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Link Between Affinity and Cu(II) Binding Sites to Amyloid- β Peptides Evaluated by a New Water-Soluble UV-Visible Ratiometric Dye with a Moderate Cu(II) Affinity

Amandine Conte-Daban^{#a,b}, Valentina Borghesani^{#a,b}, Stéphanie Sayen^c, Emmanuel Guillon^c, Yves Journaux^{d,e}, Geoffrey Gontard^{d,e}, Laurent Lisnard^{d,e}, and Christelle Hureau^{a,b,*}

^{a,b}CNRS; LCC (Laboratoire de Chimie de Coordination) ; 205, route de Narbonne, F-31077 Toulouse, France. Université de Toulouse; UPS, INPT ; LCC ; F-31077 Toulouse, France

^cUniversité de Reims Champagne Ardenne, Institut de Chimie Moléculaire de Reims (ICMR), UMR 7312 CNRS-URCA, Moulin de la Housse, BP 1039, 51687 Reims Cedex 2, France

^dSorbonne Universités, UPMC Univ. Paris 06, UMR 8232, IPCM, F-75005, Paris, France

^eCNRS, UMR 8232, Institut Parisien de Chimie Moléculaire, F-75005, Paris, France

[#] These authors contributed equally to this work.

Abstract

Being able to easily determine the Cu(II) affinity for biomolecules of moderate affinity is important. Such biomolecules include amyloidogenic peptides, such as the well-known amyloid- β peptide involved in Alzheimer's disease. Here, we report the synthesis of a new water soluble ratiometric Cu(II) dye with a moderate affinity (10^9 M^{-1} at pH 7.1) and the characterizations of the Cu(II) corresponding complex by X-ray crystallography, EPR and XAS spectroscopic methods. UV-Vis competition were performed on the A β peptide as well as on a wide series of modified peptides, leading to an affinity value of $1.6 \cdot 10^9 \text{ M}^{-1}$ at pH 7.1 for the A β peptide and to a coordination model for the Cu(II) site within the A β peptide that agrees with the one mostly accepted currently.

Copper ions play key biological roles.¹ They are essential metal ions that play an important role as catalytic centers in several processes including very fundamental ones like energy production.¹ They have also been linked to several diseases. Dyshomeostasis of Cu is very dangerous and is well documented by two lethal genetic diseases, called Wilson's and Menkes' diseases, linked to an overload of Cu and to a depletion of Cu, respectively.^{2–5} Cu imbalance has also been involved in the etiology of most of neurological disorders⁶ such as Alzheimer's disease (AD),^{7–9} Parkinson's disease (PD),^{10,11} Prion diseases.^{10,12}

At a molecular scale, two parameters are important regarding Cu and the peptides or proteins involved in the above-mentioned diseases: (i) the metal environment, i.e. the nature of the atoms surrounding the metal center and (ii) the affinity of the peptides for the metal

*Corresponding Author: christelle.hureau@lcc-toulouse.fr.

center. In the context of AD which is under focus in the present article, there are many reports on the coordination sites of the Cu(II) center to the amyloid- β peptide (A β) (for recent reviews, see refs.8,13–15) and on the Cu(II) affinity for A β (see refs.16–18 and refs. therein). Some studies also aim at relating the Cu(II) affinity of modified A β peptides with the coordination sites.19–21

Determining the affinity of Cu(II) for peptides is thus of interest, first because this is intrinsically an important parameter that mirrors the possibility to have the metal ion-peptide interaction in biological conditions, and second because it give insights into the coordination sphere of the metal center when a series of modified peptides is studied. As a direct consequence, the straightforward and accurate determination of Cu affinity for peptides is an important objective. However, this is an intricate task since it necessitates appropriate analytical tools. The complexity of such studies is well illustrated by the abundant and differing reports on the evaluation of Cu(II) affinities for the A β peptide (see refs.16–20 and refs. therein). This is also true for the +I redox state of Cu,18,22–24 which is however not under focus in the present study. While Cu(II) affinity values ranging from 10^6 M^{-1} to 10^{19} M^{-1} have been proposed,25 a relative consensus has recently been reached around a value of 10^{10} M^{-1} at pH 7.417,18 (this value corresponds to the conditional affinity value, i.e. the absolute affinity value at a given pH ; the interaction with buffer is not considered and will lead to a weaker so-called apparent affinity value).25 Hence the objective of the present study is to define a new tool appropriate for the rapid, easy and accurate evaluation of Cu(II) binding affinities to peptides.

There are mainly two ways of determining affinity values.16 The first one relies on potentiometric titrations that lead to absolute constants, from which an apparent affinity value can be calculated at any pH values.16,26 The second one consists in measurements at a given pH either by (i) isothermal calorimetry (ITC), fluorescence when a fluorophore is natively present in the peptide sequence or added for purpose and when the fluorescence is quenched by the addition of the metal ion of interest or (ii) by competition experiments with a ligand of known affinity followed by the appropriate method. Potentiometry is likely the most powerful technique since it gives access to the affinity value at any pH values but could be extremely difficult to implement in case of biomolecules and requires important quantity of the biological material. Results from ITC are far from straightforward although additional data apart from the affinity can be evaluated (enthalpy, entropy...) (reviewed in ref. 16). Competition experiments appear to be a well-suited method. With respect to Cu(II), UV-Vis and fluorescent spectroscopic tools are of interest. In a seminal paper, Wedd and coworkers have designed four peptide-based fluorophores of various Cu(II) affinities ranging from 10^8 M^{-1} to 10^{14} M^{-1} . They were further used to probe the affinity of Cu(II) for different peptides and proteins by competition monitored by fluorescence.27 While Cu(II) quenches the fluorescence of the competitor, the addition of the peptide of interest restores it, removing the Cu(II) from the probe.

In the present paper, we propose an alternative strategy that relies on a UV-visible competition experiment against an easy to synthesize Cu(II) dye with moderate affinity ($3.2 \cdot 10^9 \text{ M}^{-1}$ at pH 7.1) that is anticipated to be perfectly appropriate for evaluation of Cu(II) affinity to weakly structured peptide, such as A β , at physiological pH. By UV-Vis, the

addition of the peptide of interest on the colored Cu(II) complex induces the disappearance of the absorption bands.

Using this new dye, we have investigated more than seventeen peptidic sequences derived from the human A β peptide, including murine, N-terminally modified peptides and biologically relevant mutants. Their Cu(II) affinities were evaluated and discussed with respect to the coordination models previously proposed in the literature.^{13,21,28–30}

Experimental Section

Chemicals

Reagents were commercially available and were used as received. All the solutions were prepared in milliQ water (resistance: 18.2 M Ω .cm).

The Cu(II) ion source was CuSO₄·5H₂O, bought from Sigma-Aldrich. A stock solution was prepared at 25 mM.

HEPES buffer (sodium salt of 2-[4-(2-hydroxyethyl)piperazin-1-yl]ethanesulfonic acid) was bought from Sigma-Aldrich. A stock solution was prepared at 500 mM, pH = 7.3 in order to reach a resulting pH 7.1 and the same buffer stock solution was used over the course of the present study.

Peptides

A β ₁₆ (DAEFRHDSGYEVHHQK) and the mutants:

Ac-A β ₁₆, Ac-DAEFRHDSGYEVHHQK; D1N-A β ₁₆, NAEF-RHDSGYEVHHQK; E3Q-A β ₁₆, DAQFRHDSGYEVHHQK; D7N-A β ₁₆, DAEFRHNSGYEVHHQK; E11Q-A β ₁₆, DAEFRHDSGYQVHHQK; H6R-A β ₁₆, DAEFRRDSGYEVHHQK; H6A-A β ₁₆, DAEFRADSGYEVHHQK; H13A-A β ₁₆, DAEFRHDSGYEVAHQK; H14A-A β ₁₆, DAEFRHDSGYEVHAQK; mA β ₁₆, DAEFGHDSGFVRRHQK; R5G-A β ₁₆, DAEFGHDSGYEVHHQK; Y10F-A β ₁₆, DAEFRHDSGFVHHQK; H13R-A β ₁₆, DAEFRHDSGYEVRRHQK; R5G-H13R-A β ₁₆, DAEFGHDSGYEVRRHQK; A β ₂₈, DAEFRHDSGYEVHHQKLVFFAEDVGSNK; A β ₄₀, DAEFRHDSGYEVHHQKLVFFAEDVGSNKGAILGLMVGGVV were bought from GeneCust (Dudelange, Luxembourg) with purity grade > 95%.

Stock solutions of the peptides were prepared by dissolving powder in milliQ water (resulting pH ~ 2), except for the A β ₂₈ and A β ₄₀ peptides which were dissolved in NaOH 50 mM. Peptide concentration was determined by UV-Vis absorption of Tyr10 considered as free tyrosine (at pH ~ 2.0, $\epsilon_{276} - \epsilon_{296} = 1410 \text{ cm}^{-1} \cdot \text{M}^{-1}$; at pH ~ 12.5, $\epsilon_{293} - \epsilon_{360} = 2400 \text{ cm}^{-1} \cdot \text{M}^{-1}$).³¹ For the mutants without Tyr10, the absorption of the two Phe was used ($\epsilon_{258} - \epsilon_{280} = 390 \text{ cm}^{-1} \cdot \text{M}^{-1}$).³²

As the Cu(II) coordination sites in the A β peptide are localized in the first 16 amino acids residues,⁸ the A β ₁₆ and its counterparts were used as a model of the full-length peptides.

ABH and BAH where B corresponds to the β -alanine were bought from Protéogénix (Strasbourg, France) and stock solutions were prepared by solubilizing the peptides in milliQ water. Their concentrations were determined by UV-Vis titration with a titrated Cu(II) solution following the d-d band absorption of the complex,³³ according to ref. 34.

Ligand

The 3,4-bis(oxamato)benzoic acid ligand, **L** (see Scheme S1), was synthesized as described below and used in its sodium salt form. A 8.5 mM stock solution was prepared, increasing the pH until solubilization, determined by UV-Vis titration with a titrated Cu(II) solution following the band absorption at 330 nm of the complex (see text below and Figure 3), according to ref. 34.

Synthesis of H_5L — $\text{H}_3\text{Et}_2\text{L}$: 3,4-(diethyloxamate)benzoic acid:

The ester form of the **L** ligand was prepared following the known procedure for oxamate ligands^{35,36} instead of the previously reported synthesis.³⁷ To 5 g of 3,4-diaminobenzoic acid (32.8 mmol) in THF (200 mL) is added with strong stirring 8.25 mL of ethyl oxalyl chloride (72 mmol, 2.2 eq). The mixture is refluxed for 2 hours, filtered while hot and let to cool down to room temperature. Removal of the solvent leads to a brown precipitate. The solid is abundantly washed with water, filtered, washed with 50 % v. ethanol and then with a minimum of cold 96 % v. ethanol, and dried in air. Yield: 8.9 g (77 %). See Supporting Info for elemental analysis, NMR and IR characterizations.

Aqueous NaOH (2 M, 57 mL) was added dropwise to a suspension of $\text{H}_3\text{Et}_2\text{L}$ (8 g, 22.7 mmol) in water (400 mL). The solution was stirred for 30 min and filtered. Addition of 4 M HCl (28 mL) to the resulting solution leads to the rapid formation of a brown precipitate. After 10 min the precipitate is collected on a sintered glass filter, washed with cold water, then with 96 % v. EtOH, and finally with ether before being dried for several hours in air and overnight at 45°C in an oven. Yield: 5.4 g (64%). See Supporting Info for elemental analysis, NMR and IR characterizations.

Synthesis of $\text{Li}_3[\text{Cu}(\text{L})]$ —Suitable crystals of the copper dye, $\text{Cu}(\text{L})$, were obtained as follow: A suspension of $\text{H}_3\text{Et}_2\text{L}$ (0.100 g, 0.28 mmol) in water (10 mL) was treated with 0.71 mL of LiOH 2 M (1.42 mmol) and stirred at room temperature until complete dissolution of the ligand. An aqueous solution of $\text{Cu}(\text{NO}_3)_2\cdot 3\text{H}_2\text{O}$ (0.068 g, 0.28 mmol, 5 mL) was then added dropwise to the oxamate solution resulting in a deep blue solution of the copper complex that was stirred for 30 min at room temperature and filtered. Slow diffusion of acetone yields crystals of $\text{Li}_3[\text{Cu}(\text{L})]\cdot 5.5\text{H}_2\text{O}$ in 15 days. Yield: 0.054 g (41 % based on Cu). See Supporting Info for elemental analysis, NMR and IR characterizations.

Methods

X-Ray structure

Crystallographic data were collected on a Bruker Kappa-APEX II CCD diffractometer (CuK α , λ = 1.54178). Crystal data: purple plates, monoclinic $C2/c$, a = 26.3214(15), b = 6.7217(4), c = 20.4183(11) Å, β = 112.590(3)°, V = 3335.3(3) Å³, Z = 8, T = 200(1) K, ρ = 1.567 g.cm⁻³, $F(000)$ = 1560, $\mu_{\text{CuK}\alpha}$ = 2.293 mm⁻¹. Crystals were mounted on a Hamilton cryoloop using Paratone-N oil and placed in the cold flow produced with an Oxford Cryocooling device. Partial hemispheres of data –preselected with the APEX II software³⁸ were collected using ϕ and ω scans. Integrated intensities were obtained with SAINT+ and were corrected for absorption with SADABS;^{38,39} the structure was solved with SIR9240 and refined with SHELXL-2014/741 (WinGX software package).⁴² Data refinement gives (using 271 parameters and 6 restraints) $wR2$ = 0.2432 (2923 unique reflections), R_I = 0.0720 (2514 reflections with $I > 2\sigma(I)$), GOF = 1.128. Crystallographic details are available in CIF format, free of charge via www.ccdc.cam.ac.uk/conts/retrieving.html (or from the Cambridge Crystallographic Data Centre, 12, Union Road, Cambridge CB2 1EZ, UK; fax: (+44) 1223-336-033; or deposit@ccdc.cam.ac.uk). CCDC number 1516020.

NMR, IR and Elemental Analysis

¹H and ¹³C NMR spectra were collected on a 400 MHz Bruker Avance spectrometer at 298 K.

ATR/FT-IR spectra were collected on a Bruker TENSOR 27 equipped with a simple reflexion ATR diamond plate of the Harrick MPV2 series.

Elemental analysis was performed at the ICSN, CNRS UPR 2301, in Gif-sur-Yvette, France.

Electron Paramagnetic Resonance

Electron Paramagnetic Resonance (EPR) data were recorded using an Eleksys E 500 Bruker spectrometer, operating at a microwave frequency of approximately 9.5 GHz. Spectra were recorded using a microwave power of 20 mW across a sweep width of 150 mT (centred at 310 mT) with modulation amplitude of 0.5 mT. Experiments were carried out at 110 K using a liquid nitrogen cryostat.

EPR samples were prepared from stock solution of ligand diluted down to 0.2 mM in H₂O. 0.9 eq. of ⁶⁵Cu(II) was added from 25 mM ⁶⁵Cu(NO₃)₂ stock solution home-made from a ⁶⁵Cu foil. Samples were frozen in quartz tube after addition of 10% glycerol as a cryoprotectant and stored in liquid nitrogen until used.

X-ray Absorption Spectroscopy (XAS)

Cu(II) K-edge EXAFS (Extended X-Ray Absorption Fine Structure) and XANES (X-ray Absorption Near-Edge Structure) spectra were recorded at the BM30B (FAME) beamline at the European Synchrotron Radiation Facility (ESRF, Grenoble, France).⁴³ The storage ring was operated in 7/8+1 mode at 6 GeV with a 200 mA current. The beam energy was selected using a Si(220) N₂ cryo-cooled double-crystal monochromator with an experimental

resolution close to that theoretically predicted (namely ~ 0.5 eV FWHM at the Cu energy). The beam spot on the sample was approximately $300 \times 100 \mu\text{m}^2$ (H x V, FWHM). Because of the low Cu(II) concentration, spectra were recorded in fluorescence mode with a 30-element solid state Ge detector (Canberra) in frozen liquid cells in a He cryostat. The temperature was kept at 20 K during data collection. The energy was calibrated with Cu metallic foil, such that the maximum of the first derivative was set at 8979 eV. EXAFS Cu(II) data were collected from 8840 to 8960 eV using 5 eV step of 2 s, from 8960 to 9020 eV using 0.5 eV step of 3 s, and from 9020 to 9300 eV with a k-step of $0.05 \text{ \AA}^{-1}$ and 3 s per step. At least six scans recorded on different spots were averaged. XAS samples were prepared from stock solutions of ligand and Cu(II) diluted down to approx. 1.0 mM in buffered solution. Samples were frozen in the sample holder after addition of 10% glycerol as a cryoprotectant and stored in liquid nitrogen until used. Cu(II) photoreduction was controlled by recording successive scans at the same spot. It was considered that during the first 20 minutes of recording the photoreduction is insignificant.

The data analysis was performed using the “Multi-Platform Applications for X-ray Absorption” package, including Cherokee and Roundmidnight programs,⁴⁴ according to the standard and previously reported data analysis procedures.^{45,46} Spectra were background-corrected by a linear regression through the pre-edge region and a polynomial through the post-edge region. The backscattering phase, $\Phi_i(k, R_i)$, and amplitude, $A_i(k, R_i)$, functions were obtained using the ab initio FEFF7 code.⁴⁷ Since theoretical phase shifts were used, it is necessary to fit the energy threshold E_0 by adding an extra fitting parameter, E_0 . Moreover, the FEFF7 code was used to check if the multiple scattering of our reference compounds of known crystallographic structure is negligible in the 0–3 $\text{\AA}$ range. The estimated errors for distances and coordination numbers are $\pm 0.02 \text{ \AA}$ and $\pm 20\%$, respectively.

UV-Visible spectrophotometry

UV-vis spectra were recorded on a spectrophotometer SPECORD S600 Analytik Jena at 25°C in 1 cm path length quartz cuvette.

Competition experiments between the ligand and the peptides—The experiments have been monitored by UV-Vis in HEPES buffer 0.1 M (at a resulting pH = 7.1). The ligand (50 μM in theory) and Cu(II) (45 μM in theory) were mixed and successive additions of the different peptides (0 to 500 μM in total) were added. Each addition of the peptide is made once the thermodynamic equilibrium of the previous reaction is reached (this takes about 20 min). To be reproducible with respect to pH condition, the same stock buffer solution (pH = 7.3 to reach a resulting pH of 7.1) was used for all the experiments. The competition experiments were performed at 25°C in triplicate.

Analysis of the data—The data analysis was performed with a 2-step procedure for each experiment (corresponding to a given peptide).

Step 1. The theoretical concentrations of the ligand L and the Cu(II) were adjusted using the absorbance of ligand at 246 nm ($\epsilon_{246} - \epsilon_{450} = 18000 \text{ cm}^{-1} \cdot \text{M}^{-1}$) and the absorbance of the Cu(L) complex at 330 nm ($\epsilon_{330} - \epsilon_{450} = 18000 \text{ cm}^{-1} \cdot \text{M}^{-1}$) respectively.

Step 2. The absorbance of the competition experiments measured at 330 nm after completion of the reaction was plotted as a function of the number of peptides equivalents recalculated with respect to the real ligand concentration (see step 1). Then these data were fitted with an in-house procedure using the adjusted concentrations as starting parameters.

Absorbance was calculated according to:

$$Abs = \varepsilon_{330\text{ nm}}^L \cdot [L] + \varepsilon_{330\text{ nm}}^{CuL} \cdot [CuL] + \varepsilon_{330\text{ nm}}^P \cdot [P] + \varepsilon_{330\text{ nm}}^{CuP} \cdot [CuP]$$

where P stands for peptide, L for the ligand, and CuP and CuL for the complex between Cu(II) and peptide and the ligand, respectively.

The same equation can be expressed as:

$$Abs = \varepsilon_{330\text{ nm}}^L \cdot ([L]_0 - [Cu]_0 + [\alpha]) + \varepsilon_{330\text{ nm}}^{CuL} \cdot ([Cu]_0 - [\alpha]) + \varepsilon_{330\text{ nm}}^P \cdot ([P]_0 - [\alpha]) + \varepsilon_{330\text{ nm}}^{CuP} \cdot [\alpha]$$

where α stands for the progression of the following reaction : peptide + Cu(II) $\rightarrow$ peptide-Cu(II)

and the $[]_0$ correspond to the starting values in which the dilution due to addition of the peptide has been taken into account when required.

Cu(II) is supposed to be chelated by the peptide or by L, there is no free Cu(II).

$$\frac{K_d^{CuP}}{K_d^{CuL}} = \frac{[P] \cdot [Cu]}{[\alpha]} \cdot \frac{[CuL]}{[L] \cdot [Cu]}$$

With the starting concentrations:

$$\frac{K_d^{CuP}}{K_d^{CuL}} = \frac{[P]_0 - [\alpha]}{[\alpha]} \cdot \frac{[Cu]_0 - [\alpha]}{[L]_0 - [Cu]_0 + [\alpha]}$$

This is a quadratic equation having the form

$$a\alpha^2 + b\alpha + c = 0, \text{ with}$$

$$\alpha = \frac{-b + \sqrt{\Delta}}{2a}, \text{ with}$$

$$b = \frac{K_d^{CuP}}{K_d^{CuL}} \cdot ([L]_0 - [Cu]_0) + [P]_0 + [Cu]_0$$

$$\Delta = b^2 - 4.a.c$$

$$a = \frac{k_d^{CuP}}{k_d^{CuL}} - 1$$

$$c = -[P]_0 \cdot [Cu]_0$$

Then, the K_d^{CuP} value was adjusted to obtain the best reproduction of the experimental data.

NB: Although K_d/K_a are dimensionless values, they are given here in M or M⁻¹, respectively, for convenience. Indeed for 1:1 system, the K_d value in M gives a direct idea at which concentration the complex is predominantly formed.

NB: in a preliminary experiment, K_d^{CuL} was evaluated by competition with ABH and BAH knowing $K_d^{Cu(ABH)}$ and $K_d^{Cu(BAH)}$ 33 with the similar procedure.

Results

The competitor used in the present work to evaluate the Cu(II) affinity of a wide series of Aβ derived peptides by UV-Vis competition experiments, is an oxamate ligand (H₅L, Scheme S1, for the sake of clarity L⁵⁻ will be written L. Similarly, the [Cu(L)]³⁻ will be noted Cu(L)). Oxamate ligands have largely proved their appeal for the design of metalloligands and the subsequent preparation of multidimensional compounds with controlled architectures and properties.⁴⁸ In particular, the use of oxamate ligands and the multi-step assembly of oxamate-based complexes have successfully led to relevant objects in molecular magnetism: switchable molecules, high nuclearity coordination complexes, single-chain magnets, porous magnets.^{49,50} Recently oxamate-based systems have also shown attractiveness in broader coordination chemistry fields such as catalysis.^{51,52} Here, we will take advantage of the mild Cu(II) affinity and of the presence of an intense UV-Visible band for the complex with Cu(II) (λ = 330 nm, ε = 18000 cm⁻¹.M⁻¹).

The X-ray structure of Cu(L) is shown in Figure 1 and the crystallographic parameters are given in Table 1, top, and in the Supporting Info. The complex adopts a square-planar geometry where the L ligand coordinates the Cu(II) ion via its two oxamate groups each binding the metal center through one nitrogen and one oxygen atom. The square-planar geometry is fairly regular with a 2.4° dihedral angle between the oxamate groups and a 3.1° dihedral angle between the Cu(II) coordination plane and the phenyl plane of the ligand. The

carboxylate group of the ligand remains uncoordinated. Supramolecular arrangements of the complexes are described in the Supporting Information (Figure S1).

The persistence of the structure in aqueous solution has been probed by several complementary techniques. The most appropriate one is EXAFS by which the Cu-N and Cu-O distances have been determined in solution and are in perfect agreement with those obtained for the crystal structure (for crystallography, see Table 1, bottom and Figure S1, for EXAFS fitting, see Figure S2).

The parameters estimated from the EPR signature (Figure 2A) are also in line with the Cu(L) solid-state structure. Indeed, the $g_{\parallel}$ and $A_{\parallel}(^{65}\text{Cu})$ values equal 2.22 and $212 \times 10^4 \text{ cm}^{-1}$, respectively and are thus in line with a 2N2O equatorial binding mode according to the Peisach and Blumberg correlation.⁵³ A weak rhombicity linked to the presence of four non equivalent equatorial ligands is at the origin of the complex features observed in the $g_{\perp}$ region (coupling between $I_{\text{Cu}} = 3/2$ with $S_{\text{Cu}} = 1/2$ for the two g_x and g_y transitions). The presence of two equivalent nitrogen atoms is consistent with the five super-hyperfine lines observed on the first parallel transition, arising from the coupling between $I_{\text{N}} = 1$ with $S_{\text{Cu}} = 1/2$. The XANES spectrum of Cu(L) is given in Figure 2B. It is in line with Cu(II) complexes where the Cu(II) center lies in a square planar environment.¹

The other important parameter of this ligand is its affinity constant for Cu(II). This affinity was evaluated by the same type of competition experiments than those detailed below, performed first between L and two short peptides, namely ABH and BAH (B stands for β -alanine).³³ These peptides were studied by potentiometry to determine the impact of the 5-membered versus 6-membered metallacycles with regard to the Cu(II) affinity in an ATCUN-like (Amino Terminal CU and Nickel) sites.⁵⁴ Their conditional Cu(II) affinity values calculated at pH 7.1 from the potentiometric data equal: 3.0×10^8 and $3.7 \times 10^9 \text{ M}^{-1}$, respectively. Competition experiments with the ABH (*resp.* BAH) lead to a Cu(II) affinity value of $2.2 \times 10^9 \text{ M}^{-1}$ (*resp.* $4.1 \times 10^9 \text{ M}^{-1}$) for L. The average value of $K_a^{\text{L}} = 3.2 \pm 1.0 \times 10^9 \text{ M}^{-1}$ was used in the rest of the study.

The competition experiment between L and the A β peptides for Cu(II) binding were performed by using the peptides to remove the Cu(II) from the colored Cu(L) complex leading to a disappearance of the signature at 330 nm characteristic of the Cu(L) complex. Figure 3A shows the typical series of spectra obtained for a given peptide (here A β_{16}). L was first added (dotted line), then Cu(II) (solid line) and then successive additions of the peptide were performed (grey lines, See Experimental Section for more details). Note that the increase at 276 nm is due to Tyr10 absorbance and perfectly matches the amount of peptide added.

The absorbance at $\lambda = 330 \text{ nm}$ of the mixture was plotted as a function of the equivalent numbers of peptide added (dots in Figure 3B). Then, these data were fitted with an in-house procedure (dashed lines, for more details see Experimental section).

Experimental absorbance values and their fits are shown in Figure 3B as a matter of illustration for A β_{16} , H6A-A β_{16} and Ac-A β_{16} , (curves a, b, c, respectively). Qualitative

analysis indicates that the Cu(II) affinity of the three peptides follow the trend: $A\beta_{16} > H6A-A\beta_{16} > Ac-A\beta_{16}$. All the Cu(II) affinity values obtained in the present study are gathered in Table 2 while the corresponding reproductions of the experimental data are given in the Supporting Info (Figure S3). The accuracy of the fitting procedure is shown in the Figure S4. All the calculated Cu(II) affinity values were normalized with respect to the value of $A\beta_{16}$ for an easier comparison, as shown in Table 2 and in Figure 4.

Discussion

The $A\beta$ peptides used in this study were modified with respect to the human sequence on any potential binding amino-acids, namely the carboxylate containing amino-acid residues, the His residues and the N-terminal amine. The mutation involved between the human and murine sequences were also studied.

Human-derived peptides

The first modification is the acetylation of the N-terminal amine (Asp1) precluding the possibility of Cu(II) binding. The affinity constant of Cu(II) for this modified peptide is at least 10 times weaker than that of the $A\beta_{16}$ peptide, in line with an important contribution of the N-terminal amine in Cu(II) binding to $A\beta_{16}$, as discussed in the literature.^{28,55,56} In the second group of modified peptides, the impact of the carboxylate function is evaluated. The carboxylate group of each amino acid residues Asp1, Glu3, Asp7 and Glu11 is mutated with an amide function. Only a very weak impact is observed with a decrease in the affinity upon modification that doesn't exceed a factor of 2. In the $A\beta_{16}$ there are 4 carboxylate groups available for the Cu(II) coordination. Therefore, if one carboxylate is mutated, 3 other ones remain to complete the coordination sphere. This is in perfect agreement with the possibility of having an equilibrium between the four carboxylate groups for one binding position.⁵⁷ The high similarity of affinity values for the four carboxylate-modified peptides also suggest that there is no preference for one given carboxylate group. The last group of modified peptides consist in the His mutants (His6, His13 and His14). While His6 mutation (H6A or H6R) has a significant impact (a 5-fold decrease of the affinity value), His13 or His14 mutation has a lesser effect (with a maximal 2-fold decrease of the affinity value). This indicates that (i) none of the His is essential but (ii) the His13 and His14 can more easily exchange for the binding position on the peptide than His6 and His13 or than His6 and His14 do. This is in line with the proposition that in the main coordination site His13 and His14 exchange for one binding position while His6 is constantly bound.^{13,30,58} It is worth mentioning that similar differences have been previously determined by ITC.^{19,20} All together these data fit very well with the coordination site of Cu(II) proposed based on spectroscopic data (reviewed in refs.13,30) and reminded in Scheme 1. In such proposition, the Cu(II) ion is equatorially bound to the N-terminal amine, the adjacent carbonyl group from the Asp1-Ala2 bond, the imidazole group from His 6 and from His13 or 14 in equilibrium (see Scheme S2, top).⁸ A carboxylate-containing residue has been proposed to occupy the apical position.⁵⁷ Last, the Cu(II) affinity seems to weakly increase with the length of the peptide, in line with previous reports,^{17,59} but this result has to be taken with cautious due to the increase in the error bar when working with the $A\beta_{40}$ peptide. This

confirms that the A β ₁₆ moiety is a good model with respect to Cu(II) binding site within the full-length peptide.

Human versus murine peptides

The murine sequence differs from the human one by 3 mutations: R5G, Y10F and H13R. With respect to Cu(II) affinity values, the murine peptide is twice stronger than the human one, in line with previous measurements by direct competition,^{17,29} ITC,¹⁹ and potentiometry.⁵⁵ The main mutation responsible for such a difference is the R5G mutation. This has been previously related to the formation of a 6-membered metallacycle between the side chain of His6 and the deprotonated Gly5-His6 peptide function for the murine peptide that is not observed in the human case (see Schemes 1 and S2 bottom).^{29,30}

Conclusions

In the present study, a new ratiometric dye was synthesized and fully characterized. It was further used to determine the affinity constant of Cu(II) for several A β peptides by UV-Vis competition experiments. This was made possible by the intense ($18000\text{ cm}^{-1}\cdot\text{M}^{-1}$) LMCT band displayed at $\lambda = 330\text{ nm}$ by the Cu(L) complex. The affinity values obtained in this study for the various modified peptides agree perfectly with published coordination sites, thus validating the proposed strategy that relies on the determination of affinity constants of a wide series of modified peptides to extrapolate a coordination model for a given peptide. In addition, the designed dye could be further used to straightforwardly determine Cu(II) affinity for other flexible peptides such as alpha-synuclein involved in Parkinson's disease¹¹ or Prion protein,¹² for which the Cu(II) affinities are proposed to lie in the 10^8 - 10^{10} M^{-1} range. However, outside this range, competition with L will not give reliable values. In addition, the molar extinction coefficient is relatively low implying the use of quite large biomolecule concentration. As a consequence, to generalize this approach to the evaluation of Cu(II) affinity for any other (bio)molecules relies on the ability to modulate the Cu(II) affinities and to increase the molar extinction coefficient of oxaloamino-benzoic acid derivatives by playing with the scaffold.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Figure 1.
Ellipsoid plot (H and disorders atoms are omitted for clarity) of Cu(L).

Figure 2.

Panel A. EPR signature of Cu(L). [L] = 200 μ M, [$^{65}\text{Cu(II)}$] = 180 μ M, [HEPES] = 50 mM, pH = 7.1, T = 120 K. Inset: superhyperfine structure on the first parallel line. *Panel B.* Normalized Cu K-edge X-ray Absorption Near-Edge Structure (XANES) spectrum of Cu(L). [L] = 1 mM, [Cu(II)] = 0.9 mM, [HEPES] = 0.1 M, pH = 7.0, T = 20K.

Figure 3.

Panel A. UV-Vis spectra of a solution of L (dotted line), in the presence of 0.9 eq. of Cu(II) (solid line) and after addition of increasing amounts (approx.. 1, 3, 7 and 10 eq.) of Aβ₁₆ peptide (grey lines). The arrows indicate the variation of the UV-Vis spectra upon addition of the peptide. The increase at 276 nm is due to Tyr10 absorption. *Panel B.* Experimental absorbance at 330 nm (dots) and the best fits (dashed lines) of the Cu(L) system upon addition of peptides (Aβ₁₆ (a), H6A-Aβ₁₆ (b) and Ac-Aβ₁₆ (c)). [L] = 50 μM, [Cu(II)] = 45 μM, [peptides] = 0 to 500 μM, [HEPES] = 0.1 M, pH = 7.1, T = 25°C.

Figure 4.

Bar graph of the normalized affinity constants in log units of the A β peptides and some of their mutants for Cu(II). The reference for the normalization is the hA β ₁₆. The error bars indicate the standard deviation. [L] = 50 μ M, [Cu(II)] = 45 μ M, [peptides] = 0 to 500 μ M, [HEPES] = 0.1 M, pH = 7.1, T = 25°C. The affinity constant of the ligand is $K_a = 3.2 \pm 1.0 \times 10^9 \text{ M}^{-1}$.

Table 1
Crystallography distances between Cu(II) and their neighbours (top) and EXAFS
parameters (bottom): N = number of neighbours, R = absorber-neighbour distance, σ =
Debye-Waller factor.

N	Distance (Å)
Cu-N ₁	1.9061(55)
Cu-N ₂	1.9127(37)
Cu-O ₁	1.9588(34)
Cu-O ₄	1.9642(51)
N	Distance (Å)
2 Cu-N	1.92
2 Cu-O	1.97
σ^2 (Å ²)	R factor
0.00471	0.18 %

Table 2
Cu(II) affinity values of the A β peptides studied here. The reference for the normalization is the A β ₁₆. A Cu(II) affinity value of $3.2 \pm 1.0 \cdot 10^9 \text{ M}^{-1}$ for the competing ligand L was taken.

Peptide	$K_a (10^9 \text{ M}^{-1})^a$	$K_a(\text{peptide})/K_a(\text{hA}\beta_{16})$
A β ₁₆	1.6	1.0
A β ₂₈	1.8	1.1
A β ₄₀	2.2	1.4
Ac-A β ₁₆ ^b	< 0.1	< 0.1
D1N-A β ₁₆	1.1	0.7
E3Q-A β ₁₆	1.0	0.6
D7N-A β ₁₆	1.1	0.7
E11Q-A β ₁₆	1.3	0.8
H6R-A β ₁₆	0.3	0.2
H6A-A β ₁₆	0.3	0.2
H13A-A β ₁₆	0.9	0.6
H14A-A β ₁₆	0.8	0.5
mA β ₁₆	3.0	1.9
R5G-A β ₁₆	3.1	1.9
Y10F-A β ₁₆	1.2	0.8
H13R-A β ₁₆	0.7	0.4
R5G-H13R-A β ₁₆	2.7	1.7

^aThe error bars are evaluated in the S.I. (see Figure S4).

^bThe K_a value for Ac-A β ₁₆ is too weak to be thoroughly determined by this method. Only a higher value can be given.