


 Cite this: *New J. Chem.*, 2023, 47, 18178

 Received 15th July 2023,  
 Accepted 5th September 2023

DOI: 10.1039/d3nj03307d

rsc.li/njc

## A complex of cobalamin with an organic peroxide†

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Aquacobalamin reacts with *m*-chloroperoxybenzoic acid (MCPBA) to form a Co(III)-peroxo complex, as supported by UV-vis, NMR and resonance Raman spectra complemented by mass spectra and density functional calculations. At high MCPBA concentrations, degradation of cobalamin is observed as previously described.

### Introduction

Peroxide complexes of biological transition metal centres are in some cases reactive intermediates in complex reaction mechanisms (e.g., in peroxidases, cytochromes P450, non-heme iron/manganese/copper monooxygenases, iron and copper bleomycin, vanadium-containing peroxidases and others).<sup>1–4</sup> Stable peroxo complexes of biological transition metal centres are also known, such as with Co(III) cobalamin (Cbl(III)) or Co(II) bleomycin.<sup>5,6</sup>

Until recently, the reactivity of cobalamin with oxidizing agents has been confined to processes where, especially with strong oxidizing agents, the corrin ring is covalently modified by oxygenation or halogenation, or where Co(I) or Co(II) are oxidized to Co(III) in an outer-sphere manner.<sup>5,7–13</sup> Outer-sphere oxidants such as hexachloroiridate and peroxosulfate

have no effect on Cbl(III) (data not shown), while hydrogen peroxide does form a stable and reversible complex, assigned as Co(III)-hydroperoxo based mostly on UV-vis and NMR spectra complemented by density functional (DFT) calculations. The UV-vis spectra (and ensuing properties, such as resonance Raman spectra) of cobalamin are notoriously insensitive to changes in the exogenous axial ligand; for instance, the maxima of aquacobalamin differ in position from those of hydroxocobalamin by less than 10 nm – and the differences between hydroxo- and hydroperoxocobalamin are even smaller. By contrast, the aromatic region of the <sup>1</sup>H-NMR spectra does offer distinctively different signatures for each of these cobalamin complexes.<sup>5</sup> More recently, another example of peroxide-cobalamin interaction has been described – with *m*-chloroperoxybenzoic acid (MCPBA).<sup>14</sup> In this case Cbl was reported to act as a catalyst for MCPBA decomposition and subsequent oxidation of organic substrates. While such pseudoperoxidasic cycles have been reported previously for Cbl with hydrogen peroxide or *tert*-butyl hydroperoxide (*t*BuOOH) instead of MCPBA, in all of these previous instances activation of the peroxo bond was achieved *via* reaction with Cbl(II) rather than Cbl(III),<sup>8,9</sup> one may expect a similar mechanism in the MCPBA case. A second, more recent example of oxygen–oxygen bond cleavage by aquacobalamin has been reported for peroxomonosulfate.<sup>15</sup> In a related system – heptamethyl cobyrinate – a Co(III) adduct with MCPBA was detected at low temperature using UV-vis, IR, ESR, and ESI-MS.<sup>14</sup> Somewhat related Co-peroxo complexes of geminal diols are as stable as to have been described by X-ray crystallography.<sup>16</sup>

Reported here are data showing that while a representative organic peroxide (*t*BuOOH) brings no significant changes to the NMR and UV-vis spectra of Cbl(III), MCPBA does yield a detectable complex, identifiable as such in UV-vis and NMR spectra corroborated with DFT calculations.

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† Electronic supplementary information (ESI) available: UV-vis spectra of Cbl + *t*BuOOH mixtures, titration curves for aquaCbl reacting with MCPBA at pH 7, monitored at 490 nm and 525 nm, representative spectra of species A and B at pH 7, room temperature, resulting from the fitting of the stopped-flow data upon the reaction of aquaCbl, absorbance time course from stopped-flow UV-vis experiments at varying MCPBA concentrations monitored at 525 nm, DFT-optimized geometries for Cbl-MCPBA (A) and Cbl-*t*BuOOH (B) complexes, DFT-derived data of Cbl-MCPBA and Cbl-*t*BuOOH models, TD-DFT-derived wavelengths (nm) and oscillator strengths (OS) for Cbl-MCPBA- and Cbl-*t*BuOO- models, main orbitals responsible for the main contributors to the two main bands in the TD-DFT spectra in Cbl complexes with MCPBA and *t*BuOOH, details of resonance Raman spectra of Cbl in the presence or absence of hydrogen peroxide or MCPBA, mass spectra of Cbl with MCPBA at 4 and at 30 minutes, and Cbl structure with atom labelling employed for <sup>1</sup>H-NMR assignments. See DOI: <https://doi.org/10.1039/d3nj03307d>



## Materials and methods

Hydroxocobalamin hydrochloride (HOCbl,  $\geq 98\%$ ) was obtained from Sigma–Aldrich (Munich, Germany) and used as received and hydrogen peroxide 30% was obtained from ChemPUR. MCPBA was from Sigma–Aldrich (Munich, Germany) and was employed in the form of a 2 M stock solution in acetone. *t*BuOOH was from Sigma–Aldrich (Munich, Germany), 70% solution in H<sub>2</sub>O. The buffer solution was 50 mM phosphate, pH 7, unless otherwise specified.

UV-vis, NMR, resonance Raman stopped-flow and DFT data were obtained using a methodology previously described for the reaction of hydrogen peroxide with aquaCbl, as detailed below.<sup>5</sup>

UV-vis spectra were performed on a Cary 50 UV-vis spectrophotometer (Varian, Inc., Foster City, CA, USA). Stopped-flow spectra were collected on a Biologic SFM-300 system equipped with three syringes and sequential mixing, with a high-speed diode array detector. The stopped-flow data were processed with the SPECFIT32 software package (BioLogic system, Claix, France) using a single process for fitting. Time course graphs are listed in the Figures at 435 nm, specific for the peroxide adduct.

The NMR spectra were recorded immediately after preparing the samples, at 22 °C unless otherwise stated, diluting the sample (at concentrations indicated in text and Figure legends) with D<sub>2</sub>O (1 : 1 volume ratio), on a 400 MHz Bruker instrument. The NMR titration of aquaCbl with MCPBA was performed at 20 °C on a 500 MHz Bruker Avance spectrometer. The solvent of choice was pH 7 Britton–Robinson universal buffer prepared in D<sub>2</sub>O. 1 mM aquaCbl was titrated with incremental amounts of MCPBA to final concentrations, as indicated in Fig. S9 (ESI<sup>†</sup>) caption. A water-suppression pulse sequence was used for these measurements.

Raman spectra were measured at 22 °C immediately after preparing the samples, on a Renishaw inVia Raman spectrometer coupled with a Leica microscope. 10  $\mu$ l of the sample was dropped on a microscope slide covered with aluminum foil. The 532 nm laser line with a power of 50 mW was focused on the sample by using a 5X objective. Each spectrum is represented as an average of 4 accumulations, 4 seconds each.

High-resolution mass spectra (HRMS) were recorded on an LTQ ORBITRAP XL mass spectrometer (ThermoScientific) using positive electrospray ionization. The instrument was externally calibrated. The samples were prepared at room temperature (22 °C) and then inserted into the instrument at times indicated in the Figure legends. The following conditions were used: source voltage, 3.2 kV; sheath and auxiliary gas flow, 8 and 5 arbitrary units, respectively; vaporizer temperature 50 °C, capillary temperature 275 °C, analyzer temperature 26 °C; capillary voltage, 28 V; tube lens voltage, +110 V. The number of microscans was set to three.

For DFT calculations, the Gaussian09 software package<sup>17</sup> was employed following the methodology previously tested and described for related Cbl complexes.<sup>5</sup> The Cbl models were truncated, with the lateral substituents on the corrin as well as

the methyl groups on the benzimidazole replaced by hydrogen. Gas-phase geometries and frequency analyses were computed with the aid of the B3LYP<sup>18,19</sup> functional at the def2-SV(P)<sup>20</sup> double-zeta basis set level. Long-range interactions were accounted for by the use of Grimme's D3 dispersion correction.<sup>20</sup> TD-DFT derived<sup>21</sup> UV-Vis spectra were computed in the C-PCM solvent continuum adapted for an aqueous environment.<sup>22</sup> For the latter property the B3PW91<sup>18,23</sup> functional outperformed B3LYP and was subsequently employed. Data for the lowest-spin states are shown for each model. In terms of methodology choice for DFT calculations, previous studies from us,<sup>5,24–28</sup> but more importantly from Brunold and especially from Kozłowski on related systems have each indicated advantages for a functional or another.<sup>29–42</sup> We have, more in general, shown that even within the same class of compounds the performance of a functional can vary widely depending on the property sought, or on apparently simple changes in ligands (*e.g.*, axial ligation in heme complexes).<sup>43–46</sup> The methodology employed here was selected for its ability to best mimic trends in UV-vis spectra as described in our previous study on hydroperoxocobalamin, thus also allowing consistency between the two sets of data. In terms of general structural features, the cobalamin-peroxo adducts (unlike, *e.g.*, the case of Cbl(i)) offer no particular challenge – so that the choice of functional should affect the results at the level of detail (which, nevertheless, in predicting trends in UV-vis maxima of Cbl adducts, is essential).

## Results and discussion

As shown in Fig. 1, upon reaction with MCPBA the UV-vis spectrum of aquacobalamin (H<sub>2</sub>O Cbl<sup>+</sup>) undergoes bathochromic shifts reminiscent of the hydroperoxocobalamin adduct.<sup>5</sup> Isosbestic points at 461 nm, 340 and 367 nm suggest the formation of a single new product, which can in principle be assigned as a peroxoacid(MCPBA)-Cbl(III) complex analogous to the previously-reported cobalamin(III)-hydroperoxo complex. A 0.2 mM affinity for MCPBA binding to Cbl(III) may be deduced from the data in Fig. 1 (based on the absorbance changes at 351 nm; see also Fig. S2 (ESI<sup>†</sup>) for titration curves at 490 and 525 nm). At larger MCPBA concentrations (above 5–10 mM) a slight decrease in intensity in the visible region of the spectrum is observed, consistent with previous observations of a chemical reaction with degradation of MCPBA and generation of reactive redox species.<sup>14</sup>

No changes in the shape of the UV-vis spectrum of H<sub>2</sub>O Cbl<sup>+</sup> were observed with *t*BuOOH (ESI<sup>†</sup> Fig. S1). If indeed the Cbl complexes with hydrogen peroxide and with MCPBA entail a deprotonated peroxo group, and then the p*K*<sub>a</sub> values may be responsible for the fact that *t*BuOOH does not form such a complex: the p*K*<sub>a</sub> of *t*BuOOH (12.7) is one unit higher than that of H<sub>2</sub>O<sub>2</sub>, and distinctly higher than that of MCPBA, too (7.6). At larger *t*BuOOH concentrations ( $\geq 10$  mM), the spectrum of Cbl does decay slowly – in line with the known intrinsic instability of *t*BuOOH and possibly with catalysis from Cbl;



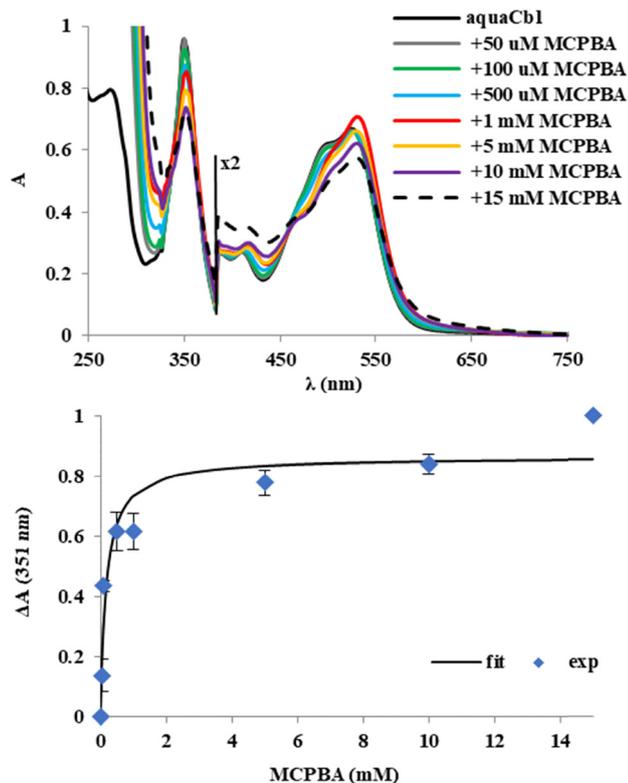


Fig. 1 UV-vis spectra of 50  $\mu\text{M}$  aquacobalamin treated with MCPBA (upper panel) and the titration curve at pH 7 monitored at 351 nm (lower panel).

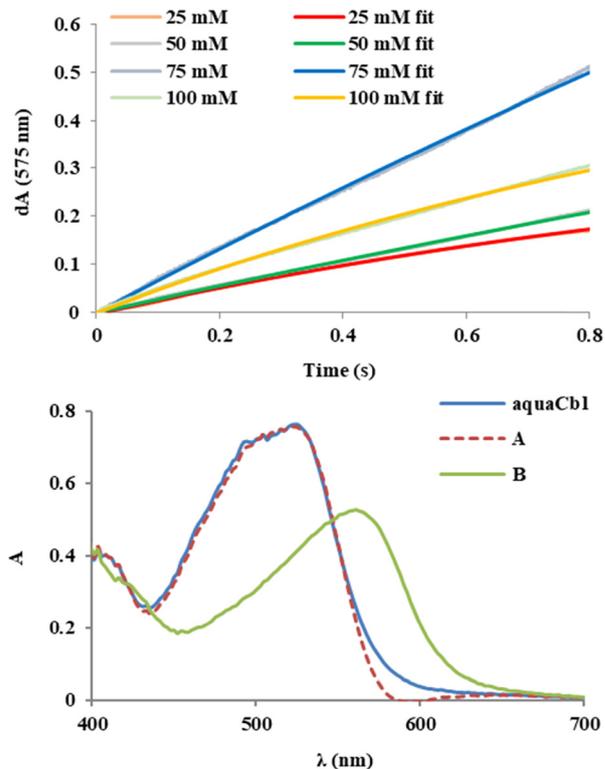


Fig. 2 Upper panel: absorbance time courses from stopped-flow UV-vis experiments monitoring the reaction of 0.3 mM aquacobalamin with varying MCPBA concentrations at 575 nm. Lower panel: representative spectra of species A and B at pH 7, resulted from fitting of the stopped-flow data 50 mM MCPBA. See also Fig. S3 and S4 (ESI $^\dagger$ ).

however, evidence for a detectable *t*BuOOH-Cbl complex as an intermediate species in such reactions, if any, has so far not been identified.

The kinetics of the aquacobalamin + MCPBA reaction may be followed with stopped-flow UV-vis spectroscopy (*cf.* Fig. 2 and ESI $^\dagger$ , Fig. S3 and S4) at MCPBA concentrations distinctly higher than those in Fig. 1. Between 25 and 75 mM MCPBA, the experimentally-observed changes in absorbance increase with the MCPBA concentration as expected. However, this range is too narrow to allow a meaningful estimation of a rate constant, and the trend is reversed at 100 mM MCPBA. At all concentrations, the data may be fitted with a single process, where species A (*cf.* Fig. 2 and Fig. S3 and S4, ESI $^\dagger$ ) is essentially identical to  $\text{H}_2\text{OCbl}^+$  and species B exhibits a maximum at 560 nm, characteristic of partially degraded corrin.<sup>5,11,12,47,48</sup> This is consistent with the behavior on Cbl in the presence of such high MCPBA concentrations more extensively described elsewhere, highlighting the occurrence of peroxide activation and ensuing oxidative activity.<sup>14</sup>

Fig. 3 shows NMR spectra collected for  $\text{H}_2\text{OCbl}^+$  mixed with *t*BuOOH and with MCPBA, respectively (with further details shown in Fig. 4, and proton labelling shown in Fig. S9, ESI $^\dagger$ ). The spectrum of the Cbl-*t*BuOOH mixture retains almost all of the features of  $\text{H}_2\text{OCbl}^+$ ; slight shifts are seen in the B7, R1 and C10 signals – possibly due to non-covalent interaction with *t*BuOOH, but decidedly not to *t*BuOO- coordination. A Cbl(III)

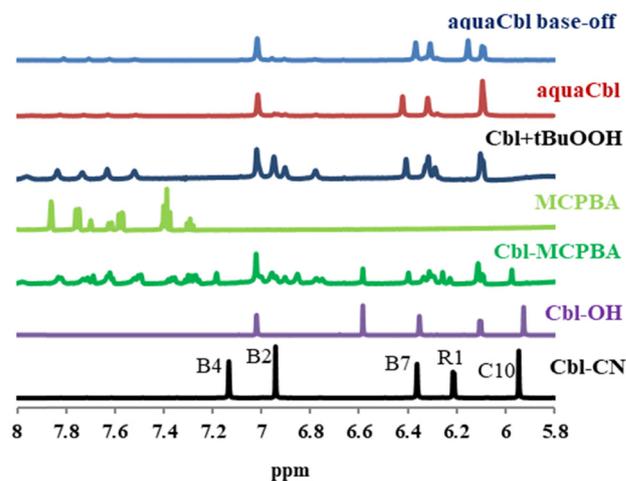


Fig. 3  $^1\text{H}$ -NMR spectra of Cbl-peroxy mixtures and of reference complexes. Peaks are assigned based as previously described.<sup>5</sup> Conditions: 5 mM Cbl(III) (aqua/hydroxo or cyano, as indicated), 5 mM ligand (MCPBA, *t*BuOOH), 50 mM phosphate buffer pH 7 for all samples except aqua base-off (pH 3, dimethylbenzimidazole deligated) and Cbl-OH (pH 10).

complex with the neutral *t*BuOOH form is less likely but cannot be entirely excluded, considering the similar coordination properties of the *t*BuOOH hydroxyl *vs.* water, and the fact that



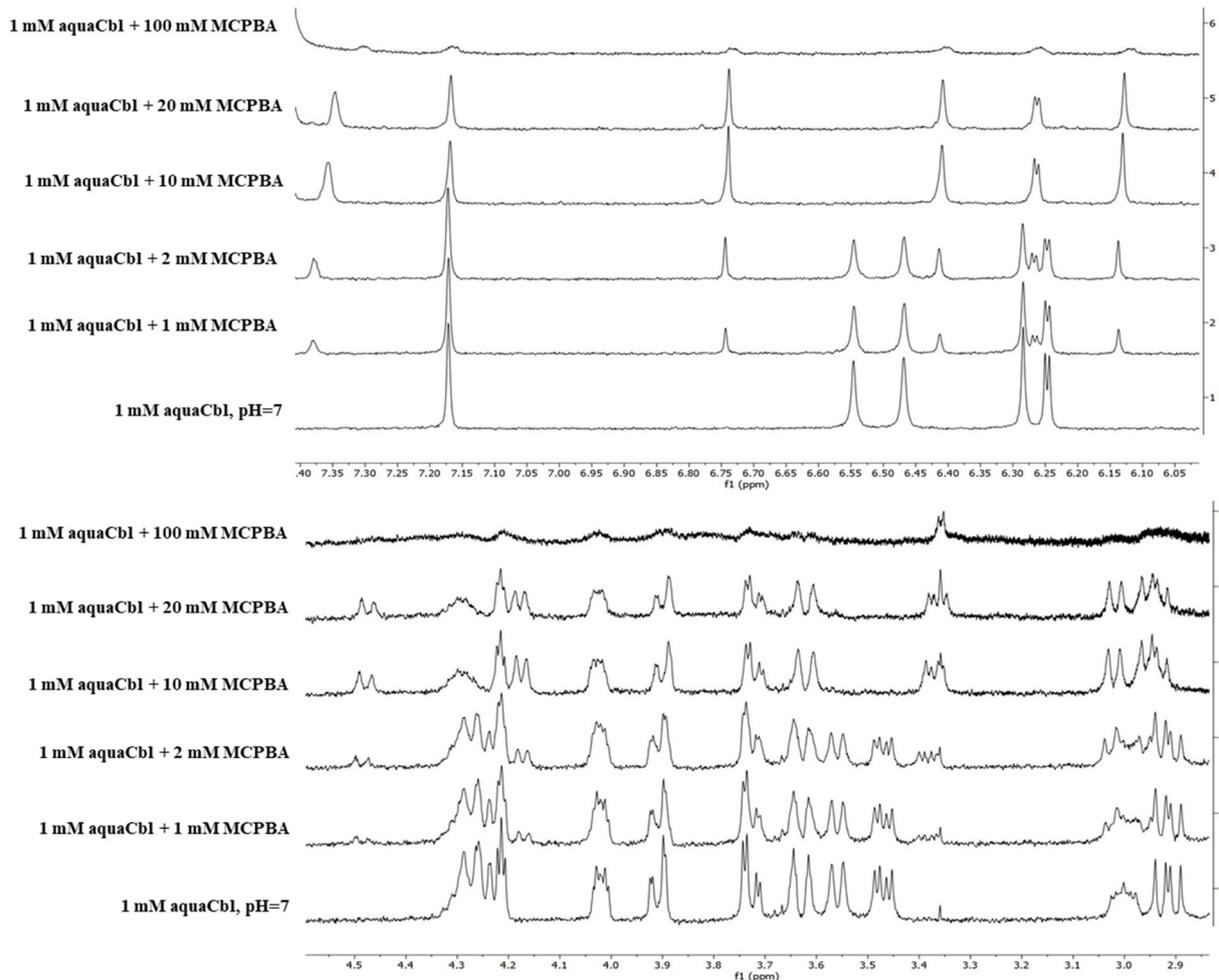


Fig. 4  $^1\text{H-NMR}$  titration of aquaCbl with MCPBA. Conditions: concentrations indicated in the figure, pH 7 in Britton–Robinson universal buffer (40 mM each borate, acetate, phosphate).

density functional theory (DFT) calculations suggest much smaller spectral differences between  $\text{H}_2\text{O}$  and  $\text{H}_2\text{O}_2$  Cbl complexes, compared to  $\text{H}_2\text{O}$  vs. a deprotonated peroxide.<sup>5</sup> By contrast, the Cbl-MCPBA NMR spectrum is distinctly different from aqua as well as from other possible candidates (hydroxo, base-off aqua). Moreover, the MCPBA signals themselves are now all shifted compared to pure MCPBA, consistent with coordination to Cbl. The minor additional signals seen in the 7.5–6 ppm region of the Cbl-MCPBA spectrum may be due to Cbl-bound MCPBA or to degradation products in line with previous proposals.<sup>14</sup> The concentration dependence of the Cbl-MCPBA  $^1\text{H-NMR}$  spectra (cf. Fig. 4) confirms the latter observation, with the Cbl signals mostly degraded at 100 mM MCPBA. The titration also confirms that (1) only one adduct with MCPBA is formed, and (2) saturation of Cbl occurs below 10 mM, in line with the UV-vis titration of Fig. 1.

DFT geometry optimization of Cbl models with a neutral MCPBA ligand bound to Co either *via* the (carboxyl)OH group or *via* the Cl atom yield almost identical energies, with the OH slightly favoured by only  $\sim 1 \text{ kcal mol}^{-1}$ , cf. Table S1 (ESI $^\dagger$ ).

Since a Co-Cl(MCPBA) complex is extremely unlikely, one may further assume that the same is true for the Co-(H)OO-C(O)-isomer. This is in line with the fact that indeed organic acids are not known to be effective ligands to Cbl. On the other hand, the geometric parameters predicted for the deprotonated MCPBA as a ligand, as well as for *t*BuOO $^-$  and *t*BuOOH, are very similar to those previously reported for  $\text{H}_2\text{O}_2$  and HOO-complexes of Cbl at the same level of theory (cf. Table S1 and Fig. S5, ESI $^\dagger$ ).<sup>5</sup> Time-dependent DFT (TD-DFT) simulations of the UV-vis spectra of Co(III)-OOC(O)-MCPBA (cf. Table S2 and Fig. S6, S7, ESI $^\dagger$ ) reveal maxima at wavelengths intermediary between those of aqua and of hydroxo Cbl, which (especially taking into account the limits of the computational method for such complexes) are in line with the experimental observations.<sup>5,44</sup> Elongation of the CoO–OH bond in a hydroperoxo peroxide-Cbl model and in a heme ferric-hydroperoxo model by 1 Å was found to entail energy costs of  $\sim 45 \text{ kcal mol}^{-1}$  in both cases, with a crossing onto the triplet/quartet state and with no major changes in partial atomic charges on the Co/Fe or on the peroxo oxygen atoms. The latter suggests, as expected,



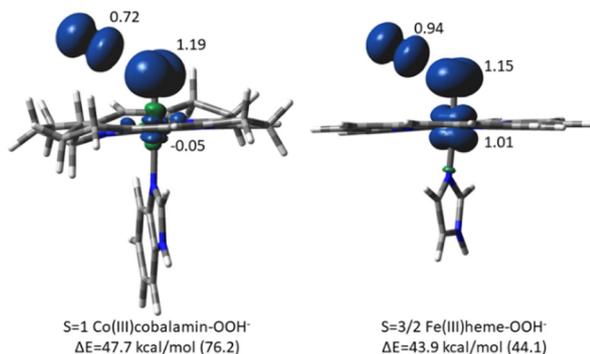


Fig. 5 Relative energies for  $S = 1$  Co(III)cobalamin-hydroperoxo and  $S = 3/2$  Fe(III)heme-hydroperoxo adducts after elongation of the O–O bonds by  $\sim 1$  Å relative to the equilibrium state. Also shown are Mulliken spin densities on the Co, Fe and oxygen atoms. Shown in parentheses are the relative energies of the low-spin states. The equilibrium geometries of the low-spin states ( $S = 0$  and  $S = 1/2$ , respectively) are taken as references. From UB3LYP/def2-SV(P)/D3 calculations.

that oxygen–oxygen bond cleavage in both cases would be homolytic (hence, no change in formal oxidation state from the Fe(III)/Co(III)-hydroperoxo stage) and would lead to a hydroxyl radical and a ferryl/chromyl unit. However, as shown in Fig. 5, while the ferryl does show the classical Fe(IV)-oxo center spin density distribution (50% on each bonding partner), the chromyl unit is clearly a Co(III)-oxyl (diamagnetic Co(III) with an  $S = 1/2$  oxyl ligand) similarly to other late transition metals such as Cu where “high-valent oxo” complexes in general tend to feature oxyl ligands, within the framework of the recently coined “oxo wall” paradigm.<sup>1,49–53</sup>

The resonance Raman spectra of Cbl are dominated by the corrin ring and relatively insensitive to the nature of the exogenous ligand. Fig. 6 shows that the resonance Raman spectra of hydroperoxo- and MCPBA-Cbl(III) are very similar to each other, and slightly different from that of  $\text{H}_2\text{OCbl}^+$ , suggesting that in both cases an anionic peroxy moiety is coordinated to the corrin. The peroxy/MCPBA vs. aqua differences are observable, especially in the signals at  $\sim 1495$ ,  $795$ ,  $725$  and  $630$   $\text{cm}^{-1}$ . Possibly of more interest is the band observed for the control MCPBA sample at  $873$   $\text{cm}^{-1}$  (cf. Fig. 6 and Fig. S8, ESI<sup>†</sup>), which is known<sup>54</sup> to be due to the O–O stretching. This band overlaps with a shoulder in the aquaCbl spectrum. Importantly, in the MCPBA + Cbl sample the shoulder at  $873$   $\text{cm}^{-1}$  is distinctly weaker than in the aquaCbl sample; an increase in intensity would have been expected in a sample where the signals of aquaCbl and MCPBA overlap physically without any chemical interaction. This may be taken as evidence against a scenario where aquaCbl and MCPBA are still intact. Instead, also importantly, a new weak shoulder at  $866$   $\text{cm}^{-1}$  appears in the MCPBA + Cbl spectrum, tentatively assignable as the O–O stretching of the Cbl-MCPBA complex. For comparison, in the  $\text{H}_2\text{O}_2$  + Cbl sample (also shown in Fig. 6 and Fig. S8, ESI<sup>†</sup>), a similar shift is seen compared to aquaCbl, but only to  $870$   $\text{cm}^{-1}$ .

In our initial attempts to employ mass spectrometry (HRMS ESI) to detect the Co(III)-hydroperoxide complex of cobalamin,

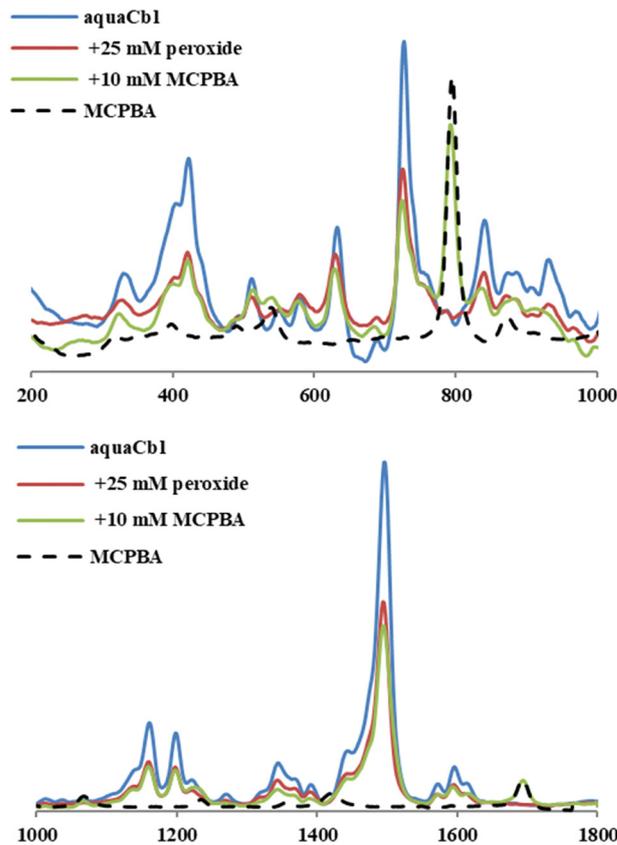


Fig. 6 Resonance Raman spectra of Cbl in the presence or absence of hydrogen peroxide or MCPBA. Conditions:  $1$  mM  $\text{H}_2\text{OCbl}^+$ ,  $25$  mM  $\text{H}_2\text{O}_2$ ,  $10$  mM MCPBA,  $50$  mM phosphate pH  $7$ ,  $22$  °C.

only the signal corresponding to pentacoordinated Cbl was identified, either in aqua<sup>5</sup> or in peroxy cobalamin. However, upon recording the ESI spectra immediately after mixing Cbl with MCPBA in water ( $\sim 2$  minutes from mixing to measurement), a signal could clearly be identified (cf. Fig. 7) corresponding to a Cbl-MCPBA adduct with the ( $m/z$ ) found at  $1500.5548$  [ $\text{M}$ ]<sup>+</sup> (vs. calculated  $1500.5565$ ), alongside the expected pentacoordinated  $1329.5716$  [ $\text{M} - \text{mCPBA}$ ]<sup>+</sup> as well as several  $z = 2$  signals for the Cbl-MCPBA. As shown in ESI<sup>†</sup> Fig. S10, in a sample incubated for 2 extra minutes before injecting into the instrument, the relative abundance of the Cbl-MCPBA peak decreases and, more importantly, several new signals appear which do not fit the mass of intact Cbl complexes and are thus taken as evidence for Cbl degradation in line with above-discussed considerations. At 30 minutes after mixing, neither the intact pentacoordinated Cbl nor the Cbl-MCPBA peaks were visible. This time evolution of the mass spectra was expected: based on the NMR titration, a  $10$  mM concentration of MCPBA was required in the HRMS experiment in order to have the cobalt sites saturated. However, as seen in the UV-vis spectra of Fig. 1 as well as in the NMR titration of Fig. 4, at  $10$ – $20$  mM there is already a decrease in intensity of the Cbl signals, suggestive of partial degradation; such degradation is expected to be accelerated



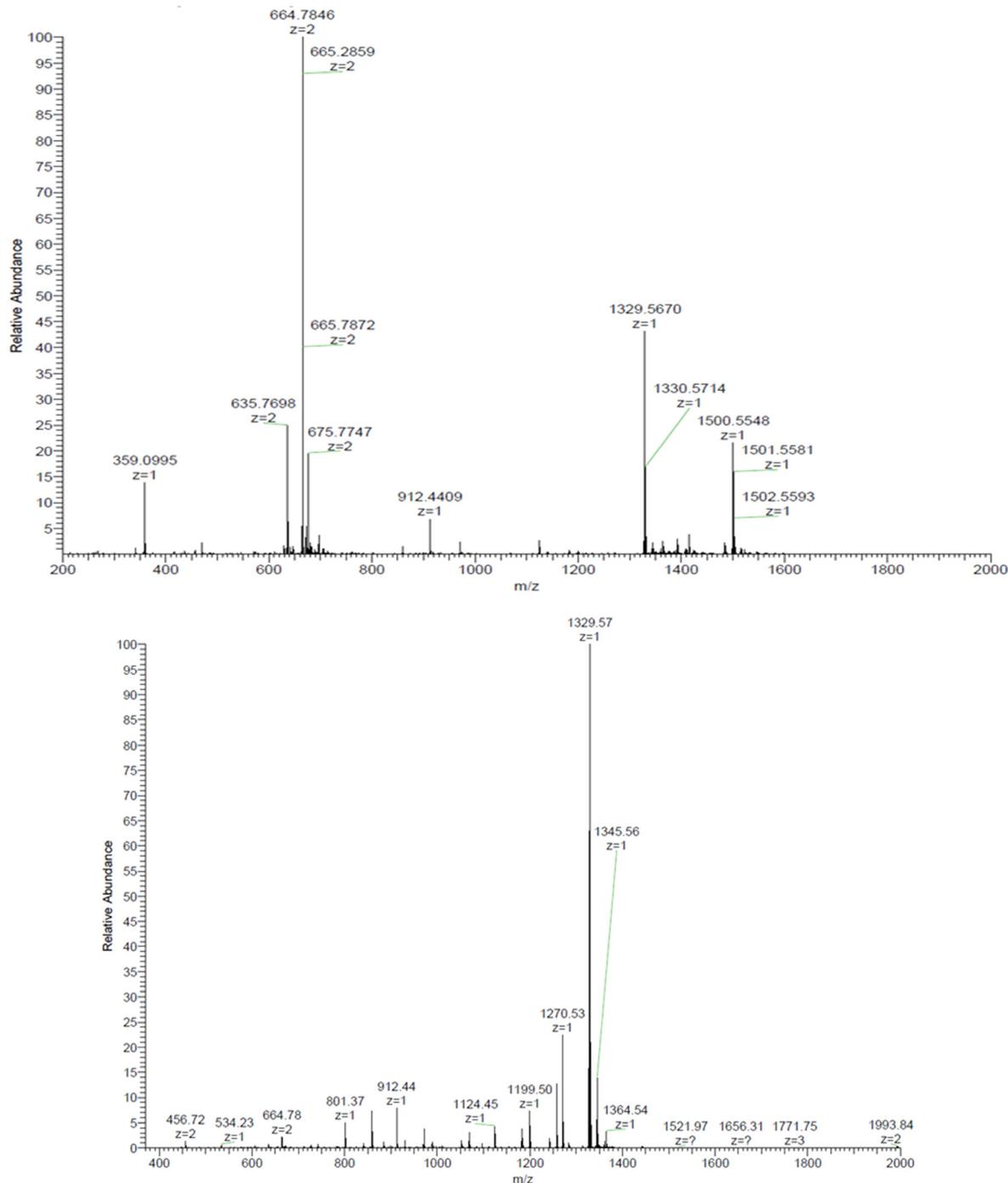


Fig. 7 Upper panel: ESI(+)-HRMS ( $m/z$ ) for the Cbl-MCPBA adduct. Calculated for  $C_{69}H_{93}CoN_{13}O_{17}PCl$  1500.5565, found: 1500.5548  $[M]^+$ , 1329.5716  $[M-mCPBA]^+$ , 675.7747  $[M-mCPBA + Na]^{2+}$ , 664.7855 5716  $[M-mCPBA + H]^{2+}$ . Lower panel: spectrum of the starting material (hydroxocobalamin). Conditions: 1 mM  $H_2OCl^+$ , 10 mM MCPBA, 50 mM phosphate pH 7.

by the HRMS protocol (see capillary and vaporizer temperatures in figure legends).

## Conclusions

To conclude, Cbl(III) forms a peroxyacid complex with deprotonated *m*-chloroperoxybenzoic acid, as supported by UV-vis,

$^1H$ -NMR and resonance Raman spectroscopic data, density functional calculations, and mass spectrometry. By contrast, *t*BuOOH does not appear to be able to form a peroxy complex with Cbl(III). These findings suggest that the peroxide coordination chemistry of cobalamin does extend beyond simply hydrogen peroxide,<sup>5</sup> but that not all peroxy compounds should be expected to bind to Cbl(III).



## Author contributions

ML: conceptualization, investigation, validation, formal analysis, visualization, writing – original draft, review and editing; CZT: conceptualization, methodology, investigation, validation, formal analysis, visualization; NH: investigation, methodology, formal analysis, visualization; AA: investigation; RS: methodology, investigation, validation; SDI: methodology, investigation; BAMV – methodology, investigation, validation; RSD – conceptualization, investigation, validation, visualization, writing – review and editing.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

Funding from the Romanian Ministry of Education and Research (projects PN-III-P4-ID-PCCF-2016-0142 and PN-III-P1-1.1-PD-2021-0279) and from the European Union Social Fund (Entrepreneurship for innovation through doctoral and postdoctoral research POCU/380/6/13/123886) is gratefully acknowledged. Prof S. V. Makarov (Ivanovo State University of Chemistry and Technology) and Artiom Gaina-Gardiuta (UBB) are thanked for helpful discussions.

## Notes and references

- R. Silaghi-Dumitrescu, *Struct. Bond*, 2013, **150**, 97–118.
- Y. Zhang and J. A. Gascon, *J. Inorg. Biochem.*, 2008, **102**, 1684–1690.
- S. P. de Visser, D. Kumar, R. Neumann and S. Shaik, *Angew. Chem., Int. Ed.*, 2004, **43**, 5661–5665.
- H. P. Hersleth, U. Ryde, P. Rydberg, C. H. Gorbitz and K. K. Andersson, *J. Inorg. Biochem.*, 2006, **100**, 460–476.
- M. Lehene, D. Plesa, S. Ionescu-Zinca, S. D. Iancu, N. Leopold, S. V. Makarov, A. M. V. Brănzanic and R. Silaghi-Dumitrescu, *Inorg. Chem.*, 2021, **60**, 12681–12684.
- F. Carrascoza, M. Surducan, L. A. Eriksson and R. Silaghi-Dumitrescu, *Inorg. Chim. Acta*, 2020, **509**, 119682.
- W. Sand, M. Dopson, G. Levicán, Á. Sandoval, A. Ferrer, J. Rivera, C. Zapata, J. Norambuena, Á. Sandoval, R. Chávez and O. Orellana, *Front. Microbiol.*, 2016, **7**, 748.
- D. S. Salnikov, S. V. Makarov and O. I. Koifman, *New J. Chem.*, 2021, **45**, 535–543.
- P. W. Johns, A. Das, E. M. Kuil, W. A. Jacobs, K. J. Schimpf and D. J. Schmitz, *Int. J. Food Sci. Technol.*, 2015, **50**, 421–430.
- G. Tsiminis, E. P. Schartner, J. L. Brooks and M. R. Hutchinson, *Appl. Spectrosc. Rev.*, 2017, **52**, 439–455.
- R. S. Dassanayake, M. M. Farhath, J. T. Shelley, S. Basu and N. E. Brasch, *J. Inorg. Biochem.*, 2016, **163**, 81–87.
- I. A. Dereven'kov, S. V. Makarov, N. I. Shpagilev, D. S. Salnikov and O. I. Koifman, *Biometals*, 2017, **30**, 57–764.
- I. A. Dereven'kov, D. S. Salnikov, S. V. Makarov, G. R. Boss and O. I. Koifman, *Dalton Trans.*, 2013, **42**, 15307–15316.
- J. Cheng, Y. Shiota, M. Yamasaki, K. Izukawa, Y. Tachi, K. Yoshizawa and H. Shimakoshi, *Inorg. Chem.*, 2022, **61**, 9710–9724.
- I. A. Dereven'kov, E. S. Sakharova, V. S. Osokin and S. V. Makarov, *Int. J. Mol. Sci.*, 2022, **23**, 11907.
- P. Kumar, S. V. Lindeman and A. T. Fiedler, *J. Am. Chem. Soc.*, 2019, **141**, 10984–10987.
- M. J. Frisch, G. W. Trucks, 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. 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. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, T. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. 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 09, Revis. E.01*, Gaussian, Inc., Wallingford CT, 2016.
- A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1988, **37**, 785–789.
- A. Schäfer, H. Horn and R. Ahlrichs, *J. Chem. Phys.*, 1992, **97**, 2571–2577.
- R. Ditchfield, *Mol. Phys.*, 1974, **27**, 789–807.
- V. Barone and M. Cossi, *J. Phys. Chem. A*, 1998, **102**, 1995–2001.
- J. P. Perdew and Y. Wang, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1992, **46**, 12947–12954.
- I. A. Dereven'kov, P. A. Ivlev, C. Bischin, D. S. Salnikov, R. Silaghi-Dumitrescu, S. V. Makarov and O. I. Koifman, *J. Biol. Inorg. Chem.*, 2017, **22**, 969–975.
- D. S. Salnikov, R. Silaghi-Dumitrescu, S. V. Makarov, R. van Eldik and G. R. Boss, *Dalton Trans.*, 2011, **40**, 9831–9834.
- M. Surducan, S. V. Makarov and R. Silaghi-Dumitrescu, *Polyhedron*, 2014, **78**, 72–84.
- I. A. Dereven'kov, D. S. Salnikov, S. V. Makarov, M. Surducan, R. Silaghi-Dumitrescu and G. R. Boss, *J. Inorg. Biochem.*, 2013, **125**, 32–39.
- D. S. Salnikov, I. A. Dereven'kov, S. V. Makarov, E. S. Ageeva, A. Lupan, M. Surducan and R. Silaghi-Dumitrescu, *Rev. Roum. Chim.*, 2012, **57**, 353–359.
- I. G. Pallares and T. C. Brunold, *Inorg. Chem.*, 2014, **53**, 7676–7691.
- P. M. Kozłowski, V. V. Nazarenko and A. A. Jarzecki, *Inorg. Chem.*, 2006, **45**, 1424–1426.
- A. P. Ghosh, P. Lodowski and P. M. Kozłowski, *Phys. Chem. Chem. Phys.*, 2022, **24**, 6093–6106.



- 32 J. Kuta, S. Patchkovskii, M. Z. Zgierski and P. M. Kozłowski, *J. Comput. Chem.*, 2006, **27**, 1429–1437.
- 33 K. Mieda-Higa, A. Al Mamun, T. Ogura, T. Kitagawa and P. M. Kozłowski, *J. Raman Spectrosc.*, 2020, **51**, 1331–1342.
- 34 H. Solheim, K. Kornobis, K. Ruud and P. M. Kozłowski, *J. Phys. Chem. B*, 2011, **115**, 737–748.
- 35 M. Kumar and P. M. Kozłowski, *Coord. Chem. Rev.*, 2017, **333**, 71–81.
- 36 A. P. Ghosh, P. Lodowski, A. Chmielowska, M. Jaworska and P. M. Kozłowski, *J. Catal.*, 2019, **376**, 32–43.
- 37 T. Andruniow, M. Z. Zgierski and P. M. Kozłowski, *J. Am. Chem. Soc.*, 2001, **123**, 2679–2680.
- 38 P. M. Kozłowski, B. D. Garabato, P. Lodowski and M. Jaworska, *Dalton Trans.*, 2016, **45**, 4457–4470.
- 39 M. D. Liptak and T. C. Brunold, *J. Am. Chem. Soc.*, 2006, **128**, 9144–9156.
- 40 T. A. Stich, N. R. Buan and T. C. Brunold, *J. Am. Chem. Soc.*, 2004, **126**, 9735–9749.
- 41 E. D. Greenhalgh, W. Kincannon, V. Bandarian and T. C. Brunold, *Biochemistry*, 2022, **61**, 195–205.
- 42 A. J. Reig, K. S. Conrad and T. C. Brunold, *Inorg. Chem.*, 2012, **51**, 2867–2879.
- 43 A. M. V. Brânzanic, U. Ryde and R. Silaghi-Dumitrescu, *J. Inorg. Biochem.*, 2020, **203**, 110928.
- 44 A. A. A. Attia, D. Cioloboc, A. Lupan and R. Silaghi-Dumitrescu, *J. Inorg. Biochem.*, 2016, **165**, 49–53.
- 45 A. A. A. Attia, A. Lupan and R. Silaghi-Dumitrescu, *RSC Adv.*, 2013, **3**, 26194–26204.
- 46 R. Silaghi-Dumitrescu and I. Silaghi-Dumitrescu, *J. Inorg. Biochem.*, 2006, **100**, 161–166.
- 47 H. M. Abu-Soud, D. Maitra, J. Byun, C. E. A. Souza, J. Banerjee, G. M. Saed, M. P. Diamond, P. R. Andreana and S. Pennathur, *Free Radicals Biol. Med.*, 2012, **52**, 616–625.
- 48 D. Maitra, I. Ali, R. M. Abdulridha, F. Shaeib, S. N. Khan, G. M. Saed, S. Pennathur and H. M. Abu-Soud, *PLoS One*, 2014, **9**, 5–12.
- 49 V. A. Larson, B. Battistella, K. Ray, N. Lehnert and W. Nam, *Nat. Rev. Chem.*, 2020, **4**, 404–419.
- 50 E. Andris, R. Navrátil, J. Jašík, M. Srnc, M. Rodríguez, M. Costas and J. Roithová, *Angew. Chem., Int. Ed.*, 2019, **58**, 9619–9624.
- 51 R. Silaghi-Dumitrescu, M. Surducan and A. Papp, *Croat. Chem. Acta*, 2014, **87**, 75–78.
- 52 A. Imre, A. Mot and R. Silaghi-Dumitrescu, *Cent. Eur. J. Chem.*, 2012, **10**, 1527–1533.
- 53 R. Silaghi-Dumitrescu, *Rev. Chim.*, 2007, **58**, 461–464.
- 54 Z. Tang, J. Xiao, F. Li, Z. Ma, L. Wang, F. Niu and X. Sun, *ACS Omega*, 2020, **5**, 10451–10458.

