Coordination frameworks assembled from Cu\textsuperscript{II} ions and H\textsubscript{2}-1,3-bdpb ligands: X-ray and magneto structural investigations, and catalytic activity in the aerobic oxidation of tetralin†

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The syntheses and crystal structures of H\textsubscript{2}-1,3-bdpb-MeOH, [Cu\textsuperscript{II}(1,3-bdpb)(OCH\textsubscript{3})\textsubscript{2}] (CFA-5) and [CuCl-(H\textsubscript{2}-1,3-bdpb)] (H\textsubscript{2}-1,3-bdpb = 1,3-bis(3,5-dimethyl-1H-pyrazol-4-yl)benzene) are described. The copper(II) containing metal–organic framework (termed Coordination Framework Augsburg University-5, CFA-5) crystallizes in the trigonal crystal system, within the space group \textit{R\textsubscript{3}} (no. 148) and the unit cell parameters are as follows: \(a = 26.839(3)\) Å, \(c = 15.8317(16)\) Å, \(V = 9876.2(19)\) Å\textsuperscript{3}. CFA-5 features a two-fold inter-penetrated 3-D microporous framework structure of cross-linked wheel-shaped \{Cu\textsuperscript{II}(pz)(OMe)\}\textsubscript{12} fundamental building units, each containing twelve copper(II) ions, \(\mu\text{-}2\)-bridging MeO\textsuperscript{-} groups and pyrazolate (pz\textsuperscript{-}) ligands. Replacing copper(II) acetate by copper(II) chloride in the synthesis leads to compound [Cu\textsuperscript{II}Cl(H\textsubscript{2}-1,3-bdpb)], which crystallizes in the orthorhombic crystal system, within the space group \textit{Pnma} (no. 62) and the unit cell parameters are as follows: \(a = 6.1784(8)\) Å, \(b = 6.1784(8)\) Å, \(c = 6.1784(8)\) Å, \(V = 1583.8(4)\) Å\textsuperscript{3}. In contrast to the former compound, CuCl(H\textsubscript{2}-1,3-bdpb) is a non-porous compound consisting of Cu\textsuperscript{II}–Cl zigzag chains expanding in the direction \{100\} and H\textsubscript{2}-1,3-bdpb ligands. CFA-5 is characterized by elemental and thermogravimetric analyses, variable temperature powder X-ray diffraction and IR-spectroscopy; and its porosity and magnetic properties are described in detail. CFA-5 shows a promising catalytic activity in the heterogeneously catalyzed aerobic oxidation of tetralin, which is compared with other catalytically active metal–organic frameworks.

Introduction

In the last decade the major driving force for the continued interest in metal–organic frameworks (MOFs) has been the promising technical applications of these materials ranging from gas storage and separation,\textsuperscript{1} drug delivery,\textsuperscript{2} sensing\textsuperscript{3} to promising technical applications of these materials ranging from gas storage and separation,\textsuperscript{1} drug delivery,\textsuperscript{2} sensing\textsuperscript{3} to promising technical applications of these materials ranging from gas storage and separation,\textsuperscript{1} drug delivery,\textsuperscript{2} sensing\textsuperscript{3}.

The large specific surface areas of up to \(7140\) m\textsuperscript{2} g\textsuperscript{-1},\textsuperscript{5} open pores with apertures of up to \(98\) Å,\textsuperscript{6} the high thermal stability and a high volume density of active sites within the framework structures suggest a host of applications in the field of heterogeneous catalysis in particular.\textsuperscript{14,7}

On searching for suitable MOF candidates for catalytic oxidation processes, copper-containing frameworks seem to be particularly interesting owing to the fact that this element is present in the active site of many metalloenzymes, such as copper-containing oxidases or oxygenases.\textsuperscript{8} Biologically inspired MOF catalysts trying to mimic structural and functional aspects of these enzymes could be applied for wide-ranging synthetic applications in the oxidation of organic intermediates containing non-activated C–H-bonds. However, until now, only a few reports have appeared in the literature where Cu-MOFs were successfully applied as oxidation catalysts. In particular, the Cu-catalyzed hydroxylation of phenol,\textsuperscript{9} oxidation of trimethylsilyl enolates to \(\alpha\)-hydroxy ketones,\textsuperscript{10} allylic oxidation of cyclohexenes,\textsuperscript{11} cross-dehydrogenative coupling reaction of ethers with 2-carbonyl-substituted phenols,\textsuperscript{12} and oxidation of benzene derivatives and benzylic compounds have been described.\textsuperscript{13} Some years ago we had started the development of Cu-containing MOF catalysts containing pyrazole ligands for the purpose of introducing oxygen functions into the C–H bonds.
of organic substrates. Our previous studies on the reactivity of CFA-2 (a Cu(i)-containing MOF) toward molecular oxygen for instance have shown that this compound is stable during oxidation and reduction of the Cu ions, suggesting its usage in liquid-phase oxidation reactions. The exploration of pyrazolate-based Cu-MOFs follows the simple rational and experimental observation that this ligand forms strong coordinative bonds with Cu(i) and Cu(II), thus leading to robust coordination polymers. Similar to carboxylate-type ligands, pyrazolates are known to form μ₂-bridges between metal ions. Unlike the former, the hydrolytic stability of N-heterocyclic ligand-based frameworks containing the latter, i.e. electron-rich transition metal ions, is exceptionally high which has been convincingly demonstrated. More speculatively, we might anticipate that pyrazolate heterocycles could mimic the coordination of Cu ions by histidine (= imidazolate) ligands, a coordination motif which is frequently observed in (multi-)copper oxidases, thus opening routes towards biomimetic catalysis.

μ₂-Bridging pyrazolate ligands have been frequently used to generate trinuclear copper (n/n) [Cu₃(μ₂-pz)₃] metallocycles, which later on were introduced as secondary building units (SBUs) into metal–organic frameworks. Polynuclear metal-locyclic coordination units of the type [{Cu₇(μ₃-OH)(μ₂-pz)}ₙ], n = 6, 8, 9, 12 and 14, have been described, which might become SBUs of the as yet undiscovered MOFs. One such example, CFA-5, a novel MOF which contains a dodecanuclear {Cu₆(μ₃-pz)(OMe)}₁₂ SBU is presented in this article. CFA-5 was characterized by elemental and thermogravimetric analyses, X-ray diffraction and IR-spectroscopy and its porosity and magnetic properties have been studied in detail. Additionally, the results of catalytic tests of CFA-5 in the aerobic oxidation of tetralin are presented and compared to other Cu-MOFs.

Results and discussion

Syntheses and characterization

H₂-1,3-bdpb (1,3-bis(3,5-dimethyl-1H-pyrazol-4-yl)benzene) was synthesized according to the published procedure. The compound CFA-5 was obtained as green hexagonal prism crystals (Fig. 1) after heating a MeOH solution of H₂-1,3-bdpb and copper(n) acetate hydrate in the presence of 2,6-lutidine. By applying microwave irradiation instead of conventional heating the reaction time was reduced drastically, from 3 d to 25 min.

Replacing copper(n) acetate by copper(n) chloride in the synthesis leads to the compound [CuCl(H₂-1,3-bdpb)] (Scheme 1).

Single crystal structure analyses

H₂-1,3-bdpb-MeOH (C₁₆H₂₂N₄·CH₃OH). H₂-1,3-bdpb-MeOH crystallizes in the monoclinic crystal system within the space group P2₁/n (no. 14). Detailed description of the crystal structure of H₂-1,3-bdpb-MeOH is presented in ESI.† [Cu₄(1,3-bdpb)(OCH₃)₂] (CFA-5). CFA-5 crystallizes in the trigonal crystal system within the space group R₃ (no. 148). The asymmetric unit consists of eighteen carbon, four nitrogen, two oxygen and two copper atoms. An Ortep style plot of the asymmetric unit of CFA-5 with atom labels is shown in ESI, Fig. S3.† CFA-5 features a 3-D two-fold interpenetrated microporous structure. The twelve metal atoms are connected by twelve μ₂-bridging MeO⁻ and by twelve pyrazolate moieties belonging to 1,3-bdpb⁻ ligands, respectively, creating dodecanuclear copper(n) rings, as shown in Fig. 2a. In the literature several examples of copper pyrazolate structures created from four up to tetradecanuclear copper rings are known. To the best of our knowledge, CFA-5 represents the first example of a copper-containing MOF in which a dodecanuclear copper(n) SBU was structurally characterized. In these SBUs, each Cu(n) center exhibits an unusual distorted tetrahedral coordination geometry (as shown in Fig. 2b). The coordination geometry of each Cu(n) (N,N',O,O') subset lies between that of the tetrahedrally coordinated Cu(i) ions and that of Cu(II) complexes featuring a square planar coordination environment. The Cu-N distances range between 1.924(5) and 1.963(5) Å, whereas the Cu-O distances range between 1.904(4) and 1.929(5) Å (see Table 1). These values are in good agreement with those found in structurally related copper compounds.

Placing a least-squares plane through the twelve Cu(n) ions of each SBU, six 1,3-bdpb⁻⁻ ligands (blue and purple ligands in Fig. 2a) are found above and the six remaining ones (black and green) are placed below this plane. Among the twelve coordinating pyrazolate moieties six are almost perpendicular to the plane passing through the Cu centers (blue and green ligands; the angle between the Cu₁₂ plane and a least-squares plane through the twelve Cu(II) ions is exceptionally high which has been convincingly demonstrated). More speculatively, we might anticipate that pyrazolate heterocycles could mimic the coordination of Cu ions by histidine (= imidazolate) ligands, a coordination motif which is frequently observed in (multi-)copper oxidases, thus opening routes towards biomimetic catalysis.

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always found opposite to the 1,3-bdpb\(^{2-}\) ligand that bridges the same pair of Cu\(^{II}\) ions. Of the twelve methoxides, six are pointing their methyl groups towards the center of the dodecanuclear SBU, thus leading to a hydrophobic internal channel surface.

Defining the wheel-shaped \(\{\text{Cu}^{II}(\text{pz})(\text{OMe})\}_\text{12}\) fragment as SBU of CFA-5, these SBUs are connected by the meta-substituted benzene ring of the 1,3-bis(3,5-dimethyl-1H-pyrazol-4-yl)benzene ligand, thus creating one-dimensional channels expanding in the \(c\)-direction of the crystal lattice (see Fig. 2c and d). Taking the van der Waals radii of hydrogen atoms (1.2 Å) into account, the narrowest channel diameter calculated between the hydrogen atoms of the MeO\(^-\) groups is 4.5 Å, whereas the pore width, calculated between carbon atoms of the methyl groups and pyrazolate rings, is 5.86 Å (see Fig. S4, ESI\(^*\)). An estimation with the program SQUEEZE reveals\(^{21}\) that the initial solvent’s accessible void volume is 1213.3 Å\(^3\), which is 12.3% of the unit cell volume (9876.2(19) Å\(^3\)).
Table 1  Selected bond lengths of CFA-5

<table>
<thead>
<tr>
<th>Atoms</th>
<th>Bond length (Å)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cu(1)−O(1)</td>
<td>1.923(4)</td>
</tr>
<tr>
<td>Cu(1)−N(4)</td>
<td>1.924(5)</td>
</tr>
<tr>
<td>Cu(1)−N(1)</td>
<td>1.925(5)</td>
</tr>
<tr>
<td>Cu(1)−O(2)</td>
<td>1.929(5)</td>
</tr>
<tr>
<td>Cu(2)−O(1)</td>
<td>1.904(4)</td>
</tr>
<tr>
<td>Cu(2)−O(2)/#1</td>
<td>1.921(4)</td>
</tr>
<tr>
<td>Cu(2)−N(3)</td>
<td>1.942(5)</td>
</tr>
<tr>
<td>Cu(2)−N(2)</td>
<td>1.963(5)</td>
</tr>
</tbody>
</table>

Symmetry transformations used to generate equivalent atoms: #1 x – y + 2/3, x + 1/3, –z + 1/3.

The compound CuCl(H$_2$-1,3-bdpb) crystallizes in the orthorhombic crystal system within the space group Pnma (no. 62). The asymmetric unit consists of one copper, one chlorine, two nitrogen and nine carbon atoms. An Ortep style plot of the asymmetric unit of CuCl(H$_2$-1,3-bdpb) with atom labels is shown in the ESI, Fig. S5.† The copper, chlorine and two carbon atoms of the phenyl ring are lying on the crystallographic mirror plane (4c in Wyckoff notation). The two Me-pz rings of each H$_2$-1,3-bdpb ligand are inclined with respect to the central benzene ring. The least-squares planes running through the atoms of the Me-pz ring and the atoms of the benzene ring, respectively, have an angle of 40.96(2)°, which is very close to the value found for H$_2$-1,3-bdpb-MeOH (41.24(5)°).

The structure is created by Cu(μ)−Cl zigzag chains expanding in the [100] direction and H$_2$-1,3-bdpb$^{2−}$ ligands. The coordination geometries of the Cu(μ) ions are tetrahedral, being composed of two chlorine and two nitrogen atoms from two H$_2$-1,3-bdpb ligands, as depicted in Fig. S6a.† The Cu−N distances range from 1.969(3) to 1.970(3) Å, while Cu−Cl distances range from 2.41(2) to 2.53(2) Å. These values are in good agreement with those found in structurally related copper compounds.26

The compound CuCl(H$_2$-1,3-bdpb) exhibits a layered structure in which the layers are shifted from each other by half of the b period (6.45 Å) (see Fig. S6b†). The van der Waals interactions between CH$_3$ groups and weak hydrogen bonds are responsible for the observed crystal packing motif. The closest non-bonding distance between hydrogen atoms of the methyl groups from closely-packed molecules is equal to 2.67(1) Å, while N−Cl distance is equal to 3.25(2) Å. The atomic coordinates and isotropic thermal parameters, selected bond lengths and angles are presented in Tables S5 and S6 (ESI†), respectively.

TGA and XRPD studies

The phase purity of CFA-5 was confirmed by XRPD measurement under ambient conditions. The experimental XRPD pattern is consistent with the simulated one as gleaned from the single crystal X-ray diffraction data, as shown in Fig. 3. Differences in peak intensities are due to occluded solvent molecules.

In addition, the thermal stability of CFA-5 was determined by thermogravimetric (TG) and VTXRPD measurements. Prior to TG measurement, the sample was heated at 100 °C under vacuum for 2 h in order to remove the occluded solvent molecules (MeOH). As shown in Fig. 4, the thermogravimetric profile of CFA-5 under nitrogen exhibits a weight loss of 6.8% between 200 and 300 °C corresponding to the loss of one MeOH molecule per formula unit. The next degradation step (~37.5%), in which a fraction of the organic ligand is lost, is observed between 300 and 600 °C. According to the XRPD data presented in Fig. 5, the sample is stable up to ca. 150 °C.

Fig. 3  Calculated and measured X-ray powder patterns for CFA-5.
At 200 °C characteristic peaks at 6.60–6.76° 2θ (corresponding to the (210) and (101) planes), 9.44 (201), 11.42 ((300),(321)), 11.80 (102), 13.54 (202), 14.82 (431), 15.05 (322), 17.47 (510), 17.58 (531) become weaker and the peaks >20° in 2θ disappear, which indicates framework damage. Subsequent heating of the sample leads to amorphization of the material.

Physisorption results

CFA-5 exhibits permanent porosity, which is confirmed by argon gas sorption. Prior to sorption measurement, the crystalline sample was heated at 100 °C under vacuum for 1 h. The argon adsorption isotherm follows type I behaviour, typical for microporous solids. The maximum uptake achieved at 77 K and p/p₀ = 0.99 is 252 cm³ g⁻¹ (Fig. 6). The micropore volume obtained from the sorption isotherm is 0.2 cm³ g⁻¹ on applying the NLDFT method.²⁷ The same value was obtained using the de Boer t-method micropore analysis.²⁸ The adsorption data were fitted to the BET equation to give a surface area of 460 m² g⁻¹, the Langmuir surface area is equal to 510 m² g⁻¹. To evaluate the pore size distribution, the argon sorption isotherms sampled at 77 K were analyzed using non-local density functional theory (NLDFT) implementing a carbon equilibrium transition kernel for argon adsorption at 77 K based on a slit-pore model.²⁹ The distribution calculated by fitting the sorption data reveals micropores with a diameter of ca. 6.6 Å (Fig. 7, which is slightly higher than the data from the crystal structure (5.86 Å)).

CFA-5 becomes amorphous after prolonged heating at T > 250 °C, as indicated by VTXRPD data. However, the Ar gas sorption measurement, performed for the sample heated at 250 °C under vacuum for 1 h, shows that the material remains porous. The maximum uptake achieved at 77 K and p/p₀ = 0.99 is 192 cm³ g⁻¹ (Fig. 6). The micropore volume obtained from the sorption isotherm is 0.1 cm³ g⁻¹ applying the NLDFT method.²⁷ The adsorption data were fitted to the BET equation to give a surface area of 241 m² g⁻¹, the Langmuir surface area is equal to 278 m² g⁻¹. The pore size distribution calculated by fitting the sorption data shows that the volume of small pores, corresponding to the channels as described in the crystallographic section (<10 Å), is strongly reduced and the maximum is shifted to a lower value (5.5 Å), whereas the volume of larger
interactions seems to be less probable. The further indicating antiferromagnetic exchange interactions. Because of the high symmetry of the compound and the known tendency of compounds consisting of multiple coupled Cu(II) centers to contain paramagnetic impurities, the aforementioned explanation of this behavior due to ferromagnetic interactions seems to be less probable. The dependence of the molar magnetization \( M_m \) on the magnetic field \( B \) up to 5 T at 2.0 K is depicted in the inset of Fig. 8. In this region, the magnetization is almost independent of the applied field and retains a value close to zero. Therefore, the ground state of the compound corresponds to an \( S = 0 \) state, i.e. singlet spin ground state.

**Catalytic studies**

Cu-MOFs mimicking copper-containing enzymes like pMMO should be able to catalyse direct oxidation of methane to methanol with molecular oxygen. However, since the binding energy of the C–H bond of methane is quite high (421 kJ mol\(^{-1}\)), a catalyst is indispensable to open reaction pathways with low activation energies comparable to the enzymatic oxidation with pMMO. Currently, solid catalysts for selective oxidation of methane with molecular oxygen in the gas phase need high reaction temperatures of more than 200 °C. These high temperatures favour total oxidation of methanol, which can be oxidised more easily than methane. Thus, for preliminary catalytic studies it is advisable to choose a substrate with an activated C–H bond like benzylic compounds, e.g. oxidation of tetralin gives tetralol and tetralon as products (see Scheme 2). As shown in Scheme 2, there are two major mechanisms. One is the radical autoxidation mechanism proceeding via tetralin hydroperoxide as an intermediate which is decomposed into the products. The latter step is by far not as simple as shown in Scheme 2, because two different propagation cycles consuming tetralin hydroperoxide in bimolecular elementary steps are postulated. The radical autoxidation can be enhanced by a catalyst through radical formation (initiation) and hydroperoxide decomposition (Haber–Weiss cycle). The other mechanism is analogous to the enzymatic mono oxygenation through a proton driven formation of a high-valent metal-oxo species from molecular oxygen which oxygenates C–H bonds forming the corresponding alcohols.

In order to benchmark catalytic results, not only CFA-5, but also \([\text{Cu}(2\text{-pymo})_2] \) and \([\text{Cu}_3(\text{BTC})_2] \) (HKUST-1) were tested. \([\text{Cu}(2\text{-pymo})_2] \) had already been used in the aerobic oxidation of tetralin by Xamena et al. They observed that \([\text{Cu}(2\text{-pymo})_2] \) significantly enhances the tetralin hydroperoxide formation according to the radical autoxidation mechanism. Ryan et al. investigated both \([\text{Cu}(2\text{-pymo})_2] \) and \([\text{Cu}_3(\text{BTC})_2] \) by DFT calculations and came to the conclusion that both Cu-MOFs the decomposition of the tetralin hydroperoxide can only occur at the outer surface. In the case of \([\text{Cu}_3(\text{BTC})_2] \) the pore window size is 5.04 Å and, thus, is smaller than the kinetic diameter of tetralin which is expected to be larger than 5.85 Å. Since the narrowest channel diameter of CFA-5 is 4.5 Å, tetralin and tetralin hydroperoxide are also not expected to access the inner surface of the MOF. In the case of \([\text{Cu}_3(\text{BTC})_2] \) the pore size is large enough (10.7 Å according to Schlichte et al.) and the coordination at the free coordination site of the Cu paddlewheel structural motif indeed occur, but the calculated activation energy was not lower than the one for the gas phase. Therefore, the inner surface of \([\text{Cu}_3(\text{BTC})_2] \) should not act as a catalyst. However, the DFT calculations...
showed that undercoordinated Cu clusters at the outer surface of Cu-MOFs should decompose tetralin hydroperoxide.\textsuperscript{35}

In Fig. 9, the measured time dependence of the conversion of tetralin over time is shown for the three catalysts CFA-5, [Cu(2-pymo)\textsubscript{2}], and [Cu\textsubscript{3}(BTC)\textsubscript{2}]. All the three catalysts exhibit an induction period which is typical for the radical autoxidation mechanism (see Scheme 2, mechanism II). CFA-5 shows an activity very similar to that of [Cu(2-pymo)\textsubscript{2}], but both catalysts display a significantly shorter induction period than [Cu\textsubscript{3}(BTC)\textsubscript{2}]. Since we verified experimentally that no uncatalysed initiation of the reaction occurred, Cu at the outer surface of the MOFs seems to be an active site in the initial formation of tetralin radicals. Xamena \textit{et al.} excluded leaching of copper.\textsuperscript{13}\textsuperscript{b} By an optimized ICP-MS measurement procedure we could detect only 3.5 ppm Cu in the solution after 24 hours which corresponds to leaching of approx. 1.5\% of Cu from the original Cu-MOF. Since Xamena \textit{et al.} have shown that copper acetate corresponding to an assumed copper leaching of 10\% gives a much slower increase of conversion over time,\textsuperscript{13}\textsuperscript{b} we also assume that copper leaching is negligible. However, this will be investigated more deeply in future work.

The observed induction period (see Fig. 9) is an indication for the radical autoxidation mechanism proceeding via tetralin hydroperoxide (THP) as an intermediate. As expected, the measured selectivities of THP are significant and reach more than 80\% in the case of [Cu\textsubscript{3}(BTC)\textsubscript{2}] (see Fig. 10). Surprisingly CFA-5 and [Cu(2-pymo)\textsubscript{2}] show a different course of THP selectivity, because for these Cu-MOFs THP selectivity is not increasing, but decreasing with progressing time. Thus, the most likely explanation might be that CFA-5 and [Cu(2-pymo)\textsubscript{2}] catalyse the decomposition of THP, whereas [Cu\textsubscript{3}(BTC)\textsubscript{2}] is not able to do this. For further verification of this conclusion, a hot filtration test was performed after 6 h reaction time. As can be seen from Fig. 11 the reaction proceeds even after removal of the catalyst due to the radical autoxidation mechanism with an increasing selectivity to THP. This is a clear evidence that CFA-5 and [Cu(2-pymo)\textsubscript{2}] catalyse the decomposition of THP whereas [Cu\textsubscript{3}(BTC)\textsubscript{2}] does not. Since the course of conversion over time after 6 h is almost the same, irrespective of whether or not the catalyst is present (not shown), this allows the conclusion that all three catalysts do not catalyse the THP formation, if the initiation has been completed (which is evidently the case after 6 h).

To sum up, based on our catalytic studies and on the published knowledge\textsuperscript{13}\textsuperscript{b,35} a reaction mechanism can be proposed as shown in Fig. 12:

- Initiation of radical THP formation by all the investigated Cu-MOFs.
- Heterogeneously catalyzed THP decomposition by CFA-5 and [Cu(2-pymo)\textsubscript{2}].

Finally, due to the proposed mechanism as shown in Fig. 12, it can be assumed that CFA-5 will be also an active catalyst for the oxidants H\textsubscript{2}O\textsubscript{2} or TBHO since it catalyzes the decomposition of hydroperoxide species. This is already known from other Cu-MOFs.\textsuperscript{13}\textsuperscript{a,c}

**Conclusions**

The work reported here focuses on the synthesis and characterization of a novel metal–organic framework constructed from wheel-shaped dodecanuclear \{Cu\textsuperscript{II}(pz)(OMe)\textsubscript{12} coordi-
nation units, created by twelve copper(II) ions connected by twelve μ₂-bridging MeO⁻ groups and pyrazolate (pz⁻) ligands. The compound remains porous after solvent removal, as confirmed by TG, VT-XRPD and gas sorption. The magnetic measurements reveal dominant strong antiferromagnetic exchange interactions within the compound. Catalytic tests of CFA-5 in the aerobic oxidation of tetralin show a similar performance as with [Cu(2-pymo)₂]. Both Cu-MOFs catalyse the CFA-5 in the aerobic oxidation of tetralin show a similar performance as with [Cu(2-pymo)₂]. Both Cu-MOFs catalyse the decomposition of tetralin hydroperoxide after 6 h reaction time and initiate its formation within the first 6 h reaction time. For both Cu-MOFs it can be suggested that low coordinated Cu centers at the outer surface are responsible for the observed activity since in both cases the substrate cannot access the pore system. Both Cu MOFs contain diazaheterocyclic linkers which might be relevant for their catalytic activity. Under the applied test conditions [Cu₃(BTC)₂] with its carboxylate linkers, does not seem to form catalytically active Cu centers, neither at the inner nor at the outer surface.

**Experimental**

**Materials and general methods**

Commercially available reagents of analytical grade were used as received without further purification. [Cu(2-pymo)₂] was synthesized according to the published procedure.³⁷ [Cu₃(BTC)₂] was purchased from Sigma-Aldrich (Basolite C-300).

**Synthesis of H₂-1,3-bdpb.** H₂-1,3-bdpb was synthesized according to the published procedure.²⁰ H₂-1,3-bdpb is soluble in DMF, DMAc, DMSO, EtOH, and MeOH. ¹H-NMR (400 MHz). C₁₆H₁₈N₄ calcd: C: 72.15, H: 6.81, N: 21.04; found C: 71.93, H: 6.80, N: 20.49. IR: (cm⁻¹) 3162 w, 3121 w, 3068 w, 3006 w, 2960 w, 2919 w, 1609 m, 1581 m, 1525 w, 1456 m, 1436 s, 1415 w, 1401 w, 1380 w, 1365 w, 1314 w, 1288 m, 1275 w, 1239 w, 1148 w, 1071 w, 1035 m, 1010 s, 981 w, 891 w, 836 m, 800 s, 784 w, 760 s, 710 s, 700 m, 640 m, 586 w, 506 m, 447 w, 406 w. The IR spectrum of H₂-1,3-bdpb is shown in Fig. S7 (ESI†). For the single-crystal X-ray diffraction experiment H₂-1,3-bdpb was recrystallized from MeOH.

**Synthesis of [Cu₂(1,3-bdpb)(OCH₃)₂]** (CFA-5)

Solvothermal method. A mixture of Cu(OAc)₂·H₂O (11 mg, 0.055 mmol) and H₂-1,3-bdpb (15 mg, 0.056 mmol) was dissolved in MeOH (4 mL). 2,6-Dimethylpyridine (2,6-lutidine) (0.05 mL) was added and the solution was placed in a glass tube (10 mL). The tube was closed with a cap and heated at 120 °C for 3 d and subsequently cooled to room temperature. The green crystals were filtered off by suction and washed thoroughly with MeOH. The synthesis can be similarly performed at larger quantities (upscale factor: 50). Yield: 12 mg, 96% (based on Cu(OAc)₂·H₂O). [Cu₂(C₁₆H₁₈N₄)(OCH₃)₂] calcd: C: 47.67, H: 4.89, N: 12.35; found C: 47.44, H: 4.83, N: 12.28. IR: (cm⁻¹) 1605 w, 1584 w, 1539 w, 1492 w, 1424 m, 1377 w, 1331 m, 1301 w, 1244 m, 1167 w, 1130 w, 1063 w, 1034 w, 900 w, 816 m, 776 s, 709 s, 700 m, 662 w, 547 s, 447 s. The IR and UV/Vis spectra of CFA-5 are shown in Figs S7 and S8, † respectively.

**Microwave irradiation method.** A mixture of Cu(OAc)₂·H₂O (11 mg, 0.055 mmol) and H₂-1,3-bdpb (15 mg, 0.056 mmol) was dissolved in MeOH (3 mL). 2,6-Dimethylpyridine (2,6-lutidine) (0.05 mL) was added and the solution was placed in a Pyrex sample tube (10 mL). The tube was closed with a cap and placed in a microwave synthesizer (CEM, Discover S). The resulting mixture was heated to 150 °C at 300 W for 25 min and cooled to room temperature. The green microcrystalline material was filtered off and washed with MeOH. Yield: 12 mg, 96% (based on Cu(OAc)₂·H₂O). This material exhibited the same analytical results as the ones obtained by the solvothermal method.

**Synthesis of CuCl(H₂-1,3-bdpb).** A mixture of CuCl₂ (9 mg, 0.067 mmol) and H₂-1,3-bdpb (15 mg, 0.056 mmol) was dissolved in MeOH (4 mL). 2,6-Dimethylpyridine (2,6-lutidine) (0.05 mL) was added and the solution was placed in a glass tube (10 mL). The tube was closed with a cap and heated at 120 °C for 3 d and cooled to room temperature. The colorless crystals were filtered off by suction and washed thoroughly with MeOH. Yield: 9 mg, 44% (based on H₂-1,3-bdpb). CuCl (C₁₆H₁₈N₄Cl) calcd: C: 52.60, H: 4.97, N: 15.34; found C: 52.40, H: 5.04, N: 14.94. IR: (cm⁻¹) 3228 m (→NH), 1604 w, 1591 m, 1552 w, 1507 m, 1477 w, 1452 m, 1431 m, 1410 w, 1396 w, 1378 w, 1362 w, 1300 w, 1239 s, 1161 m, 1099 w, 1068 w, 1042 w, 1025 m, 1012 m, 980 w, 889 w, 808 s, 789 m, 779 m, 761 s, 713 s, 675 s, 658 s, 610 s, 585 s, 517 s, 454 w. The IR spectrum of 3 is shown in Fig. S7. †

**Microwave irradiation method.** A mixture of CuCl₂ (9 mg, 0.067 mmol) and H₂-1,3-bdpb (15 mg, 0.056 mmol) was dissolved in MeOH (2 mL). 2,6-Dimethylpyridine (2,6-lutidine) (0.05 mL) was added and the solution was placed in a Pyrex sample tube (10 mL). The tube was closed with a cap and...
placed in a microwave synthesizer (CEM, Discover S). The resulting mixture was heated to 140 °C at 100 W for 30 min and cooled to room temperature. The colorless microcrystalline material was filtered off and washed with MeOH. Yield: 11 mg, 54% (based on H2-1,3-bdpb). This material exhibited the same analytical results as the ones obtained by the solvothermal method.

**Single-crystal X-ray diffraction**

X-ray data for the single crystal structures determinations of H2-1,3-bdpb-MeOH, CFA-5 and CuCl(H2-1,3-bdpb) were collected on a Bruker D8 Venture diffractometer. Intensity measurements were performed using monochromated (doubly curved silicon crystal) MoKα radiation (0.71073 Å) from a sealed microfocus tube. Generator settings were 50 kV, 1 mA. Data collection temperature was −173 °C. APEX2 software was used for preliminary determination of the unit cell.38 Determination of integrated intensities and unit cell refinement were performed using SAINT.39 The structures were solved and refined using the Bruker SHELXTL Software Package.40 Selected crystal data and details of structure refinements for H2-1,3-bdpb-MeOH, CFA-5 and CuCl(H2-1,3-bdpb) are provided in Table 2. Complete crystallographic data for the structures H2-1,3-bdpb-MeOH, CFA-5 and CuCl(H2-1,3-bdpb) reported in this paper have been deposited in the CIF format with the Cambridge Crystallographic Data Center, 12 Union Road, Cambridge CB21EZ, UK as supplementary publication no. CCDC 1006742-1006744.

**Table 2** Crystal data and structure refinement for compounds H2-1,3-bdpb MeOH, CFA-5 and CuCl(H2-1,3-bdpb)

<table>
<thead>
<tr>
<th>Compound</th>
<th>H2-1,3-bdpb MeOH</th>
<th>CFA-5</th>
<th>CuCl(H2-1,3-bdpb)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Empirical formula</td>
<td>C16H18N4</td>
<td></td>
<td>C16H18N4ClCu</td>
</tr>
<tr>
<td>Formula</td>
<td>[Cu2(C16H16N4)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>(OCH3)2]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>M/g mol⁻¹</td>
<td>298.39</td>
<td>365.33</td>
<td>365.33</td>
</tr>
<tr>
<td>T/K</td>
<td>100(2)</td>
<td>100(2)</td>
<td>100(2)</td>
</tr>
<tr>
<td>Wavelength/Å</td>
<td>0.71073</td>
<td>0.71073</td>
<td>0.71073</td>
</tr>
<tr>
<td>Crystal system</td>
<td>Monoclinic</td>
<td></td>
<td>Orthorhombic</td>
</tr>
<tr>
<td>Space group</td>
<td>P2₁/m (no. 11)</td>
<td></td>
<td>Pmm (no. 62)</td>
</tr>
<tr>
<td>a/Å</td>
<td>4.7416(3)</td>
<td>26.839(3)</td>
<td>6.1784(8)</td>
</tr>
<tr>
<td>b/Å</td>
<td>12.1325(7)</td>
<td>26.839(3)</td>
<td>12.9041(17)</td>
</tr>
<tr>
<td>c/Å</td>
<td>14.0000(8)</td>
<td>15.8317(16)</td>
<td>19.863(6)</td>
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<tr>
<td>α/°</td>
<td>90</td>
<td>90</td>
<td>90</td>
</tr>
<tr>
<td>β/°</td>
<td>90</td>
<td>90</td>
<td>90</td>
</tr>
<tr>
<td>γ/°</td>
<td>90</td>
<td>90</td>
<td>90</td>
</tr>
<tr>
<td>V/Å³</td>
<td>802.19(8)</td>
<td>9876.2(19)</td>
<td>1583.8(4)</td>
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<td>Z</td>
<td>2</td>
<td>18</td>
<td>4</td>
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<tr>
<td>D, g cm⁻³</td>
<td>1.235</td>
<td>1.372</td>
<td>1.532</td>
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<tr>
<td>μ/mm⁻¹</td>
<td>0.080</td>
<td>1.955</td>
<td>1.549</td>
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<tr>
<td>F(000)</td>
<td>320</td>
<td>4176</td>
<td>752</td>
</tr>
<tr>
<td>θ Range/°</td>
<td>2.92 to 26.36</td>
<td>2.65 to 25.00</td>
<td>3.16 to 25.00</td>
</tr>
<tr>
<td>Refls. collected</td>
<td>16 984</td>
<td>19 551</td>
<td>12 815</td>
</tr>
<tr>
<td>Refls. unique</td>
<td>1705</td>
<td>3874</td>
<td>1460</td>
</tr>
<tr>
<td>R(int)</td>
<td>0.0634</td>
<td>0.1439</td>
<td>0.0391</td>
</tr>
<tr>
<td>GoofF</td>
<td>1.13</td>
<td>1.036</td>
<td>1.124</td>
</tr>
<tr>
<td>R1 [I &gt; 2σ(I)]/R1</td>
<td>0.0521</td>
<td>0.0679</td>
<td>0.0440</td>
</tr>
<tr>
<td>wR2 (all data)</td>
<td>0.1049</td>
<td>0.1515</td>
<td>0.1243</td>
</tr>
<tr>
<td>Largest diff. peak and hole/Å⁻³</td>
<td>0.244 and</td>
<td>0.763 and −0.925</td>
<td>0.448 and −0.441</td>
</tr>
</tbody>
</table>

Physical methods

Fourier transform infrared (FTIR) spectra were recorded with ATR in the range 4000–400 cm⁻¹ on a Bruker Equinox 55 FT-IR spectrometer. The following indications are used to characterize absorption bands: very strong (vs), strong (s), medium (m), weak (w). Elemental analyses (C, H, N) were carried out on a Perkin-Elmer 2400 Elemental Analyzer. Thermogravimetric analysis (TGA) was performed with a TGA Q500 analyser in the temperature range of 25–800 °C under flowing nitrogen at a heating rate of 10 K min⁻¹. The argon sorption isotherms at −196 °C up to p/p₀ = 1 were measured using a Quantachrome Autosorb-I ASI-CP-8 instrument. High purity gas was used for the adsorption experiments (argon 99.999%). Ambient temperature X-ray powder diffraction (XRPD) patterns were measured with a Seiffert XRD 3003 TT diffractometer equipped with a Meteor 1D detector operated at 40 kV, 40 mA, CuKα (λ = 1.54247 Å) with a scan speed of 10 s per step and a step size of 0.02° in 2θ. The variable temperature XRPD experiment data were collected in the 2θ range of 5–60° with 0.02° steps, with a Bruker D8 Advance diffractometer equipped with a Lynxeye linear position-sensitive detector, an MRI TCPU1 oven, in transmission geometry. The sample was loaded into a capillary (Hilgenberg) made from special glass no. 10, 0.5 mm diameter and 0.01 mm wall thickness. The patterns were recorded in a temperature range from 30 to 250 °C, in the 5–60° 2θ range, with one step per 1 s, and an angular step width of 0.02° in 2θ. Temperature program between measurements: heating rate (0.5 °C s⁻¹), then 10 min isothermal. Magnetic susceptibility data of CFA-5 were recorded using a Quantum Design MPMS-5XL SQUID magnetometer. The polycrystalline samples were compacted and immobilized into cylindrical PTFE capsules. The susceptibility data were acquired as a function of the field (0.1–5.0 T) and temperature (2.0–290 K). All data were corrected for the contribution of the sample holder and the diamagnetic contributions of CFA-5 calculated from tabulated values [χ_{m}(CFA-5) = −2.8 × 10⁻⁹ m² kg⁻¹ per asymmetric unit].

**Catalytic investigations**

The method used to determine the catalytic activity is based on the work of Xamena et al.11b Catalytic tests were carried out in a three-necked flask equipped with a reflux condenser and a gas inlet tube. Before each experiment, the glassware was passivated for a period of 12 h with 1 M aqueous sodium hydroxide solution, subsequently neutralized by rinsing with 1 M hydrochloric acid for 30 min and finally washed with demineralized water and dried at 80 °C in an oven. With this passivation procedure, uncatalysed initiation of the reaction could be avoided. The reaction temperature was set to 90 °C and the applied molar ratio of tetralin/metal was always 2000 : 1. 20 g of tetralin were used both as a solvent and a reactant. The required amount of catalyst was calculated from the previously mentioned ratio. Synthetic air at a flow rate of 0.5 mL s⁻¹ was adjusted by a needle valve. A HP 6890N GC with AutoSampler.
was used to analyze the reaction mixture. All samples were measured five times and the results were averaged. Peroxide concentration was determined by the TPP method. In time-resolved experiments, small samples of the reaction solution were taken after 6, 24 and 30 hours.

Conversion ($X$) of tetralin and selectivity ($S$) to tetralin hydroperoxide were calculated according to the following formulas:

\[
X_{\text{Tetralin}} = \frac{n_{\text{Tetralin,0}} - n_{\text{Tetralin}}}{n_{\text{Tetralin,0}}}
\]

\[
S_{\text{THP}} = \frac{n_{\text{THP}}}{n_{\text{Tetralin,0}} - n_{\text{Tetralin}}}
\]

The sum of selectivities to THP, tetralol and tetralone was always larger than 95%. Thus, the consecutive oxidation of tetralol and tetralone was negligible and the carbon balance fulfilled.

**Acknowledgements**

This work was supported by a grant from the Ministry of Science, Research and Art Baden-Württemberg and by the German Research Foundation (DFG) within Priority Program “Porous Metal–Organic Frameworks” (SPP 1362, MOFs).

**Notes and references**


38 APEX2 Version 2011.6, Bruker AXS Inc.