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A Di copper Peptidyl Complex Mimics the Activity of Catalase, a Key Antioxidant Metalloenzyme

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ABSTRACT: Catalases (CAT) are antioxidant metalloenzymes necessary for life in oxygen-metabolizing cells to regulate H₂O₂ concentration by accelerating its dismutation. Many physio-pathological situations are associated with oxidative stress resulting from H₂O₂ overproduction during which antioxidant defenses are overwhelmed. We have used a combinatorial approach associated with an activity-based screening to discover a first peptidyl di copper complex mimicking CAT. The complex was studied in detail and characterized for its CAT activity both in solution and in cells using different analytical methods. The complex exhibited CAT activity in solution and, more interestingly, on HyPer HeLa cells which possess a genetically encoded ratiometric fluorescent sensors of H₂O₂. These results highlight the efficiency of a combinatorial approach for the discovery of peptidyl complexes that exhibit catalytic activity.

INTRODUCTION

Aerobic organisms generate their energy through the reduction of O₂ to H₂O. During this process, reactive oxygen species (ROS) are continuously formed. The redox cascade initially forming O₂^{•−} leads to the subsequent formation of H₂O₂. H₂O₂ then dissociates to the hydroxyl radical (•OH) in mitochondria and/or peroxisomes. ROS play a key role in a range of different mechanisms such as cell proliferation and cell differentiation.^{1–3} They are very reactive molecules with strongly oxidizing properties. The most reactive and damaging is •OH. Although H₂O₂ and O₂^{•−} are not as reactive, they can become cytotoxic if their intracellular concentration exceeds a limiting threshold, overwhelming the protective pathways and leading to oxidative stress. Oxidative stress is known to be implicated in many physio-pathological situations.^{4–9} Although the presence of ROS is not always the sole cause of these different diseases, it nevertheless is a major contributor. It is, therefore, essential to control ROS concentrations in order to limit their deleterious effects. Antioxidant defenses protect biological systems from free radical toxicity and prevent damage due to excessive concentrations of ROS. This can be achieved either by controlling the production of ROS and/or by controlling the elimination of ROS. Antioxidant defenses include stoichio-

metric scavengers such as vitamins C and E, glutathione and metalloenzymes like superoxide dismutase (SOD),^{10–15} catalase (CAT),^{16,17} glutathione peroxidase (GPx), and glutathione reductase (GR).^{18,19}

Despite the fact that higher concentrations of H₂O₂ than O₂^{•−} or OH• are tolerated in cells, the intracellular concentrations of H₂O₂ must be tightly controlled at 1–100 nM.³ CAT is one of the enzymes involved in the regulation of H₂O₂ concentration by accelerating its dismutation.²⁰ During this reaction, one molecule of H₂O₂ is reduced into two molecules of H₂O, and a second molecule of H₂O₂ is oxidized into O₂.²¹ CAT is present in all aerobic species where it has been sought for.¹⁷ To date, two classes of catalase have been identified: a monofunctional heme CAT (HemeCAT), and a non-heme dinuclear manganese CAT (MnCAT).^{20–23} The dismutation of H₂O₂ consists of two bi-electronic processes, requiring a bi-electronic catalyst. In the case of the HemeCAT, the iron and porphyrin ligand both participate to the two electron exchanges, cycling between an (P)Fe^{III} resting state and a (P⁺)Fe^{IV}=O high oxidation state, (also called compound I).^{24,25} MnCAT was more recently discovered in lactic acid bacteria^{23,26,27} which lack heme and cytochrome. It has been demonstrated that MnCAT possesses a binuclear manganese active site which cycles between the reduced state Mn^{II}-Mn^{II} and the oxidized state Mn^{III}-Mn^{III},

affording the two electrons exchange required for the dismutation reaction.^{20,22,23}

Small molecular complexes mimicking CAT have been developed to reduce oxidative stress in pathological conditions.^{28–30} These bioinspired mimics are thus mainly mononuclear (Mn or Fe) porphyrin derivatives or dinuclear manganese complexes analogous to the active site of HemeCAT or MnCAT, respectively.²⁹ Dinuclear Mn complexes bioinspired from MnCAT are alkoxo, phenoxo, or oxo/carboxylato bridged complexes. In an excellent review, Signorella and Hureau have reported that besides the redox potentials of the metal ions key to catalytic H₂O₂ dismutation, the efficacy of the complexes is associated with the presence of an intramolecular base to assist in deprotonation. In addition, the presence of a vacant site on the Mn ion is required to coordinate H₂O₂, but the complex stability is reduced by the protonation of the bridging ligand.³¹ Few mononuclear copper complexes^{30,32–36} have been reported as CAT mimics as well but examples of dinuclear copper complexes are very rare.^{37–40} Generally, CAT mimics have been studied in organic solvents or have exhibited activity only at pH above 8. So far, contrarily to SOD mimics, no peptidyl complexes mimicking CAT have been reported, whatever the metal ion.^{41–44} This may be explained by the difficulty to rationally design short peptides able to bind two metal cations (to afford the two required electrons) with appropriate redox potentials and affinities. However, peptidyl ligands are of great interest since their synthesis is very versatile, they are biocompatible and water soluble. But it seems highly complicated to design Mn-based peptidyl complexes since peptide affinity for Mn(II) is usually in the 10³–10⁶ range at physiological pH even for quite long sequences.^{45–47} In contrast, short peptide sequences have been described to bind Cu(II) with high affinities (ca. 10¹²–10¹⁵).^{42,48,49} In this manuscript, we describe implementation of a combinatorial approach to synthesize a library of copper complexes, associated with an activity-based screening. This led to the discovery of the first peptidyl di copper complex mimicking CAT redox chemistry. The selected dinuclear copper complex was studied in detail and characterized for its CAT activity in solution and in cells. Very interestingly, despite moderate intrinsic catalysis constants, this complex was efficacious in a cellular assay.

RESULTS AND DISCUSSION

Screening process

To generate a large number of small peptidyl metal complexes, a peptide library was synthesized on a solid support by a split-and-pool combinatorial method leading to a one-bead-one-peptide library (Scheme S1). Inspired by the method developed by B. Imperiali who discovered a lanthanide binding tag⁵⁰ which we have recently implemented for the discovery of peptidyl SOD mimics efficient in cells,⁴³ the peptide library was designed to generate peptides that strongly bind copper. Peptides of the library have a general sequence Ac-PX₂X₃KHX₆LH-OH where X indicates places where diversity was introduced. Amino acids possessing coordinating side chains (aspartate or glutamate, tyrosine and histidine) and non-coordinating side chains (glycine and arginine) at positions X_n were chosen to explore the properties of the 5³ peptides (Figure 1A). After side chain deprotection using trifluoroacetic acid, the supported peptides were incubated in a solution of CuSO₄ (10 mM) in HEPES buffer (50 mM, pH 7.5) for 2 hr. The beads were then washed with buffer and EDTA (25 mM, pH 8), a strong competitor⁵¹ for Cu(II) chelation, to increase differential selectivity. This additional washing step allows for the removal of more loosely bound copper ions and/or the release of copper from less inert complexes. However, note that this step does not ensure that the remaining complexes are more stable or inert than the EDTA-Cu complex since it is a heterogenous process. To perform the screening of the peptidyl complexes library, the beads carrying the peptidyl complexes were immobilized in a 2% agarose gel according to the method developed by Nitz et al. (Figure 1B).⁵⁰ The gel was then irradiated using UV light (254 nm) for 15 minutes to release the portion of peptidyl complexes that were linked to the resin via a photocleavable linker (ca. 10%, Figure 1A). The diffusion of the complexes from the bead within the gel increased their accessibility and the surface area where the potential CAT mimics are present. To screen the peptidyl metal catalysts according to their CAT activity, a protocol inspired from the functional assay used to characterize CAT on electrophoretic gels was implemented.⁵²

Figure 1. (A) Peptide library on bead. Beads are Tentagel macrobeads. Two linkers were introduced: 10% of 4-nitrophenylalanine cleaved under UV irradiation and 90% of 4-(hydroxymethyl)benzoic acid (HMBA) cleaved in basic conditions. (B) Scheme of peptidyl complexes screening using a colorimetric activity-based assay.

The gel was incubated with a solution of H_2O_2 (0.015 %) for 10 min and subsequently with two ferric complexes, $\text{K}_3\text{Fe}(\text{CN})_6$ and FeCl_3 . In the presence of H_2O_2 , $\text{K}_3\text{Fe}(\text{CN})_6$, an orange-yellow solution, is reduced to $\text{K}_4\text{Fe}(\text{CN})_6$, which then reacts with FeCl_3 to give Prussian blue ($\text{K}_3\text{Fe}^{\text{III}}(\text{Fe}^{\text{II}}(\text{CN})_6)$), which has a deep blue color.⁵² The medium around the bead is depleted in H_2O_2 when a CAT mimic is present, because of the fast dismutation of H_2O_2 . This prevents the reduction of $\text{K}_3\text{Fe}(\text{CN})_6$ and the formation of Prussian blue. A colorless area around the bead thus indicates that an active CAT mimic is present. It is worth mentioning that when the beads were not incubated in the copper solution, no positive results were obtained. This supports the fact that if iron complexes were formed from the $\text{K}_3\text{Fe}(\text{CN})_6$ and FeCl_3 , they are not active for H_2O_2 dismutation. Using this procedure, sixteen beads from the pool of 600 tested were selected, removed from the gel, and extensively washed with EDTA to eliminate the copper. The peptides were then cleaved from the resin and analyzed by MALDI-TOF mass spectrometry (Figure 1B). To enable the unambiguous elucidation of the peptide sequence using a single mass spectrometric analysis, the method developed by Griesinger *et al.*,⁵³ adapted by Imperiali *et al.*⁵⁰ which we used previously⁴³ was employed (Figure. S1). Among the sixteen beads collected, the primary sequence could be determined for twelve peptides (Table S1, A). Interestingly, at positions X_2 and X_3 , coordinating amino acids were mainly found with histidine being the more often observed amino acid at position X_2 and tyrosine at position X_3 . On the contrary, at position X_6 non-coordinating amino acids were mainly observed (Table S1, B). Importantly, the peptide sequence PHYKHRLH (called CATm1) was found twice and a third, with a sequence (PHYKHGLH) that differs only by the non-coordinating amino acid at position X_6 was also found. CATm1 was selected for further investigations.

Hit complex characterization

The CATm1- Cu^{2+} complex was first investigated in solution in a MOPS buffer (50 mM), a fairly low complexing buffer, at pH 7.5 by Electron Paramagnetic Resonance (EPR) spectroscopy. Frozen solutions with different CATm1: Cu^{2+} ratios ranging from 1:0.5 to 1:5 were prepared and recorded at low temperature ($T = 110$ K). The EPR spectrum obtained in the presence of 0.5 equivalent of Cu^{2+} , where the observed parameters were $g_{\parallel} = 2.26 > g_{\perp} = 2.06$ and $A_{\parallel} = 0.0178 \text{ cm}^{-1}$, indicates a square planar geometry (Figure 2A).⁵⁴⁻⁵⁶ This spectrum suggests a coordination site very similar to the copper coordination site of mixed N,O ligand.^{57,58} Upon addition of Cu^{2+} , the intensity of the signal increased and a second set of signals appeared gradually. This suggests that a first complex may be formed and that above one equivalent, a second copper ion is also bound (Figure 2A). A titration performed in MOPS buffer (50 mM, pH 7.5) by UV-vis circular dichroism spectroscopy confirmed this hypothesis: upon addition of Cu^{2+} , the bands at 230 nm and at 260 nm increased up to two equivalents and reached a plateau (Figure 2B).

The key parameter for an efficient catalysis of H_2O_2 dismutation is the two-electron redox potential of the CAT. The redox potential must lie between the potentials of the $\text{O}_2/\text{H}_2\text{O}_2$ and $\text{H}_2\text{O}_2/\text{H}_2\text{O}$ couples that are 0.28 V and 1.35 V vs. NHE, respectively.⁵⁹ The CATm1- Cu^{2+} complex was examined by cyclic voltammetry in MOPS buffer (50 mM) at pH 7.5. For a 1:2 CATm1: $\text{Cu}(\text{II})$ ratio, the mixture exhibited an

anodic potential that differs from those of Cu^{2+} and CATm1 in solution, providing evidence that at least one complex is formed (Figure 3A and S4). The non-Nernstian shape of the waves is more likely due to a slow electron transfer, which is often the case for peptidyl copper complexes.^{60,61} A titration was performed by electrochemistry (Figure 3A and B). Upon addition of Cu^{2+} , the current corresponding to the anodic potential (at ca. 0.57 V vs. NHE) increased linearly up to ca. 2 equivalents (Figure 3C left). Interestingly, after 2 equivalents, the slope was modified (Figure 3C), showing that after a 1:2 ratio, the system behaved differently. The continuous increase of the current after 2 equivalents may be attributed to MOPS-Cu complexes. In addition, above ca. 1:2 ratio, the current at the potential corresponding to anodic re-dissolution of Cu^0 adsorbed onto the electrode appeared at ca. 0.24 V/NHE, highlighting the fact that the peptide was unable to bind more than two copper cations (Figure 3B yellow to red). The titration was also performed in TRIS buffer (50 mM, pH 7.5), which is considered to bind copper with a higher affinity than MOPS buffer.^{62,63} Interestingly, the same trend was observed except that the peak corresponding to the anodic redissolution appeared before 2 equivalents, probably because of the presence of a TRIS-Cu complex (Figure S5).

Figure 2. (A) Electron Paramagnetic Resonance spectra in MOPS buffer (50 mM, pH 7.5, 25°C, glycerol 10%) of a mixture CATm1: $\text{Cu}(\text{OAc})_2$ at different ratios, $[\text{CATm1}] = 200 \mu\text{M}$, $T = 110\text{K}$. The inset shows a zoom with the new set of signals indicated with stars. Note that $\text{Cu}(\text{OAc})_2$ was silent in our conditions as already reported.⁶⁴ (B) CATm1 titration with Cu^{2+} by UV-vis circular dichroism in MOPS buffer (50 mM, pH 7.5, 25°C), $[\text{CATm1}] = 200 \mu\text{M}$, $T = 25^\circ\text{C}$. The inset shows the evolution of the two maxima at 230 and 260 nm upon Cu^{2+} addition. The plateau is reached after two equivalents of Cu^{2+} .

All together these results suggest that at pH 7.5, CATm1 is able to bind 2 copper(II) cations with different affinities. The reduction and oxidation potentials of the complex were measured at 0.12 V/NHE and 0.57 V/NHE, respectively, and are thus in the correct range to catalyze H_2O_2 dismutation. In the same way, the titration performed by cyclic voltammetry at pH 7.0 in MOPS buffer (50 mM) indicated that, at this pH, a 1:2

complex was also formed with a second site showing a weaker affinity constant (Figure S6).

Figure 3. Cyclic voltammetry in MOPS buffer (50 mM, pH 7.5, 25°C). (A) Voltammograms of CATm1 (1 mM), CATm1:Cu²⁺ 1:2 (1 mM) and Cu(OAc)₂ (2 mM). (B) Titration of CATm1 (1 mM) by Cu(OAc)₂ (C) Evolution of the current at the anodic potential 0.57 V vs NHE (left) and at the potential corresponding to the anodic re-dissolution (0.24 V, right). Working electrode: Glassy carbon (3 mm diameter), counter electrode: Platinum, reference electrode: SCE. Scan rate of 100 mV/s, the small arrow indicates the scanning direction and arrows in bold, the evolution of the waves upon addition of Cu(OAc)₂.

Dinuclear complex stability

In order to obtain further insight into the stability of this 1:2 complex, the apparent dissociation constants were determined by isothermal titration calorimetry (ITC) at pH 7 and 7.5 in MOPS buffer (50 mM). At pH 7, the thermogram obtained was successfully fitted with a “2 independent sites” model leading to apparent dissociation constants of about $K_{d1} = 1.0 \pm 0.3 \cdot 10^{-6}$ and $K_{d2} = 1.1 \pm 0.2 \cdot 10^{-5}$, thus confirming that the peptide is able to bind two copper(II) ions with a second site exhibiting a weaker affinity constant (Figure S7, Table 1). The first complexation reaction has a large favorable enthalpic component ($\Delta H_1 = -47.2 \pm 0.7 \text{ kJ mol}^{-1}$) and an unfavorable entropic component ($-\Delta S_1 = 12.9 \pm 0.1 \text{ kJ mol}^{-1}$), whereas it is the opposite for the second site ($\Delta H_2 = 14.0 \pm 0.9 \text{ kJ mol}^{-1}$ and $-\Delta S_2 = -42.3 \pm 0.6 \text{ kJ mol}^{-1}$) leading to overall Gibbs free energies of -34.3 and $-28.3 \text{ kJ mol}^{-1}$ for the first and the second site, respectively. The first complexation reaction is more favorable than the second one which is entropically driven

(Figure S7, Table 1). At pH 7.5, Cu(OAc)₂ titration measured by ITC did not allow for the measurement of the binding constants of the two sites (Figure S8 A).

Table 1. Thermodynamic parameters for complexation of Cu(OAc)₂ with CATm1 measured using ITC (average of two independent experiments $\pm$ SEM) at pH 7.0, fluorescence spectrometry at 25°C (average of three independent experiments $\pm$ SEM) and mass spectrometry at pH 7.5. Conditional dissociation constants were calculated considering competition with the buffer (see SI).⁶⁵

	pH 7.0 MOPS	pH 7.5	
		MOPS	NH ₄ HCO ₂
1st Site			
n_1	0.88 ± 0.08		
ΔH_1 (kJ mol ⁻¹)	-47.2 ± 0.7		
ΔS_1 (J mol ⁻¹ K ⁻¹)	-43.3 ± 0.4		
$^{app}K_{d1}$	$1.0 \pm 0.3 \cdot 10^{-6}$	$2.8 \pm 0.6 \cdot 10^{-6}$	$2.5 \cdot 10^{-9[a]}$
$^{cond}K_{d1}$	$4.8 \pm 1.4 \cdot 10^{-9}$	$7.9 \pm 0.5 \cdot 10^{-9}$	$2.5 \cdot 10^{-9[a]}$
2nd Site			
n_2	1.04 ± 0.04		
ΔH_2 (kJ mol ⁻¹)	14.0 ± 0.9		
ΔS_2 (J mol ⁻¹ K ⁻¹)	142 ± 2		
$^{app}K_{d2}$	$1.1 \pm 0.2 \cdot 10^{-5}$	$8.5 \pm 4.9 \cdot 10^{-6}$	$1.0 \cdot 10^{-8[a]}$
$^{cond}K_{d2}$	$5.3 \pm 1.0 \cdot 10^{-9}$	$2.5 \pm 1.4 \cdot 10^{-8}$	$1.0 \cdot 10^{-8[a]}$

[a] Apparent dissociation ($^{app}K_d$) constants used to fit the data obtained by ESI-MS at 25°C. Refer to Fig. 4 for more details.

In order to get insights onto the weaker binding site, ITC experiments were performed starting from a 1:1 CATm1:Cu(OAc)₂ mixture. The stoichiometry extracted from the binding isotherm was about 0.25 (Figure S8 C to E), thus indicating that the heat exchanges do not correspond to the complexation of the second copper ion. This suggests that at 1:1 CATm1:Cu(OAc)₂ ratio, a mixture of complexes is already present in solution, and thus that the two binding sites have affinities too close to be distinguished in ITC. It has been recently shown that mass spectrometry can reliably and accurately provide both speciation and binding constants for non-covalent protein complexes and peptidyl or proteyl metal complexes.^{66–69} Consequently, a titration by electrospray ionization mass spectrometry (ESI-MS) was performed in ammonium formate (10 mM) at pH 7.5 to obtain the relative binding constants. Upon addition of copper(II) acetate to a solution of CATm1 (20 μ M), complexes with two stoichiometries corresponding to CATm1:Cu 1:1 and 1:2 were identified. After each addition of Cu(II), the ESI-mass spectrum was measured. Representative spectra are given in Figure 4A. The relative abundances of CATm1, CATm1:Cu 1:1, and CATm1:Cu 1:2 were normalized at each step in the ESI-MS titration and the mole equivalents of Cu(II) bound were calculated from the mass spectra to generate the speciation as a function of Cu(II) added (Figure 4B). The speciation curves obtained were simulated using Hyperquad Simulation and Speciation (HySS) to afford the relative stepwise apparent dissociation constants for the first copper(II) (K_{d1}) and the second copper(II) (K_{d2}) bound to the peptide. The corresponding apparent dissociation constants used to fit the data in Figure 4 were in the range of 10^{-8} – 10^{-9} with the first association constant being 4 times higher than the second. In the absence of a standard, the absolute values of these K_d are not determined. The speciation curves obtained using CuSO₄ instead of Cu(OAc)₂ were fitted using these constants and matched perfectly, confirming the reproducibility of the relative binding constants from the normalized ESI-mass spectral data (Figure S11). In addition, it shows that the copper counter-ion does not influence the stability of the complexes. In order to determine the apparent values of

these dissociation constants, titrations were performed by fluorescence spectroscopy in a MOPS which is a competitive buffer. Indeed, a clear quenching of tyrosine fluorescence was observed upon addition of copper acetate as it has been already observed for others peptides that bind copper(II).⁶⁵ The titration curves were successfully fitted with a 1-to-2 ligand-to-metal complex model using HypSpec software leading to apparent stepwise dissociation constants of $2.8 \pm 0.6 \cdot 10^{-6}$ and $8.5 \pm 4.9 \cdot 10^{-6}$ for the first and second site, respectively (Figure S12). K_{a1} is 3 times greater than K_{a2} , which is consistent with the relative constants found by mass spectrometry. In addition, by considering the competition with MOPS,⁶⁵ the conditional dissociation values fall in the 10^{-8} - 10^{-9} range and are thus in line with the values used for the fits of the mass spectrometry data.

Figure 4. Titration of CATm1 (20 μ M) by copper acetate monitored by electrospray ionization mass spectrometry in ammonium formate (10 mM) at pH 7.5. a) Representative normalized spectra at ratio CATm1:Cu(II) 1:0, 1:0.3, 1:1, 1:1.8 from top to bottom. b) Normalized abundance of each species calculated from the mass spectra during the titration. The experimental speciation was fitted by a simulated speciation using Hyperquad Simulation and Speciation (HySS) at the same concentration. From the simulated speciation, HySS provides relative cumulative (β) binding constants for the first (β_1) and second (β_2) Cu(II) binding to the peptide. From these cumulative binding constants, the relative stepwise binding constants (K_A) for the first (K_1) and second (K_2) Cu(II) bound to the peptide were determined with K_1 being 4 times larger than K_2 , respectively. In the absence of a standard, the absolute values of K_A were not determined.

Catalytic activity in solution

In order to determine the intrinsic catalase-like activity of the redox-active peptidyl complex derived from CATm1, O_2 formation was monitored in solutions containing H_2O_2 and the catalyst. O_2 was measured by polarography using a Clark-type electrode.²⁹ This was performed at CATm1:Cu(II) 1:2 ratio and with two copper salts ($CuO(Ac)_2$ and $CuSO_4$) leading to apparent kinetic constants (Figure S13). The measurement of the initial rates $v_0 = k_{cat}[catalyst]_0[H_2O_2]_0$, which can be written $v_0 = k_{obs}[catalyst]_0$ under pseudo first order conditions depending on the catalyst concentration, enabled the determination of k_{obs} for O_2 apparition. The measured k_{obs} for the 1:2 peptide:Cu²⁺ ratio are higher than for the copper salts, which indicates that the peptidyl copper complex is a better catalyst than the copper salt for H_2O_2 dismutation (Table 2). The O_2 formation reaction from H_2O_2 follows a Michaelis-Menten catalytic behavior, as does the natural enzyme.⁷⁰ Lineweaver-Burk double reciprocal plot allows k_{cat} and K_M determination since the x intercept corresponds to $-1/K_M$ and the slope to K_M/V_{max} with $V_{max} = k_{cat}[catalyst]_0$ (Figure S14). The measured k_{cat} is for O_2 evolution and thus corresponds to half of k_{cat} of H_2O_2 dismutation (owing the stoichiometric coefficients in $2 H_2O_2 \rightarrow O_2 + 2 H_2O$).

Table 2. Parameters describing the catalysis of H_2O_2 dismutation in MOPS (50 mM, pH 7.5, 25°C).

	$k_{obs}^{[a]}$ (s ⁻¹)	K_M (M)	$k_{cat}^{[b]}$ (s ⁻¹)	$k_{cat}/K_M^{[b]}$ (M ⁻¹ .s ⁻¹)	Ref.
$CuSO_4$	$3.2 \cdot 10^{-3}$				This work
CATm1: $CuSO_4$ 1:2	$2.0 \cdot 10^{-2}$	$1.9 \cdot 10^{-2}$	$7.5 \cdot 10^{-2}$	4.0	This work
$Cu(OAc)_2$	$7.4 \cdot 10^{-3}$				This work
CATm1: $Cu(OAc)_2$ 1:2	$1.3 \cdot 10^{-2}$	$2.1 \cdot 10^{-2}$	$1.2 \cdot 10^{-1}$	5.5	This work
Catalase	$1.1 \cdot 10^3$				This work
$Cu(N-baa)_2(phen)^{[c]}$		$5.2 \cdot 10^{-2}$	$6.6 \cdot 10^{-2}$	1.3	33
$[Cu(HL)](ClO_4)_2^{[d]}$		$1.7 \cdot 10^1$	$1.5 \cdot 10^{-3}$	$8.9 \cdot 10^{-5}$	35
$Cu_2(pxdiprbrtaccn)Cl_4^{[e]}$		1.5	1.24	0.8	40
Catalase		$8.3 \cdot 10^{-2}$	$2.6 \cdot 10^5$	$3.1 \cdot 10^6$	70

T. thermophiles

[a] k_{obs} are pseudo first order constant for O_2 evolution measured with $[H_2O_2] = 4.9$ mM.

[b] k_{cat} are first order rates for H_2O_2 dismutation and K_M the Michaelis-Menten constants. In this work, they have been measured with CATm1:Cu²⁺ 1:2 mixture at 100 μ M.

[c] N-baaH: N-benzoylanthranilic acid, phen: 1,10-phenanthroline. The reactions were performed in DMF at 20°C

[d] HL: 1,3-bis[(2-aminoethyl)amino]-2-propanol. The reactions were performed at 25°C in TRIS buffer, pH 7.2.

[e] pxdiprbrtaccn: 1,4-Bis(4,7-diisopropyl-1,4,7-triazacyclonon-1-ylmethyl)benzene. The reactions were performed in phosphate buffer (0.01 M, pH = 7.4) at 25°C.

The apparent second order constant (k_{cat}/K_M) for the 1:2 complexes are similar, whatever the copper salt used, and are higher than the values reported for other copper complexes and in particular for those measured in aqueous solutions, where it is more challenging to measure this activity (Table 2). However, the apparent second order constants of CATm1:Cu 1:2 complexes are much lower than those of purified natural catalase, showing that there is still room for improvement. Additionally, the turnover number (TON), the turnover frequency (TOF), and the maximum yield of H_2O_2 dismutation were calculated and are reported in Table S2 (SI). The TON is about 3 which is slightly weaker than for most manganese porphyrins but higher than for many other Mn complexes with CAT-like activities.²⁹ During O_2 monitoring by polarography, when O_2 evolution reached a plateau, the dismutation of H_2O_2 could have been again fasten by the addition of fresh catalyst (Figure S15). This suggests that the catalyst was inactivated

during the process, which in turn, may explain the low calculated TON. This has been confirmed by mass and UV-vis spectrometry experiments (Figure S16 and S17). By ESI mass spectrometry, upon addition of H_2O_2 (20 equivalents), catalyst oxidation and fragmentation were rapidly observed (Figure S16). UV-vis spectrometry enabled the determination of the pseudo first order rate of catalyst degradation, which was found one order of magnitude slower than pseudo first order rate of H_2O_2 dismutation (Figure S17). This clearly supports

the fact that the catalyst can cycle several times before being inactivated. To note, as already reported, the 1:1 complex has been investigated for its SOD activity in cuvette using the McCord and Fridovich assay.⁴³ In this assay, the IC_{50} of the 1:1 complex was 372 nM which was only 1.6 time better than CuSO_4 (IC_{50} = 603 nM). This shows that this complex has a very weak SOD activity. This illustrates that the strategy allowed for the selection of an efficacious catalyst for the target-activity only.

Figure 5. H_2O_2 levels in HeLa Hyper cells monitored by fluorescence for 60 min after treatment with H_2O (control cells), CATm1 peptide (250 μM), CuSO_4 (500 μM), complex CATm1- Cu^{2+} 1:2 (250 μM). The right panel represents the mean of the ratio for 2 or 3 independent experiments.

Catalytic activity in cells

To go further, this new catalyst was assayed on HeLa cells which express the protein HyPer intracellularly. HyPer is a genetically encoded ratiometric fluorescent sensor of H_2O_2 based on a circularly permuted yellow fluorescent protein (cpYFP) integrated into the regulatory domain of the bacterial H_2O_2 sensing protein OxyR (OxyR-RD).⁷¹ In presence of H_2O_2 , thiols of HyPer are oxidized, leading to the formation of disulfide bridges that modify the conformation of the protein and thus its spectral properties. The ratiometric modification of the excitation spectrum of HyPer can consequently allow monitoring H_2O_2 levels by measuring the ratio $I_{(491/530)}/I_{(405/530)}$.^{72–74} This ratio was monitored for 60 min for several cells (typically 10 to 25 cells) before and after addition of CATm1:Cu 1:2 mixture at 250 μM and the corresponding controls (CATm1 at 250 μM , or CuSO_4 at 500 μM) in MOPS buffer (30 mM, pH 7.5). To note, no additional H_2O_2 was added, the observed effects are thus modifications of the basal level of H_2O_2 . Since H_2O_2 is mainly produced at the membranes and can easily diffuse through,⁷⁵ an efficient catalase mimic that remains outside the cells may induce a decrease of the H_2O_2 concentration inside the cells. This is the case of HeLa HyPer cells incubated with catalase that does not translocate into cells but leads to a fluorescence ratio decrease.⁷⁶ We have previously reported the use of Hyper HeLa to quantify the activity of a range of Mn-based anti-oxidants.⁷⁴ The intensity ratios ($I_{(491/530)}/I_{(405/530)}$) normalized against the values from the cells before the addition of the tested compounds are reported in Figure 5. As shown in Figure 5, the H_2O_2 concentration decreased rapidly after addition of the 1:2 complex and

was then stabilized around 28% after 40 min. For the CATm1 peptide alone, a slight decrease of the fluorescence ratio was also observed, but after 20 min the H_2O_2 concentration increased to the initial level. Addition of CuSO_4 led to a decrease of the H_2O_2 concentration, but only by 19%. Significantly, these data strongly support the fact the 1:2 complex is efficacious for H_2O_2 dismutation in a cellular context.

In conclusion, this innovative strategy combining a combinatorial approach with an activity-based screening of the peptidyl metal complexes library led to the discovery of an efficacious di-Cu(II) complex with CAT activity. As far as we know, this is the first peptidyl complex mimicking the anti-oxidant properties of CAT reported so far. Its characterization was performed using a range of techniques (CD, ITC, CV, fluorescence and ESI-MS), and its activity was determined in solution using a Clark electrode and in a cellular assay. The cellular assay results are very encouraging as many potential metal ligands exist in the cell culture media that could have competed with the peptide and could have led to its destruction. The biological activity of this class of complexes may be improved if a higher stability could be achieved and work towards this goal is currently in progress in the lab.

EXPERIMENTAL SECTION

General

All chemicals and solvents were of synthesis grade and were used as received without further purification. Common solvents for solid support synthesis and L-amino acids were obtained from either Sigma Aldrich, Novabiochem or Iris biotech GMBH. CATm1 peptide was synthesized on an Advanced ChemTech automated synthesizer at the

IBPS SU, FR3631 peptide synthesis core facility and purified by HPLC as described.

MALDI-TOF mass spectra were recorded on a Voyager DE-Pro MALDI-TOF mass spectrometer (Applied Biosystems) in positive mode using as matrix a solution of α -Cyano-4-hydroxycinnamic acid (CHCA) at 10 mg/mL in $\text{CH}_3\text{CN}:\text{H}_2\text{O}$ (50:50) containing 0.1% TFA. Calibration was performed using external standards (Proteomix 4, LaserBio Labs Sofia-Antipolis, France). Electrospray ionization mass spectrometry was performed on a Bruker MicrOTOF II with the following parameters: capillary voltage= 4200 V, Nebulizer= 2.0 Bar, Dry gas = 4.0 L/min, Dry Temp = 200 °C, Capillary exit voltage = 100 V. Analytical HPLC was performed on an Agilent Technologies 1200 Series equipped with a multiple wavelength absorbance detector, using a Proto 200 C18 3 μm Higgins Analytical Inc. 100 x 4.6 mm column. Preparative HPLC was performed on a Waters 600 HPLC Pump equipped with a Waters 2487 dual wavelength absorbance detector, using a Column VP 260/16 C18 Htec 5 μm . UV-vis spectrometry was performed on a Cary 300 bio in a double beam mode with the buffer in the reference cell at 25°C. Electron Paramagnetic Resonance (EPR) data were recorded using an Elexsys E500 Bruker spectrometer, operating at a microwave frequency of approximately 9.44 GHz. Spectra were recorded using a microwave power of 20 mW across a sweep width of 250 mT (centered at 310 mT) with modulation amplitude of 0.5 mT. Experiments were carried out at 110 K using a liquid nitrogen cryostat. Circular dichroism (CD) spectra (215-300 nm) were collected on a J-810 spectropolarimeter (Jasco, Tokyo, Japan). Cyclic Voltammetry measurement were performed using a MetrOhm potentiostat (AUTOLAB model). All measurements were carried out in the indicated buffer. Glassy carbon (GC), a platinum plate and SCE KCl saturated were used as working electrode, counter and reference electrode, respectively; all potential values are reported versus NHE reference. Isothermal titration calorimetry (ITC) experiments were performed on a TA Instrument (New Castle, DE) NanoITC calorimeter and analyzed using the provided software. The ability of the complex to catalyze H_2O_2 dismutation was measured by polarography. Titrations by fluorescence were performed on a Jasco FP-8300 spectrofluorimeter. An O_2 -sensitive Clark-type electrode OD InLab 605-ISM purchased from Mettler Toledo was used. Prior to measurements, the electrode was calibrated in ambient air then in a provided 0% dissolved oxygen solution.

Procedures and data analysis

Peptide library synthesis on solid support (Figure S1)

Linkers coupling and first amino acid grafting. The peptides library was synthesized manually on solid support by using standard Fmoc chemistry on 1 gram of Tentagel macrobeads (loading: 0.24 mmol/g, 280-300 μm). First, orthogonal linkers (6 equiv.) coupling was performed on a solution of HMBA and 3-N-Fmoc-amino-3-(2-nitrophenyl)propionic acid (Fmoc - ANP) linkers (10:1) with HOBt (1 equiv.) and DIC (1 equiv.) in DMF for 1h. Standard deprotection conditions of Fmoc group were employed (20% piperidine in DMF for 1 min then 10 min at room temperature) followed by washings with NMP. Then the first amino acid was introduced by treatment with the symmetrical anhydride of β Ala (6 equiv.) in the presence of 4-dimethylaminopyridine (0.1 equiv.) in DMF for 1h.

Amino acid coupling reactions (without capping). Coupling reactions were performed in NMP (3 mL) with HOBt (3 equiv.) and HBTU (3 equiv.) as coupling agents and in basic conditions (diisopropylethylamine, DIEA, 6 equiv.). The resin was shaken for 45 to 60 min at room temperature. After each reaction, solvents and soluble reagents were removed under vacuum and the resin was washed 5 times with NMP. Completion of the reaction was monitored by the Kaiser test, which indicates the presence of free amine by a blue coloration of the bead. The amino acid coupling was repeated as many times as necessary until a negative test Kaiser was obtained. Then, Standard deprotection conditions of Fmoc group were employed (20% piperidine in DMF for 1 min then 10 min at room temperature) followed by washings with NMP (5 x 3 mL).

Amino acid coupling reactions (with capping). By using Bibilo software, the positions in the library that required capping were identified, then manual calculation of truncated peptide molar mass allowed for the determination of which commercially available capping

reagent combination was required to obtain a non-degenerate mass ladder. The twelfth coupling required simultaneous capping since it is a mass encoding step. Coupling then involved treatment with a mixture of the desired amino acid and mass cap (85:15, 10 equiv.), N,N-diisopropylcarbodiimide (DIC), and N-hydroxybenotriazole (HOBt) in 10:10 ratio in DMF for 1 hour. The mass caps were: BocD for E, BocW for H, BocF for Y, BocI for R and BocA for G.

Split and pool strategy. The ninth, twelfth and the thirteenth coupling were performed after splitting the resin. The resin was split into 5 approximately equal portions. Then, each batch reacted with a different amino acid using standard coupling conditions. After the reaction, all five batches were pooled together and the next coupling was performed.

A final acetylation was performed on the last amino acid (Proline) by a solution of acetic anhydride in dichloromethane (DCM) (3 mL, 10/90 v:v) for 1 h at room temperature. Solvents and soluble reagents were removed by filtration. The resin was washed with DCM (5 x 3 mL) then by methanol (3 x 3 mL). The resin was dried under vacuum for 1h. Finally, deprotection of the lateral chains was carried out in acidic conditions with TFA/ H_2O /Trisopropylsilane (95%/2.5%/2.5%) for 2 h at room temperature. The resin was then washed sequentially by TFA, DCM and HEPES buffer.

Complexes formation

Beads (ca. 50 mg) were inserted in PP syringe with a frit and were washed with HEPES (50 mM, pH 7.5) solution (3 x 1 mL). The beads were incubated for 2h at 60°C with 2 mL of a $\text{Cu}(\text{OAc})_2$ solution (10 mM) in HEPES (50 mM, pH 7.5). After incubation, the solution was removed by filtration and the beads were washed with HEPES (50 mM, pH 7.5, 1 x 1 mL), EDTA (25 mM, 1 min, 1 x 1 mL) then again with HEPES (50 mM, pH 7.5, 3 x 1 mL) solution to increase the selectivity pressure.

Combinatorial screening and selection

The beads (approximately 50-100) were introduced in a small beaker and molten agarose (10 mL, 2% in HEPES 50 mM, pH 7.5) was added. The solution was poured into a Petri dish, so that the beads were evenly distributed in the dish. The solution was kept at room temperature for 10 minutes to allow the formation of a gel. The agarose gel was then placed under a UV lamp (electronic dual light transilluminator) at 254 nm during 10-15 min to cleave the photocleavable linker. The gel was then incubated with a H_2O_2 solution (10 mL, 0.015%, 10 min) then rinsed extensively with H_2O milliQ. The water was poured off. A 2% FeCl_3 solution (0.6 mg in 30 mL H_2O milliQ) and a 2% potassium ferricyanide solution (0.6 mg in 30 mL H_2O milliQ) were poured together directly on top of the gel. The solution became blue and the gel was incubated for 3 minutes then extensively washed by H_2O milliQ. Beads with white halos were cut from the agarose gel with a tip of a 1 mL pipette and transferred to a 1.5 mL eppendorf tube. The residual agarose surrounding the selected bead was melted away in distilled H_2O (1 mL) at 110°C. The selected bead was subsequently washed with ethylenediaminetetraacetate (0.5 M; pH 7.0), then extensively with H_2O for the metal decoordination. Peptides were cleaved from the bead by treatment with NH_4OH (28%, 50 μL) overnight at room temperature. After cleavage, the solution was freeze-dried, and the peptide residue resuspended in an H_2O milliQ (100 μL) as a library stock solution.

Sequence determination by MALDI-TOF mass spectrometry

1 μL of peptide stock solutions was mixed with 1 μL of matrix solution and 1 μL of the mixture solution was placed onto the MALDI plate and allowed to crystallize. Peptides sequencing was carried out with the Software developed by C. Griesinger et al.⁵³ (Figure S4).

CATm1 purification

The peptide was purified by reverse phase HPLC on a VP260/16 semipreparative column with a linear gradient from 5 to 30% acetonitrile in water both containing 0.1% TFA over 30 min. High purity (> 95%) was confirmed by analytical HPLC (Proto 200 C18 column from 5% to 100% acetonitrile in water and 0.1% TFA over 10 min) and the expected mass was found by MALDI-TOF mass spectrometry: $[\text{M}+\text{H}]^+ = 1128.4$ (Figure S2 and S3).

Peptide stock solution preparation

Stock solution of the peptide was prepared by dissolving peptide powder in milliQ water. Stock solution concentration was determined by measuring the UV-vis spectrum of a diluted solution (1 μ l in 449 μ l of milliQ water) and using the extinction coefficient of the tyrosine Tyr considered as free tyrosine ($\epsilon_{276} = 1280 \text{ cm}^{-1}\text{M}^{-1}$).

Electron Paramagnetic Resonance spectroscopy (EPR)

EPR experiments were carried out at 110 K using liquid nitrogen cryostat. EPR samples (200 μ l) were prepared from stock solution of peptide diluted down to 0.2 mM in MOPS buffer (0.5 M, pH 7.5). Cu^{2+} was added from 50 mM stock solution of CuSO_4 in water to form the complexes. Samples were frozen in an ethanol/liquid nitrogen solution in quartz tube after addition of 10% glycerol as a cryoprotectant and stored in liquid nitrogen until used.

Titration by circular dichroism (CD)

Spectra of solutions of CATm1 (200 μ M) in MOPS buffer (50 mM, pH 7.5) were recorded before and after successive additions of CuSO_4 . A 1 mm cell was used and the spectra were recorded with a Peltier temperature controller set at 298 K under constant nitrogen flush. The wavelength range was set from 300 nm to 215 nm. The scan rate, the sensitivity, and bandwidth were respectively 200 $\text{nm}\cdot\text{min}^{-1}$, 0.125 s, and 1 nm. Each spectrum was an average of three scans.

Cyclic voltammetry experiments

All measurements were carried out in a buffer (MOPS or TRIS at pH 7.5 or 7). Glassy carbon (GC), a platinum plate and SCE KCl saturated were used as the working electrode, counter and reference electrode, respectively. All potential values are reported versus the SCE reference electrode. Dissolved dioxygen was removed by bubbling the solution with dinitrogen gas. Between each measurement the working electrode was washed with distilled water and polished with sand paper.

Isothermal titration calorimetry (ITC)

Titration at pH 7: titrations were performed by injecting 10 μ l aliquots of 5 mM $\text{Cu}(\text{OAc})_2$ in MOPS buffer (50 mM; pH 7) into the calorimeter cell containing the peptide CATm1 solution (ca 0.5 mM) diluted in MOPS buffer (50 mM; pH 7), with 5 min between two injections. The experiments were performed at 25 $^{\circ}\text{C}$. Data were analyzed using the program NanoAnalyze provided by TA Instruments.

Titration at pH 7.5: titrations were performed by injecting 10 μ l aliquots of 5 mM or 10 mM $\text{Cu}(\text{OAc})_2$ in MOPS buffer (50 mM; pH 7.5) into the calorimeter cell containing the peptide CATm1 solution (ca 0.5 mM) or 1:1 CATm1: $\text{Cu}(\text{OAc})_2$ mixture (at 1 or 2 mM) diluted in MOPS buffer (50 mM; pH 7), with 5 min or 10 min between two injections. The experiments were performed at 25 $^{\circ}\text{C}$. Data were analyzed using the program NanoAnalyze provided by TA Instruments. (See SI for more details).

Titration by electrospray ionization mass spectrometry

1 mM $\text{Cu}(\text{OAc})_2\cdot\text{H}_2\text{O}$ was titrated into 20 μ M CATm1 peptide solution in a 10 mM ammonium formate solution at pH 7.5. After each addition of $\text{Cu}(\text{II})$, the sample was measured using electrospray ionization mass spectrometry. The abundance of apo CATm1, CATm1- Cu_1 , and CATm1- Cu_2 were normalized from each step in the ESI-MS titration. The amount of $\text{Cu}(\text{II})$ bound was calculated from the mass spectra to generate the speciation as a function of $\text{Cu}(\text{II})$ added. This speciation was simulated using Hyperquad Simulation and Speciation (HySS). The same concentrations of peptide and $\text{Cu}(\text{II})$ as the experimental data were used in the HySS simulation. 2.0 mol. eq. of $\text{Cu}(\text{II})$ were added in the simulated titration. HySS uses cumulative (β) binding constants and the simulated cumulative constant for the first (β_1) and second (β_2) $\text{Cu}(\text{II})$ binding to the peptide were adjusted until the simulation fit the experimental data. From these cumulative binding constants, the stepwise binding constants for the first (K_1) and second (K_2) $\text{Cu}(\text{II})$ bound to the peptide were found to be $10^{8.6}$ and 10^8 , respectively. ESI-mass spectral data have been reported to accurately represent the speciation following metal binding into various protein systems.^{68,69,77,78} It appears that the binding of metals typically does not impart significant differences in the ionization efficiency which allows for the quantitation of the species present. With respect to the results presented in this paper, the binding constants were calculated from a simulation of the data in Figure 4. These same binding constants were used to fit the results of a separate titration

where 5 mM CuSO_4 was titrated into 20 μ M CATm1 in 0.5 mM ammonium formate buffer at pH 7.56. The quality of the fit shown in Figure S11 is essentially identical to the fit shown in Figure 4, confirming the reproducibility of the binding constants from the normalized ESI-mass spectral data. In addition, the amount of copper bound to the peptide can be determined from the ESI-mass spectral data and this compared to the mol. equivalences of $\text{Cu}(\text{II})$ added to the solution (Figure S9). The ratio of $\text{Cu}(\text{II})$ added to $\text{Cu}(\text{II})$ recovered is nearly 1:1 indicating that the CATm1 speciation obtained by ESI-MS is representative of the solution speciation.

Titration by fluorescence spectrometry.

$\text{Cu}(\text{OAc})_2\cdot\text{H}_2\text{O}$ was titrated into CATm1 peptide solution in a 50 mM MOPS buffer at pH 7.5 at 25 $^{\circ}\text{C}$. After each addition of $\text{Cu}(\text{II})$, the emission spectrum (average of 2 accumulations) upon excitation at 275 nm was recorded between 280 and 400 nm ($\text{slit}_{\text{exc}} = \text{slit}_{\text{em}} = 5 \text{ nm}$, scan rate = 200 nm/min). Before recording each spectrum, it was ensured that the thermodynamic equilibrium was reached (stable fluorescence intensity). The intensity at the maximum of tyrosine emission (303 nm) was used to generate the titration curves which were then fitted using HypSpec software. HypSpec determines cumulative (β) binding constants for the first (β_1) and second (β_2) $\text{Cu}(\text{II})$ binding to the peptide. From these cumulative binding constants, the stepwise binding constants for the first (K_1) and second (K_2) $\text{Cu}(\text{II})$ bound to the peptide were found. The provided K_d values are the average of three independent titrations $\pm$ standard error of the mean (SEM).

Catalase activity assay using a Clark-type electrode

The experiments were carried out in MOPS buffer (50 mM, pH 7.5) at 25 $^{\circ}\text{C}$ in a micro cell sealed with a rubber septum to avoid the introduction of O_2 from the air. The dissolved dioxygen was removed by bubbling the solution with dinitrogen gas. Hydrogen peroxide solution (15%, 1 μ l) was added. The dioxygen from the spontaneous dismutation of hydrogen peroxide was removed by bubbling the solution with dinitrogen gas. Finally, sample was injected through the septum to the stirred H_2O_2 solution. Reaction rates were determined by measurement of the O_2 evolved upon time as described in reference.²⁹ The initial rates v_0 were measured from the slope of linear fits (from 0 to 20 s) of dioxygen formation at several complex concentrations. Then k_{obs} corresponds to the slope of the linear fit of $v_0 = f[\text{catalyst}]$ (Figure S11). The O_2 formation reaction from H_2O_2 follows a Michaelis-Menten catalytic behavior similar to the natural enzyme. Michaelis-Menten constants K_m , V_{max} and k_{cat} were determined using the Lineweaver-Burk double reciprocal plot: $1/v_0 = f(1/[\text{H}_2\text{O}_2])$ for which the x intercept corresponds to $-1/K_m$, the slope to K_m/V_{max} and $V_{\text{max}} = k_{\text{cat}} [\text{O}_2][\text{catalyst}]_0$. Since during dismutation two H_2O_2 lead to the formation of a single O_2 molecule $k_{\text{cat}} = 2 \times k_{\text{cat O}_2}$. To generate these plots, the initial rates at a catalyst concentration of 100 μ M were measured at several $[\text{H}_2\text{O}_2]$ concentrations (Figure S12). Complementary kinetics parameters were determined at 100 μ M of the catalysts in 4.9 mM H_2O_2 solution in MOPS (50 mM) at pH 7.5 (Table S2). The maximal yield of O_2 production (in %) was calculated from the measured $[\text{O}_2]_{\text{max obs}}$ values using the following equation: $\text{Yield O}_2 (\%) = (2[\text{O}_2]_{\text{max obs}} / [\text{H}_2\text{O}_2]_0) \times 100$. The TON was calculated as maximal number of O_2 moles produced per mole of a catalyst, $\text{TON} = [\text{O}_2]_{\text{max obs}} / [\text{Catalyst}]_0$. The TOF values in s^{-1} which represent the ratio of initial rates and concentrations of catalysts were calculated as followed: $\text{TOF} = v_0 / [\text{Catalyst}]_0$.

Experiments on HeLa HyPer cells

Cell culture. Stable cell line HeLa HyPer1 was prepared using the HeLa Flp-In cell line, which was kindly provided by Stephen Taylor⁷⁹ and cultured at 37 $^{\circ}\text{C}$ under a 5% CO_2 atmosphere. HyPer1 expression in this stable cell line was controlled by doxycycline, added 24 h after seeding. The cells were cultured for an additional 24 h before being processed for analysis.

Pharmacological treatments. Cells were incubated with or without antioxidant during 1 hr at 37 $^{\circ}\text{C}$ under a 5% CO_2 atmosphere.

Imaging. Imaging was performed with a CSU-W1 Yokogawa spinning disk coupled to a Zeiss Axio Observer Z1 inverted microscope equipped with a sCMOS Hamamatsu camera and a 63 $\times$ objective (63 $\times$ /1.4 oil WD: 0.17 mm) oil objective. DPSS 100 mW 405 nm and

150 mW 491 nm lasers and a 525/50 bandpass emission filter were used.

H₂O₂ levels quantification and statistical analysis. Images were processed with the Fiji software, to obtain the HyPer1 ratio of the emission at 530 nm (491/530)/(405/530). HyPer ratio was then measured for several cells, and normalized to the ratio value of the control condition. Data were analyzed using GraphPad Prism 8 and expressed as the mean $\pm$ standard error of the mean (SEM). Statistical significance was calculated using an ordinary one-way ANOVA followed by Tukey's multiple comparison test.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website.

Supplementary Scheme, figures, and tables, detailed experimental procedures, MALDI-TOF spectra, ITC thermograms, HPLC chromatograms, fluorescence images are provided.

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All authors have given approval to the final version of the manuscript.

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Notes

The authors declare no competing financial interest.

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Efficiently selected! A combinatorial approach associated with an activity-based screening led to the discovery of the first peptidyl di-copper complex mimicking Catalase activity. This complex accelerates H₂O₂ dismutation into O₂ and H₂O in test tubes and in cells.