -plus  $\text{HCO}_3^-$ . Dissolution in  $\text{KOH}$  plus  $\text{H}_2\text{O}_2$  followed by precipitation *via* addition of alcohol, per the synthesis regenerates the  $\text{A}_3\text{V}(\text{O}_2)_4$  materials. We have performed this reaction to



demonstrate its feasibility. Since the synthesis is robust and single step; the source of vanadium and the  $\text{CO}_3^{2-}$  present in the reprocessing reaction does not alter the product. The challenge to implementing this in an industrial setting is the cooling required for maximum yield of product.

## Conclusions

In this study, we have experimentally and computationally demonstrated that reactive alkali vanadium peroxide compounds (containing up to 5% adventitious superoxide) undergo stabilization *via* direct air capture of  $\text{CO}_2$ , forming  $\text{VO}(\text{CO}_3)(\text{O}_2)_2^{3-}$ . The re-release of vanadium-bound  $\text{CO}_2$  by heating (*i.e.* regeneration step) is simultaneous with vanadium-bound peroxide, with a low release temperature of 250 °C. Alkali carbonate and alkali bicarbonate are also products of carbon capture, especially with long aging times. However, the K-analogue shows evidence that the carbon capture is always *via* the vanadium-centre, followed by migration of the carbonate to the alkali. In addition, we provide evidence that vanadium catalyses lower temperature release of alkali-bound carbonate (520 °C *vs.* 850 °C). These studies all show that the alkalis and vanadium are synergistic in both capture and release of atmospheric  $\text{CO}_2$ . The reactivity of vanadium peroxides increases with alkali counteraction size ( $\text{Cs} > \text{Rb} > \text{K}$ , as shown here); yet the underlying role of the alkalis is still poorly understood. Ongoing experiments and computation include detailed analysis of the Li and Na analogues of vanadium tetraperoxide to better understand the complex role of the alkalis. We are also investigating the DAC performance of the  $d^0$  transition metal peroxide neighbours including Nb, Ta, Mo, W and Ti by computation and experiment. Emerging trends in reactivity will inform and enable the use of a range of peroxide-functionalized inorganic oxides in DAC.

## Data availability

Crystallographic data for compounds 2 and 3 have been deposited at the CCDC under 2264013 and 2264012, respectively, and can be obtained from <https://www.ccdc.cam.ac.uk/structures/>. Computational input files and structures can be accessed through the public GitHub repository ([https://github.com/tjz21/DAC\\_peroxovanadates](https://github.com/tjz21/DAC_peroxovanadates)) or through Zenodo (<https://doi.org/10.5281/zenodo.10266510>).

## Author contributions

EGR methodology, investigation, visualization and writing; ZM methodology and investigation, JSH investigation, visualization and writing, TL investigation, KB investigation, EDW investigation and writing, CRS investigation, TJZ writing and supervision, MN conceptualization, supervision, writing and visualization, project administration, funding acquisition.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 K. O. Yoro and M. O. Daramola, in *Advances in Carbon Capture*, ed. M. R. Rahimpour, M. Farsi and M. A. Makarem, Woodhead Publishing, 2020, pp. 3–28, DOI: [10.1016/B978-0-12-819657-1.00001-3](https://doi.org/10.1016/B978-0-12-819657-1.00001-3).
- 2 D. Shindell and C. J. Smith, Climate and air-quality benefits of a realistic phase-out of fossil fuels, *Nature*, 2019, **573**, 408–411.
- 3 T. Wilberforce, A. Olabi, E. T. Sayed, K. Elsaid and M. A. Abdelkareem, Progress in carbon capture technologies, *Sci. Total Environ.*, 2021, **761**, 143203.
- 4 A. Gambhir and M. Tavoni, Direct Air Carbon Capture and Sequestration: How It Works and How It Could Contribute to Climate-Change Mitigation, *One Earth*, 2019, **1**, 405–409.
- 5 S. Deutz and A. Bardow, Life-cycle assessment of an industrial direct air capture process based on temperature-vacuum swing adsorption, *Nat. Energy*, 2021, **6**, 203–213.
- 6 P. Gabrielli, M. Gazzani and M. Mazzotti, The role of carbon capture and utilization, carbon capture and storage, and biomass to enable a net-zero- $\text{CO}_2$  emissions chemical industry, *Ind. Eng. Chem.*, 2020, **59**, 7033–7045.
- 7 X. Shi, Y. Lin and X. Chen, Development of sorbent materials for direct air capture of  $\text{CO}_2$ , *MRS Bull.*, 2022, **47**, 405–415.
- 8 R. W. J. Edwards and M. A. Celia, Infrastructure to enable deployment of carbon capture, utilization, and storage in the United States, *Proc. Natl. Acad. Sci. U.S.A.*, 2018, **115**, E8815–E8824.
- 9 N. McCann, D. Phan, X. Wang, W. Conway, R. Burns, M. Attalla, G. Puxty and M. Maeder, Kinetics and Mechanism of Carbamate Formation from  $\text{CO}_2(\text{aq})$ , Carbonate Species, and Monoethanolamine in Aqueous Solution, *J. Phys. Chem. A*, 2009, **113**, 5022–5029.
- 10 F. O. Ochedi, J. Yu, H. Yu, Y. Liu and A. Hussain, Carbon dioxide capture using liquid absorption methods: a review, *Environ. Chem. Lett.*, 2021, **19**, 77–109.
- 11 Y. Hu, Y. Guo, J. Sun, H. Li and W. Liu, Progress in MgO sorbents for cyclic  $\text{CO}_2$  capture: a comprehensive review, *J. Mater. Chem. A*, 2019, **7**, 20103–20120.
- 12 P. E. Sánchez Jiménez, A. Perejón, M. Benítez Guerrero, J. M. Valverde, C. Ortiz and L. A. Pérez Maqueda, High-performance and low-cost macroporous calcium oxide based materials for thermochemical energy storage in concentrated solar power plants, *Appl. Energy*, 2019, **235**, 543–552.



- 13 N. McQueen, P. Kelemen, G. Dipple, P. Renforth and J. Wilcox, Ambient weathering of magnesium oxide for CO<sub>2</sub> removal from air, *Nat. Commun.*, 2020, **11**, 3299.
- 14 M. Samari, F. Ridha, V. Manovic, A. Macchi and E. J. Anthony, Direct capture of carbon dioxide from air via lime-based sorbents, *Mitig. Adapt. Strateg. Glob. Chang.*, 2020, **25**, 25–41.
- 15 D. Fu, Y. Park and M. E. Davis, Zinc Containing Small-Pore Zeolites for Capture of Low Concentration Carbon Dioxide, *Angew. Chem., Int. Ed.*, 2022, **61**, e202112916.
- 16 D. Fu, Y. Park and M. E. Davis, Confinement effects facilitate low-concentration carbon dioxide capture with zeolites, *Proc. Natl. Acad. Sci. U.S.A.*, 2022, **119**, e2211544119.
- 17 R. Gonzalez-Olmos, A. Gutierrez-Ortega, J. Sempere and R. Nomen, Zeolite versus carbon adsorbents in carbon capture: A comparison from an operational and life cycle perspective, *J. CO<sub>2</sub> Util.*, 2022, **55**, 101791.
- 18 C. C. Turvey, S. A. Wilson, J. L. Hamilton, A. W. Tait, J. McCutcheon, A. Beinlich, S. J. Fallon, G. M. Dipple and G. Southam, Hydrotalcites and hydrated Mg-carbonates as carbon sinks in serpentinite mineral wastes from the Woodsreef chrysotile mine, New South Wales, Australia: Controls on carbonate mineralogy and efficiency of CO<sub>2</sub> air capture in mine tailings, *Int. J. Greenhouse Gas Control*, 2018, **79**, 38–60.
- 19 D. Suescum-Morales, J. R. Jiménez and J. M. Fernández-Rodríguez, Review of the Application of Hydrotalcite as CO<sub>2</sub> Sinks for Climate Change Mitigation, *Chem. Eng.*, 2022, **6**, 50.
- 20 S. A. Didas, S. Choi, W. Chaikittisilp and C. W. Jones, Amine-Oxide Hybrid Materials for CO<sub>2</sub> Capture from Ambient Air, *Acc. Chem. Res.*, 2015, **48**, 2680–2687.
- 21 A. Cherevotan, J. Raj and S. C. Peter, An overview of porous silica immobilized amines for direct air CO<sub>2</sub> capture, *J. Mater. Chem. A*, 2021, **9**, 27271–27303.
- 22 M. Ding, R. W. Flaig, H.-L. Jiang and O. M. Yaghi, Carbon capture and conversion using metal-organic frameworks and MOF-based materials, *Chem. Soc. Rev.*, 2019, **48**, 2783–2828.
- 23 C. G. Piscopo and S. Loebbecke, Strategies to Enhance Carbon Dioxide Capture in, *Met.-Org. Frameworks*, 2020, **85**, 538–547.
- 24 X. Shi, H. Xiao, H. Azarabadi, J. Song, X. Wu, X. Chen and K. S. Lackner, Sorbents for the Direct Capture of CO<sub>2</sub> from Ambient Air, *Angew. Chem., Int. Ed.*, 2020, **59**, 6984–7006.
- 25 H. Jakob, S. Leininger, T. Lehmann, S. Jacobi and S. Gutewort, Peroxo Compounds, Inorganic, in *Ullmann's Encyclopedia of Industrial Chemistry*, 2007, DOI: [10.1002/14356007.a19\\_177.pub2](https://doi.org/10.1002/14356007.a19_177.pub2).
- 26 M. Dakanali, E. T. Kefalas, C. P. Raptopoulou, A. Terzis, G. Voyiatzis, I. Kyrikou, T. Mavromoustakos and A. Salifoglou, A New Dinuclear Ti(IV)–Peroxo–Citrate Complex from Aqueous Solutions. Synthetic, Structural, and Spectroscopic Studies in Relevance to Aqueous Titanium(IV)–Peroxo–Citrate Speciation, *Inorg. Chem.*, 2003, **42**, 4632–4639.
- 27 M. Grzywa, M. Różycka and W. Łasocha, Crystal structure of potassium tetraperoxomolybdate (VI) K<sub>2</sub>[Mo(O<sub>2</sub>)<sub>4</sub>], *Powder Diffr.*, 2005, **20**, 203–206.
- 28 M. Grzywa, W. Nitek and W. Łasocha, Synthesis, characterization and crystal structures of lithium tetraperoxomolybdate (VI) tetrahydrate Li<sub>2</sub>[Mo(O<sub>2</sub>)<sub>4</sub>]·4H<sub>2</sub>O and lithium tetraperoxotungstate (VI) tetrahydrate Li<sub>2</sub>[W(O<sub>2</sub>)<sub>4</sub>]·4H<sub>2</sub>O, *J. Mol. Struct.*, 2007, **828**, 111–115.
- 29 H. Haeuseler, M. Wagener and H. Müller, Preparation, Crystal Structure, and Vibrational Spectra of Rubidiumperoxoniobate (Rb<sub>3</sub>NbO<sub>8</sub>), *Z. Naturforsch. B*, 1997, **52**, 1082–1086.
- 30 G. Wehrum and R. Hoppe, On the constitution of peroxotantalates (V) with alkali metals: on the structure of K<sub>3</sub>Ta(O<sub>2</sub>)<sub>4</sub>, *Z. Anorg. Allg. Chem.*, 1993, 619.
- 31 R. Stomberg and C. Brosset, The Crystal Structure of Potassium Perchromate, *Acta Chem. Scand.*, 1960, **14**, 2.
- 32 T.-J. Won, C. L. Barnes, E. O. Schlemper and R. C. Thompson, Two crystal structures featuring the tetraperoxovanadate (V) anion and a brief reinvestigation of peroxovanadate equilibria in neutral and basic solutions, *Inorg. Chem.*, 1995, **34**, 4499–4503.
- 33 Z. W. Yu, P. A. Jansson, B. I. Posner, U. Smith and J. W. Eriksson, Peroxovanadate and insulin action in adipocytes from NIDDM patients. Evidence against a primary defect in tyrosine phosphorylation, *Diabetologia*, 1997, **40**, 1197–1203.
- 34 V. Conte and B. Floris, Vanadium and molybdenum peroxides: synthesis and catalytic activity in oxidation reactions, *Dalton Trans.*, 2011, **40**, 1419–1436.
- 35 D. V. Kravchuk, N. N. Dahlen, S. J. Kruse, C. D. Malliakas, P. M. Shand and T. Z. Forbes, Isolation and reactivity of uranyl superoxide, *Angew. Chem., Int. Ed.*, 2021, **60**, 15041–15048.
- 36 D. V. Kravchuk and T. Z. Forbes, Thermodynamics and Chemical Behavior of Uranyl Superoxide at Elevated Temperatures, *ACS Mater. Au*, 2022, **2**, 33–44.
- 37 P. Schwendt, J. Tatiersky, L. Krivosudský and M. Šimuneková, Peroxido complexes of vanadium, *Coord. Chem. Rev.*, 2016, **318**, 135–157.
- 38 P. Schwendt and D. Uskert, Thermal-Decomposition of Potassium Mu-Oxo- and Mu-Hydroxo-Bis(Oxo-Diperoxovanadate), *Chem. Zvesti*, 1981, **35**, 229–233.
- 39 G. Arrambide, J. Rivadeneira, S. Etcheverry, B. Parajón-Costa, D. Gambino and E. Baran, Spectroscopic Behavior and Biological Activity of K<sub>3</sub>[VO(O<sub>2</sub>)<sub>2</sub>CO<sub>3</sub>]·H<sub>2</sub>O, *Biol. Trace Elem. Res.*, 2009, **132**, 176–183.
- 40 G. Zou, Z. Lin, H. Zeng, H. Jo, S.-J. Lim, T.-S. You and K. M. Ok, Cs<sub>3</sub>VO(O<sub>2</sub>)<sub>2</sub>CO<sub>3</sub>: an exceptionally thermostable carbonatoperoxovanadate with an extremely large second-harmonic generation response, *Chem. Sci.*, 2018, **9**, 8957–8961.
- 41 I. Knopf, T. Ono, M. Temprado, D. Tofan and C. C. Cummins, Uptake of one and two molecules of CO<sub>2</sub> by the molybdate dianion: a soluble, molecular oxide model system for carbon dioxide fixation, *Chem. Sci.*, 2014, **5**, 1772–1776.



- 42 M. Grzywa and W. Łasocha, Crystal structure of potassium and ammonium tetraperoxovanadates (V), *Z. Kristallogr. - Cryst. Mater.*, 2007, **222**, 95–98.
- 43 A. Arteaga, L. Zhang, S. Hickam, M. Dembowski, P. C. Burns and M. Nyman, Uranyl–Peroxide Capsule Self-Assembly in Slow Motion, *Eurasian J. Chem.*, 2019, **25**, 6087–6091.
- 44 D. V. Kravchuk, N. N. Dahlen, S. J. Kruse, C. D. Malliakas, P. M. Shand and T. Z. Forbes, Isolation and reactivity of uranyl superoxide, *Angew. Chem., Int. Ed.*, 2021, **60**, 15041–15048.
- 45 M. Grzywa, W. Łasocha and D. Rutkowska-Żbik, Structural investigation of tetraperoxo complexes of Mo (VI) and W (VI): X-ray and theoretical studies, *J. Solid State Chem.*, 2009, **182**, 973–982.
- 46 K. Coenen, F. Gallucci, B. Mezari, E. Hensen and M. van Sint Annaland, An in-situ IR study on the adsorption of CO<sub>2</sub> and H<sub>2</sub>O on hydrotalcites, *J. CO<sub>2</sub> Util.*, 2018, **24**, 228–239.
- 47 S. Clark, M. Segall, C. Pickard, P. Hasnip, M. Probert, K. Refson and M. C. Payne, First Principles Methods Using CASTEP, *Z. Kristallogr. - Cryst. Mater.*, 2005, **220**, 567.
- 48 D. Bayot, M. Devillers and D. Peeters, Vibrational spectra of eight-coordinate niobium and tantalum complexes with peroxo ligands: A theoretical simulation, *Eur. J. Inorg. Chem.*, 2005, **2005**, 4118–4123.
- 49 E. Mutlu, T. Cristy, S. W. Graves, M. J. Hooth and S. Waidyanatha, Characterization of aqueous formulations of tetra- and pentavalent forms of vanadium in support of test article selection in toxicology studies, *Environ. Sci. Pollut. Res.*, 2017, **24**, 405–416.
- 50 G. W. T. M. J. Frisch, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery Jr, J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman, and D. J. Fox, *Gaussian 16, Revision A. 03*, Gaussian, Inc., Wallingford CT, 2016, 3.
- 51 W. Ma, Y. Qiao, N. Theyssen, Q. Zhou, D. Li, B. Ding, D. Wang and Z. Hou, A mononuclear tantalum catalyst with a peroxocarbonate ligand for olefin epoxidation in compressed CO<sub>2</sub>, *Catal. Sci. Technol.*, 2019, **9**, 1621–1630.
- 52 M. Yamashita, K. Goto and T. Kawashima, Fixation of both O<sub>2</sub> and CO<sub>2</sub> from air by a crystalline palladium complex bearing N-heterocyclic carbene ligands, *J. Phys. Chem. B*, 2005, **127**, 7294–7295.
- 53 T. A. Manz and B. Yang, Selective oxidation passing through η<sup>3</sup>-ozone intermediates: applications to direct propene epoxidation using molecular oxygen oxidant, *RSC Adv.*, 2014, **4**, 27755–27774.
- 54 D. Wang, J. Zou, H. Cai, Y. Huang, F. Li and Q. Cheng, Effective degradation of Orange G and Rhodamine B by alkali-activated hydrogen peroxide: roles of HO<sub>2</sub><sup>–</sup> and O<sub>2</sub>, *Environ. Sci. Pollut. Res.*, 2019, **26**, 1445–1454.
- 55 B. H. Bielski, G. G. Shiue and S. Bajuk, Reduction of nitro blue tetrazolium by CO<sub>2</sub>- and O<sub>2</sub>-radicals, *J. Phys. Chem.*, 1980, **84**, 830–833.
- 56 P. A. Giguere and K. B. Harvey, An Infrared Study of Hydrogen Bonding in Solid H<sub>2</sub>O<sub>2</sub> and H<sub>2</sub>O–H<sub>2</sub>O<sub>2</sub> Mixtures, *J. Mol. Spectrosc.*, 1959, **3**, 36–45.
- 57 V. N. Shetti, M. J. Rani, D. Srinivas and P. Ratnasamy, Chemoselective Alkane Oxidation by Superoxo–Vanadium(V) in Vanadosilicate Molecular Sieves, *J. Am. Chem. Soc.*, 2006, **110**, 677–679.
- 58 K. Sobańska, A. Krasowska, T. Mazur, K. Podolska-Serafin, P. Pietrzyk and Z. Sojka, Diagnostic Features of EPR Spectra of Superoxide Intermediates on Catalytic Surfaces and Molecular Interpretation of Their g and A Tensors, *Top. Catal.*, 2015, **58**, 796–810.
- 59 D. Panda, V. Kulkarni and S. K. Singh, Evaluation of amine-based solid adsorbents for direct air capture: a critical review, *React. Chem. Eng.*, 2023, **8**, 10–40.
- 60 W. Gao, S. Liang, R. Wang, Q. Jiang, Y. Zhang, Q. Zheng, B. Xie, C. Y. Toe, X. Zhu, J. Wang, L. Huang, Y. Gao, Z. Wang, C. Jo, Q. Wang, L. Wang, Y. Liu, B. Louis, J. Scott, A.-C. Roger, R. Amal, H. He and S.-E. Park, Industrial carbon dioxide capture and utilization: state of the art and future challenges, *Chem. Soc. Rev.*, 2020, **49**, 8584–8686.
- 61 E. Dönges, *Lithium and Sodium Peroxides*, Academic Press, NY, 1963.

