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Metal ion mediated transition from random coil to β-sheet and aggregation of Bri2-23, a natural inhibitor of Aβ aggregation.

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Abstract

Furin-dependent maturation of the BRI2 protein generates the BRI2-23 fragment that is able to arrest the aggregation of amyloidβ, the peptide implicated in Alzheimer’s disease (AD). BRI2-23 contains cysteines at positions 5 and 22 which are likely to bind to metal ions such as Cu(I). Metal ions may play a role in the etiology of neurodegenerative disorders such as AD, and in this work we explore the metal ion induced folding and aggregation of BRI2-23 using Hg(II) and Ag(I) as spectroscopic probes with structural and ligand preferences similar to those of Cu(I), while not displaying redox activity under the experimental conditions. In general, interaction of BRI2-23 with soft metal ions changes structural properties and solution behavior of the peptide that tune to increasing metal to peptide stoichiometry. Potentiometric, $^{199m}$PAC and ESI-MS data indicate that addition of up to 0.5 equivalents of Hg(II) to BRI2-23 yields two-coordinated HgS$_2$ structure at the metal site. While free peptide is inherently unstructured, the presence of Ag(I) and Hg(II) gives rise to β-sheet formation. NMR spectroscopy supports the formation of β-sheet structure in the presence of 0.5 equivalents of Hg(II), and displays an interesting and marked change in the TOCSY spectra when increasing the Hg(II) to peptide stoichiometry from 0.5 to 0.7 equivalents, indicating the equilibrium between two structural analogues of the complex. Addition of more than 0.7 equivalents of Hg(II) give rise to line broadening, presumably reflecting aggregation. This is further supported by ThT fluorescence studies showing that BRI2-23 peptide does not aggregate over 24 hours, while addition of over 0.7 equivalent of Ag(I) or Hg(II) leads to increase of fluorescence, indicating that these metal ions induce aggregation. Thus, a model integrating all data into a coherent picture is that metal ion binding to the two thiolates gives rise to folding of the peptide into a structure that is prone to aggregation, forming aggregates with a considerable amount of β-sheet. Molecular dynamics simulations initiated with structures that agree with NMR data additionally support this model.
**Keywords:** BRI2, aggregation inhibitor, probe, Cu(I), Hg(II), Aβ.

**Introduction**

BRI2, also known as *ITM2b*, is a type II transmembrane protein of unknown function abundantly expressed in neuronal tissues. It is 266 amino acids long and can be divided into a large C-terminal extracellular domain, a transmembrane domain, and a short N-terminal cytoplasmatic domain. Mutations of the *ITM2b* gene result in expression of the protein precursors ABriPP and ADanPP.\(^1\) Like the amyloid precursor protein (APP), BRI2 is subject to enzymatic processing by a number of proteolytic enzymes, including prohormone furin-like convertases able to cleave the WT BRI2 protein near the C-terminus to produce a 23-amino acid peptide, called BRI2-23 (Figure 1). Similarly to WT BRI2, also ABriPP and ADanPP proteins undergo to proteolytic cleavage at the C-terminal region generating ABri and ADan peptides (Figure 1).\(^2,3\) These peptides constitute the hallmarks of the Familiar British Dementia (FBD) and Familiar Danish Dementia (FDD) disorders, that share many mechanistic and pathological similarities with Alzheimer’s disease (AD).\(^1\) This process is initialized by the activation of the convertase in *trans* region of the Golgi apparatus and continues once the secretory vesicle merges with the cellular membrane. Although both the Golgi lumen and extracellular space represent the exoplasmic face of the membrane, they differ in redox potential, with the lumen being more reductive environment.\(^4,5\)
Figure 1. Schematic representation of BRI2 protein, its disease related forms and the products of their enzymatic processing by furin.

The accumulation of ABri and ADan triggers a complex pathological cascade of events leading to neurodegeneration,\textsuperscript{6,7} while BRI2-23 preserves its solubility.\textsuperscript{8} Both ABri and ADan peptides exist in reduced or oxidized form due to the presence of two cysteine residues within the peptide sequence, which are able to form disulphide bridges.\textsuperscript{9} However, the consequences of the oxidation are found to be contradictory with respect to the peptides’ susceptibility to form oligomers and to aggregate.\textsuperscript{10} While intramolecular disulphide bridge promotes the β sheet formation and aggregation with subsequent fibrilation in ABri,\textsuperscript{11} the amyloid-like aggregates of ADan are mainly formed by the reduced form of peptide.\textsuperscript{10} These contradictory results imply that the upstream formation of non-fibrillar soluble oligomers rather than insoluble aggregates represent the pathogenic species cause neurodegeneration.\textsuperscript{10,12}

Contrary to the ABri and ADan behavior, the wild type peptide, BRI2-23 has no toxic effects, yielding neither oligomers nor aggregates of high order. Interestingly, BRI2-23 has been found to inhibit Aβ aggregation \textit{in vitro} and \textit{in vivo}.\textsuperscript{13} This inhibitory effect is probably
mediated by the fragment containing –FENKF– sequence, that is analogous to that found in peptidic inhibitors of amyloid aggregation (–KLVFF–). Moreover, a peptide encompassing a similar sequence has been demonstrated to interact with the glycine-zipper segment of Aβ1-40 (–G33XXXG37–) that is critical for the formation of stable β sheet structure. In addition, the BRI2-23 peptide transported to the plasma membrane, is able to interact with APP reducing the access of α-, β- and γ-secretases to their respective cleavage sites. As a result, the release of any APP metabolite, including Aβ, is reduced.

Transition metal ions, like Cu, Zn and Fe are implicated in many neurodegenerative disorders and are believed to influence either the aggregation of amyloidogenic proteins or the formation of reactive oxygen species (ROS). Metal ions are involved in the aggregation of ABri and ADan peptides as well. On the other hand nothing is known about the metal binding abilities of BRI2-23 and more important whether metal ions affect its structure and aggregation ability. BRI2-23 contains several residues, like Cys, His, Glu (Figure 1) which might interact with metal transition ions. In particular, the presence of two cysteines points out a possible Cu(I)-S2 coordination geometry and high Cu(I)-BRI2-23 affinity.

Herein, we aim at investigating the interaction between Cu(I) and BRI2-23 by combining experimental and theoretical methods. In order to elucidate the role, if any, played by the cuprous ion we used Hg(II) or Ag(I), as spectroscopic probes of Cu(I), while Molecular Dynamics calculations (MD) were performed by using Cu(I). The choice to use Hg(II) and Ag(I) as Cu(I) probes, was due to the fact that Cu(I) is a redox active metal which can easily oxidize to Cu(II). For this reason sample preparation is usually performed in inert atmosphere and in presence of reducing agents such as ascorbic acid or dithionite. Unfortunately, such procedure does not completely exclude the presence of Cu(II), which although very small can interfere with Cu(I)-BRI2-23 binding in two different ways: (i) by interacting with the peptide and by oxidizing Cys residues. Moreover, Hg(II) and Ag(I) ions
are commonly used to investigate Cu(I) sites, as demonstrated by previous studies.\textsuperscript{30-41} By applying a multi techniques approach we characterize the metal coordination sphere, the binding affinity and stoichiometry, the speciation profiles and the structural rearrangements of BRI2-23 induced by the metal ion.

**Experimental section**

*Peptide Synthesis and Purification*

Peptides were synthesized on an Activotec Activo-P11 automated peptide synthesizer, on 0.25 mmol scale using standard Fmoc chemistry.\textsuperscript{42} Fmoc-Ser(tBu)-Wang resin was used as the solid support so that the resulting peptides would have unprotected C-terminus as in natural system. Cleavage from the resin was performed for 120 min in a 90% trifluoroethanol (TFA) solution containing 5% thioanisole, 2% anisole, and 3% ethanedithiol as free radical scavengers. After precipitation with cold ether, the peptides were redissolved in water and lyophilized to obtain a fluffy off-white powder. The solid was redissolved in 10% acetic acid and purified by reversed phase HPLC on a Varian Prostar HPLC with a preparative C18 column (Varian Pursuit XRs C 18) with a semi-linear gradient of 0.1% TFA in water to 0.1% TFA in 9:1 CH\textsubscript{3}CN/H\textsubscript{2}O over 45 min. The identity of the peptides was verified by ESI (electrospray ionization) mass spectrometry.

*ThT fluorescence spectroscopy assay*

The fluorometric assay was performed on FP-6500 JASCO spectrofluorometer. The peptide Bri2-23 was analyzed in presence and in absence of 0.7 equivalents of Ag(I) and Hg(II). 50 mM phosphate buffer solutions were used to dissolve the peptide to a final concentration of 10 \textmu M. Thioflavin-T was added to all analyzed systems to give a 10 \textmu M final
concentration. All the samples were monitored immediately after the preparation and after one day of incubation at room temperature.

**Mass spectrometry**

High-resolution mass spectra were obtained on a BrukerQ-FTMS spectrometer equipped with an Apollo II electrospray ionization source with an ion funnel. The mass spectrometer was operated in the positive ion mode with the following parameters: scan range m/z 400–1600, dry gas-nitrogen, temperature 170ºC, ion energy 5 eV. Capillary voltage was optimized to 4500 V to obtain the highest S/N ratio. Changing the voltage (±500 V) did not significantly affect the optimized spectra. The samples (metal/ligand in a 1:1 stoichiometry, \( c_{\text{CuII}} = 1 \times 10^{-4} \text{M} \)) were prepared in 9:9:2 MeOH/H\(_2\)O/DMSO mixture in carbonate buffer pH 6.5. Variation of the solvent composition down to 5% of MeOH did not change the speciation. The sample was infused at a flow-rate of 3 mL/min\(^{-1}\). The instrument was calibrated externally with the Tunemix™ mixture (Bruker Daltonik, Germany) in quadratic regression mode. Data were processed by using the Bruker Compass Data Analysis 4.0 program. The mass accuracy for the calibration was higher than 5 ppm, enabling together with the true isotopic pattern (SigmaFit) an unambiguous confirmation of the elemental composition of the obtained complex.

**Circular dichroism spectroscopy**

CD spectra were recorded at Jasco J-815 spectropolarimeter and the temperature was controlled with PTC-423S temperature controller. A quartz cell with 1 cm optical path was used. The spectra range was 190-260 nm with a resolution of 0.1 nm and a bandwidth of 1 nm. A scan speed of 20 nm/min with 1 s response time was employed. Baseline spectra were subtracted from each spectrum and data were smoothed with the Savitzky-Golay method. Data were processed using Origin 7.0 spread sheet/graph package. The direct CD
measurements (θ, in millidegrees) were converted to mean residue molar ellipticity, using the relationship mean residue $\Delta \varepsilon = \theta / (33000 \times c \times l \times \text{number of residue})$. 10 µM solution of apo and metal (Ag (I) and Hg (II)) bound Bri2-23 either in phosphate buffer (50 mM) and at acidic pH (~3.0) were analyzed.

**Potentiometry**

Stability constants for protons and Hg(II) complexes of cysteine and BRI2-23 were calculated from titration curves carried out at 298K using a total volume of 1.5 cm$^3$. The metal ion concentration was $5 \times 10^{-4}$ mol×dm$^{-3}$ and the metal to ligand ratio was 1:2. Competition experiments were carried out with $5 \times 10^{-4}$ mol×dm$^{-3}$ BRI2-23 peptide in the presence of 1 equivalent of cysteine and 1 equivalent of Hg(II) after 1 hour of equilibration time was allowed before the titration started. NaOH was added from a 0.500 cm$^3$ micrometer syringe which was calibrated by both weight titration and the titration of standard materials. The pH-metric titrations were performed at 298K in 0.1 mol dm$^{-3}$ NaCl on a MOLSPIN pH-meter system using a Mettler Toledo InLab semi micro combined electrode calibrated in hydrogen concentrations using HCl. The HYPERQUAD program was used for stability constant calculations. Standard deviations were computed by HYPERQUAD and refer to random errors only. They are, however, a good indication of the importance of a particular species in the equilibrium.

**Isothermal titration calorimetry**

Calorimetric titrations of the peptide with Hg(II) ions were performed by isothermal titration microcalorimetry (ITC) using Nano-ITC Instrument (Ta Instruments, USA). Experiments were carried out at 298K in distilled water at pH 3.0. Peptide concentration was 0.5 mM (950µL sample cell), while the metal ion concentration 5.0 mM (in the syringe). Both solutions were degassed for at least 15 min. by using TA Instruments degassing station, and
then stored under nitrogen to minimize sample oxidation. Automated titrations were performed until saturation, up to Hg(II)/peptide mole ratio of about 3. Experiment was repeated three times. Heats of dilution and mixing for each experiment were measured by titrating Hg(II) solution into distilled water at pH 3.5. The effective heat of each peptide metal ion interaction was corrected for dilution and mixing effects. Heats of bimolecular interactions were obtained by integrating the peaks of each injection. The data were analyzed using the NanoAnalyze software v. 2.3.6, using “multiple sites” binding model. $\Delta H^o$ and the corresponding binding constant, $K_a$, molar free energy of binding, $\Delta G^o$, and the molar entropy change, $\Delta S^o$, were attained from the fundamental equations of thermodynamics: $\Delta G^o = -RT \ln K_a = \Delta H^o - T(\Delta S^o)$.

Perturbed Angular Correlation Spectroscopy

The following stock solutions were prepared and used for the PAC experiments: BRI2-23 (2 mM, concentration determined by the Ellman’s test),\textsuperscript{45} phosphate buffer of pH 3.0 and pH 7.5 (0.93M, and HgCl\textsubscript{2} (3.0 mM). The final samples contained 50 mM of the proper buffer, 200 $\mu$M or 100 $\mu$M BRI2-23, 100 $\mu$M of HgCl\textsubscript{2} in 55% sucrose. Water (150 mL) treated so as to lower the concentration of metal ions was placed in a small teflon cup sitting on a copper device, frozen in liquid nitrogen and mounted at the ISOLDE GLM beam line (at CERN) in a vacuum chamber. Radioactive $^{199m}$Hg was implanted in the ice typically for 1 hour. The radioactive $^{199m}$Hg was produced by irradiating a liquid lead target with 1 GeV protons and selected using an on line mass separator. Detailed information regarding sample preparation and data collection followed the protocol described in Iranzo et al.\textsuperscript{46} Fits were carried out with 300 points disregarding the first 10 points due to systematic errors in these, 0.05038 ns/channel, 0.981 ns time resolution. The pH presented in the table is the pH in the sample at 274 K.
**NMR Spectroscopy**

NMR experiments were carried out at 14.1 T at controlled temperature (± 0.1 K) on a Bruker Avance 600 MHz equipped with a Silicon Graphics workstation. Suppression of residual water signal was achieved by excitation sculpting, using a selective square pulse on water 2 ms long. Proton resonance assignment was obtained by TOCSY and NOESY experiments. HSQC experiments were carried out with standard pulse sequences. Spectral processing was performed on a Silicon Graphics O2 workstation using the XWINNMR 3.6 or TOPSPIN 3.1 software. Solutions of apo and metal (Ag(I) and Hg(II)) bound Bri2-23 0.3mM either at acidic (~3.0) and physiological pH were analyzed. The intensities of NOESY cross-peaks, referenced to cross-peaks related to proton pairs at fixed distances were converted into proton-proton distance constraints; the constraints were used to build a pseudopotential energy for a restrained simulated annealing (SA) calculation in torsional angle space. In particular, we performed the calculation with the program DYANA,\(^{47}\) with 300 random starting structures of the peptide and 10000 steps of SA.

**Molecular Dynamics Simulations**

The MD simulations were performed with the GROMACS 3.3.0 software package\(^ {48}\) using the GROMOS 96 force field\(^ {49}\) and the flexible SPC water model. The initial structure was immersed in a periodic water box of triclinic shape (1.50 nm thickness) and neutralized with 1 CL- counterion. Electrostatic energy was calculated using the particle mesh Ewald method.\(^ {50}\) Cutoff distances for the calculation of the Coulomb and Van der Waals interaction were 0.9 and 1.0 nm, respectively. After energy minimization using a steepest decent method, the system was subject to equilibration from 0 to 298 K and normal pressure for 30 ps. The system was coupled to the external bath by the Berendsen pressure and temperature coupling.\(^ {51}\) No positional restrain was given, only two distance restrain for the Cys5-Cu(I) and
Cys 22-Cu(I) binding (low = 1.8, up1= 2.3, up2 = 2.5). An average structure was refined further using a steepest decent energy minimization. For simulating the Cysteine-copper(I) bond, CYS2 residue were selected instead of CYS residue in Gromos 43a2 force field to mimic the oxidized form, ready for metal ion binding. Moreover Arginine (+), Histidine (+) and Lysine (+) were considered in the protonated form like in acidic pH, the total charge of the peptide with Cu(I) ion was of 1+.

The starting point of the MD simulation analysis were the NMR output structures obtained through the program DYANA after integrating the NMR 2D NOESY spectra to obtain structural constrains. Seven of the best 30 DYANA structures were randomly selected as the starting points for MD simulation. The Cys-Me binding site was firstly minimized with the steepest descend method to obtain the estimated distance between copper and the cysteine sulfur. The distance restraints were then applied as constrains for MD simulation analysis using the correct Force Field for this type of Me-bound cysteines (CYS2 for gromos43a1).

100ns of simulation was performed for each structure to reach the RMSD convergence and maintain it for at least 20ns. The seven analyzed structures do not quickly reach the RMSD stabilization, so during the 50ns of simulation a steepest descend energy minimization was done every 5ns, also because of the undefined positional restrain given to the system and because of the intrinsic unstructured part of the peptide.

Results

ThT fluorescence spectroscopy assay

The thioflavin-T (ThT) fluorimetric assay is widely used to underline the presence of stable β sheet secondary structure for peptides or proteins.\textsuperscript{52,53} It is also known, especially for amyloidogenic peptides, that a steady increase over time of the ThT fluorescent signal can be
correlated to formation of precipitating complexes. In order to compare the β sheet structural propensity of the peptide Bri2-23, the variation of ThT fluorescence of Bri2-23 peptide in presence and in absence of Ag(I) and Hg(II) metal ions was analyzed. Immediately after the metal additions, similar values of ThT fluorescence were measured (Figure 2). On the contrary the same experiments repeated after 24 hours showed an increase of ThT fluorescence of Bri2-23 solutions containing metal ions only. In particular, at t=24 hours, the apo peptide gave almost the same ThT fluorescence as measured at t=0, while the binding to Ag(I) or Hg(II) ions resulted in 30-50% of fluorescence enhancements. This behavior supports the propensity of the metal bound Bri2-23 peptide to form β sheet structure in solution, and confirms that a similar structural reorganization of the peptide is induced by Ag(I) and Hg(II).

![Figure 2](image_url)

Figure 2. Relative fluorescence ThT intensity at 480 nm measured at t=0 and t=24 hrs in 50mM phosphate buffer for free Bri2-23 10 µM and in presence of 0.7 equivalents of Ag(I) and Hg(II), respectively; c_{ThT} = 10 µM.

**Mass spectrometry**
The results of mass spectrometric analyses indicate that at slightly acidic pH conditions apo BRI2-23 (data not shown) and its mercurated complex (m/Z = 943.8 and 1415.2) exist as monomers (Figure 3). No species of higher ligand content with respect to metal have been detected, but this may be because it is difficult to detect polymeric forms of ligand or its complexes. To reduce the system’s aggregation susceptibility in the presence of metal, the mixture used in experiments contained 10% of DMSO. Interestingly, in absence of DMSO (data not shown) no spectra was obtainable for the peptide in the presence of Hg(II), possibly indicating aggregation. The MS data can be modeled with doubly deprotonated BRI2-23 and one Hg(II) ion bound, presumably reflecting coordination by the two thiolates.

![Image of ESI-MS spectra](image_url)

Figure 3. ESI-MS spectra of Hg(II) complex of BRI2-23 peptide at pH 6.5 in carbonate buffer (1mM). c_{BRI2-23} = 1 \times 10^{-4}M; Hg(II)/BRI2-23 ratio 1:1; MeOH/H_2O/DMSO = 9:9:2.

Circular dichroism spectroscopy

At both physiological and acidic pH apo Bri2-23 displays typical random coil CD spectra (Figure 4A), with the main absorption at 198 nm characteristic of flexible and disordered protein. Upon Hg(II) and Ag(I) addition the CD spectra undergo changes in both
the shape and in the wavelength of the main adsorption bands (Figures 4B and 4C). The two metals gave very similar changes, showing a negative absorption at 208-210nm. The difference spectra (Figure 4D) obtained by subtracting the CD spectra of the apo peptide from the metal complexes have the typical shape of those of β sheet proteins, thus supporting the ThT results indicating that Bri2-23 forms β sheet structure in presence of both Hg(II) and Ag(I) ions. It cannot be excluded that the CD spectroscopic changes originate from LMCT bands appearing upon the binding of the metal ions to BRI2-23. However, the change towards a β sheet like spectrum upon metal ion binding is remarkable, and appears qualitatively similar for both metal ions.

Figure 4. CD spectra recorded at pH 2.5 (3mM HCl) and 7.0 (50mM phosphate buffer) for 10µM apo Bri2-23 (A), 10µM Bri2-23 in presence of 1.0 equivalents Ag(I) (B); 10µM Bri2-23 in presence of 1.0 equivalents Hg(II) (C) and difference CD spectra for mercurated peptide and its apo form (D).

Potentiometric studies
Potentiometric titration reflects that in the measured pH range unprotected EASNCFAIRHFENKFAVETLICS peptide (BRI2-23) behaves like H₃L acid (Table I). The nine protonation constants correspond to consecutive proton binding to ε-amino group of Lys, thiolate groups of two Cys, terminal α-amino group, imidazole nitrogen of His, γ-carboxylate of three glutamate residues and terminal α-carboxylate group. The protonation of guanidinyl function of Arg residue cannot be followed.

Table 1. Protonation constants for 1mM BRI2-23 peptide and cysteine at T = 298.2 K and I = 0.1 M in NaCl; c_{BRI2-23} = 1x10⁻³, c_{Cys} = 1x10⁻³.

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Table 2. Stability constants for the Hg(II) complexes of cysteine, BRI2-23 and their ternary complexes at Hg(II)/BRI2-23/cysteine ratio 1:1:1; T = 298.2 K and I = 0.1 M in NaCl; c_{BRI2-23} = 5x10⁻⁴, c_{Cys} = 5x10⁻⁴, c_{Hg(II)} = 5x10⁻⁴.

<table>
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Since mercury(II) is known for its extreme high affinity to sulphhydryl-containing ligands, the reasonable explanation of the interactions between the BRI2-23 ligand and Hg(II) obligate us to emphasize that the total complexation of the metal ion with this ligand is essentially complete below pH 2. Therefore, complex formation constants could not be determined directly from potentiometric titration and a competitive potentiometric experiments were carried out with cysteine as competitor ligand under our experimental conditions. First, the protonation constants for cysteine were determined from potentiometric titrations followed by determination of stability constants for its mercuric complexes where the formation constant reported by Stricks and Kolthoff for Hg(Cys)$_2$ species was used as the fixed value. Constants determined for cysteine can be assigned to amine (10.21), thiolate (8.24) and carboxylate (2.12) protonations (Table 1). As in previous studies five complex species have been found within measured pH range all engaging two ligands in Hg(II) sequestration (Table 2). Subsequently, potentiometric titration at a 1:1:1 Hg(II)/Cys/BRI2-23 ratio were carried out. Slow stabilization in the competition region resulted in relatively high standard deviations reported for the calculated complex species. The best fit results are presented in Table 2. Comprehensive analysis of potentiometric data reveals the presence of 13 complex species, five (HgH$_3$Cys$_2$, HgH$_2$Cys$_2$, HgHCys$_2$, HgCys$_2$, HgH$_4$(Cys)$_2$) representing binary Hg(II)
species of cysteine, five binary complexes of mercury(II) bound BRI2-23 (HgH₆L, HgH₅Bri, HgH₄Bri, HgH₃Bri, HgH₂Bri) and three ternary complex species (HgH₂CysL, HgHCysL, HgCysL) that coexist in alkaline pH range (Figure 5).

![Species distribution diagram for Hg(II) complexes of BRI2-23 peptide in the presence of Cys at a 1:1:1 Hg/BRI2-23/Cys ratio; T = 298.2 K and I = 0.1 M in NaCl; c_{BRI2-23} = 5x10^{-4}, c_{Cys} = 5x10^{-4}, c_{Hg(II)} = 5x10^{-4}.]

Although BRI2-23 is a moderately weak competitor of cysteine, we are able to determine stability of its mercuric complexes in ternary system. Usually, the most reliable measure that allows for direct assessment of affinity of various ligands toward metal ion is log K*, protonation corrected stability constant. Taking into account the relatively high standard deviation we are able to quantify the log K* for BRI2-23 Hg(II) complexes to vary between 0.5 and 2. This discrepancy is the consequence of structural rearrangements of peptide chain that mutually result from metal binding and changes in proton concentration. Extensions of peptide sequence with structure-rearranging domain usually increase the stability of its metal complexes. Analogous estimate of log K* for cysteine complexes of general formula HgHₓCys₂ gives value exceeding 23. In consequence BRI2-23 could hardly compete with cysteine for Hg(II) binding. Furthermore, if this measure is critical factor, such a high
difference in stability would abolish formation of ternary complexes that have been determined by potentiometric titrations. Against all odds, mass spectrometric analysis confirms the occurrence of ternary complexes in alkaline pH range (see figures 1S and 2S for details).

The experiments performed in ternary system has been followed by corresponding titrations carried out for binary system with the aim of testing the system’s behavior in absence of any competing ligand. As we know from competition experiments, BRI2-23 is hardly able to compete with cysteine for metal binding in wide pH range. To reproduce the constraints of ternary system and to keep the homogeneity of the complex species formed in solution (vide supra) metal pool accessible for BRI2-23 sequestration was kept at a half of the concentration of ligand. Moreover, formation constant calculated for HgH₂BRI complex was used as the predetermined value. The best fit results are shown in Table 3. Analysis of potentiometric data reveal the formation of seven complex species HgH₆L, HgH₅L, HgH₄L, HgH₃L, HgH₂L, HgL and HgL (Figure 6). Stability constants for corresponding complex species formed in acidic and neutral pH range correlate well with values calculated for ternary system (Table 2), while formation constants for species HgHL and HgL, present in alkaline pH range give reasonable log K* values. In addition, available potentiometric, mass spectrometric and NMR data do not indicate formation of complex species with two ligand molecules binding Hg(II) (Figure 6).

Table 3. Stability constants for the Hg(II) complexes of BRI2-23 at Hg(II)/BRI2-23 ratio 1:2; T = 298.2 K and I = 0.1 M in NaCl; c_{BRI2-23} = 1x10⁻³, c_{Hg(II)} = 5x10⁻⁴.

<table>
<thead>
<tr>
<th>Species</th>
<th>log β</th>
<th>log K</th>
<th>log K*</th>
</tr>
</thead>
<tbody>
<tr>
<td>HgH₆L</td>
<td>55.10(2)</td>
<td>0.73</td>
<td></td>
</tr>
<tr>
<td>HgH₅L</td>
<td>52.09(1)</td>
<td>3.01</td>
<td>1.22</td>
</tr>
<tr>
<td>HgH₄L</td>
<td>48.65(1)</td>
<td>3.44</td>
<td>1.91</td>
</tr>
<tr>
<td>HgH₃L</td>
<td>44.01(1)</td>
<td>4.64</td>
<td>2.03</td>
</tr>
<tr>
<td>HgH₂L</td>
<td>37.72f</td>
<td>6.29</td>
<td>2.10</td>
</tr>
</tbody>
</table>
\[
\begin{array}{cccc}
\text{HgHL} & 29.54(1) & 8.18 & 1.49 \\
\text{HgL} & 18.59(2) & 10.95 & 0.59 \\
\end{array}
\]

\(^a\) L indicates BRI2-23 peptide as a ligand

\(^b\) \(\log K^* = \log [\text{CuH}_{j}L] - \log [\text{H}_{n}L] \) (where the index \(j\) corresponds to the number of the protons in the coordinated ligand to metal ion and \(n\) corresponds to the number of protons coordinated to ligand).

\(^c\) Calculated in ternary system.

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Figure 6. Species distribution diagram for Hg(II) complexes of BRI2-23 peptide in the at a 1:2 Hg/BRI2-23 ratio; \(T = 298.2\) K and \(I = 0.1\) M in NaCl; \(c_{\text{BRI2-23}} = 1 \times 10^{-3}\), \(c_{\text{Hg(II)}} = 5 \times 10^{-4}\).

**Isothermal titration calorimetry**

As we know from potentiometric experiments the total complexation of the metal ion with BRI2-23 is essentially complete below pH 2 and subsequent complex species differ only in the protonation state of groups not involved in Hg(II) binding. Therefore, comprehensive thermodynamic analysis of the interaction of BRI2-23 with Hg(II) by the isothermal titration calorimetry experiments have been performed in acidic pH range (see supplementary information for details).

**Perturbed angular correlation spectroscopy (PAC)**
$^{199}$Hg PAC spectroscopy was conducted in order to elucidate the metal site coordination geometry. The PAC parameters derived from analysis of the experimental data, Figure 4S and 7, are summarized in Table 1S. With a peptide:Hg(II) stoichiometry of 2:1 at pH 3.0 one nuclear quadrupole interaction (NQI) dominates the spectra. This NQI is compares well with previous observations of HgS$_2$ coordination geometries. The same NQI is dominating at peptide:Hg(II) stoichiometry of 1:1 at pH 3.0. At pH 7.4 the signal is still dominated by that of the HgS$_2$ coordination geometry, but it changes slightly, with line broadening especially of the second peak at ~ 2.7 rad/ns, and a minor decrease in frequency. The line broadening implies either internal dynamics at the metal site or that more than one NQI, and thus more than one coordination geometry, is present. Thus, a second NQI was introduced in the data analysis, see Table 1S, and the fitted parameters fall in-between reference data for HgS$_2$ and HgS$_3$ model systems, and might indicate that a fraction of the Hg(II) ions are in coordination geometries with coordination number higher than 2. The analysis is difficult, and other fits of equal quality may exist.

Figure 7: Fourier transformed $^{199}$Hg PAC data for BRI2-23 in 50mM phosphate buffer, pH 3.0 or 7.4 with addition of sucrose (55%) ; $c_{\text{Hg(II)}} = \text{1x10}^{-9}$ M, Hg(II)/BRI2-23 ratio 1:1 or 1:2. Blue: Fourier transformed experimental data; Black: Fit with the parameters presented in
Table 1S. Vertical blue lines are added to aid the eye, at the frequencies recorded at pH 3.0 with 2:1 BRI2-23 to Hg(II) stoichiometry, reflecting a typical HgS$_2$ coordination geometry.

*Nuclear Magnetic Resonance spectroscopy*

In order to characterize the metal coordination sphere of BRI2-23 and to better understand the conformational rearrangements of the peptide, the NMR behavior of both *apo* and metal bound forms was analyzed. Addition of Ag(I) or Hg(II) ions yielded very similar NMR spectra (Figure 5S), suggesting similar coordination features for both metals. However, the spectra of the silver complex were broader than the mercury ones, such that successive analyses were carried out on of Hg(II) systems only.

Hg(II) titration experiments were performed to determine the metal binding stoichiometry, up to 1.8. Hg(II) equivalents were added to BRI2-23 solutions. Hg(II) additions up to 0.9 equivalents resulted in change of NMR parameters of selected proton and carbon NMR resonances, on the contrary higher metal concentrations did not significantly further affect the spectra, except for increased line broadening of NMR signals, probably due to the presence of intermolecular species or soluble aggregates. The addition of 0.5 Hg(II) equivalents causes the reduction of NMR resonances belonging to the *apo* form with the simultaneous appearance of new peaks corresponding to the metal-bound form (Figure 8A), indicating the occurrence of a slow exchange regime (respect to the NMR time scale) between the free and bound states. The presence of both *apo* and metal bound signals observed at that condition strongly indicate that 1:1 complexes, rather than bis-complexes, are formed. Further addition of Hg(II) results in (i) the complete disappearance of the NMR resonances of the *apo* Bri2-23 and (ii) an increase of the intensity of the peaks corresponding to the metal complex (Figure 8B). Furthermore, the correlations observed upon the addition of 0.5 metal equivalents are well superimposed to those corresponding to either the free or metal bound
Bri2-23 (Figure 6S). In addition, the NMR spectra revealed the occurrence of two Hg(II) bound forms in slow exchange respect to the NMR time scale. The two forms (hereafter called form X and Y) exhibited clear and diverse NMR signals for residues 9-21, while showed similar chemical shift for the first 8 residues. From the evaluation of the relative intensities of the NMR signals belonging to the two bound forms, the form X was found to be more abundant than the other. The full $^1$H assignments of Bri2-23 and the two Hg(II) bound forms (X and Y) are reported in Tables 2S, 3S and 4S).

![2D $^1$H-$^1$H TOCSY spectra (NH-Hα region) of Bri2-23 0.5mM in H$_2$O:D$_2$O 90:10 at 298K and pH 3.0 in presence of A) 0.5 equivalents of Hg(II) ions; B) 0.7 equivalents of Hg(II) ions.](image)

After the complete assignment of the NMR spectra of the apo and Hg(II) bound Bri2-23 forms, the Chemical Shift Index (CSI) and the Chemical Shift Variation induced by the metal...
ion for the most abundant species (form X) was calculated. The CSI, calculated on the chemical shift of the Hα protons, is reported in Figure 9. The results obtained for the apo peptide confirm the random coil nature of Bri2-23, as previously detected from CD analysis. On the other hand the CSI obtained for the metal bound form demonstrates a behavior typical of a β strand structured peptide, especially for the regions from Asn-4 to Arg-9 and from Val-17 to Cys-22. Similar results were obtained by the analysis of the chemical shift variation induced by Hg(II) on NH and Hα protons (Figure 10). Both metal bound forms show large chemical shift variations, especially in the regions from 4 to 8 and from 16 to 22. In order to better determine the metal binding donor atoms, the Hg(II) induced chemical shift variations on all of side-chain protons were calculated as well (Figure 10). The protons exhibiting the largest changes belong to Hβ of both Cys-5 and Cys-22, strongly supporting the metal coordination to thiolate groups.

![Figure 9. Chemical Shift Index of Bri2-23 apo form (A) and of the most abundant Hg(II)/Bri2-23 complex X (B).](image-url)
Figure 10. Chemical Shift Variation induced by Hg(II) on Bri2-23 protons.

The analysis of 2D $^1$H-$^1$H NOESY spectra was also performed in order to determine the three dimensional structure the Bri2-23-Hg(II) complex. 70 NOEs were thus converted in proton-proton distance constraints to be used for structure calculation by using the DYANA program. However no a clear cut arrangement of the peptide was obtained from those calculations (bb-RMSD: 4.38 +/- 0.94 Å, and an average target function of 1.08 +/- 0.023).

**Molecular Dynamics Simulations**

The structural propensity of the Bri2-23-Hg(II) complex was investigated by Molecular Dynamics Simulations (MD) performed on seven random structures selected from the thirty ones generated from DYANA calculation (Figure 7S). Combination of experimental and theoretical techniques has been previously applied to investigate metal binding to amyloidogenic model peptides.$^{61-64}$ For each of the seven structures shown, a 100ns of MD
simulation was performed using the simulation program GROMACS\textsuperscript{31}. Contrary to the NMR analysis, we decided to run MD by considering Cu(I) instead of Hg(II) or Ag(I) as the metal bound ion. This choice was mainly due to the fact that Cu(I) is the real metal of interest and that Hg(II) and Ag(I) were just used as probes for Cu(I) in the spectroscopic analysis. Among the seven analyzed structures only five easily reached bb-RMSD convergence after 100 ns (Figure 11, structures A, B, C, D and G). The stability of each simulated systems was controlled also through the output pdb files, where backbone overlap occurred in the last 20 ns confirming the secondary structure stability (Figure 11). For structures E and F, after 100 ns of simulation, bb-RMSD did not converge. This can happen when i) the system attains a random coil secondary structure, so the bb will not be stable at all, or ii) the major part of the bb remain stable over time while the C-and N-terminus (which could be the random part of the sequence) have a high mobility. To monitor this behavior the fluctuation of single residues over time was calculated for structures E and F. As shown in Figure 8S, we found that the N- and C-termini are much more unstable than the central part of the sequence (especially for structure F). For these reasons we decided to include in our analysis also the results obtained from structures F and E.

Ribbon representations are shown in Figure 11. Structures A, B, C, E and G present a well defined conformation, composed by a short N- and C-termini random part, followed by a short \(\beta\)-sheet (parallel or anti-parallel) and a central domain constituted by turn and bend elements. In contrast, structures D and F do not possess an intrinsic \(\beta\)-sheet frame, and they just show the features of a simple but well organized conformation characterized by bend, turn and \(\beta\)-bridge secondary structure.
Figure 11. Backbone RMSD, secondary structure evolutions as function of time. (upper panel) and the snapshots from the last part of the MD trajectories of Hg(II)-BR12-23 molecules (lower panel).

From a statistical point of view, 5 of the 7 analyzed structures show a well defined β sheet arrangement, demonstrating the high propensity of Cu(I) bound peptide to adopt a partial β sheet secondary structure, confirming the analysis of the CD, ThT and NMR experimental data (section 1, 2 and 3). The obtained data support the existence of a β sheet region located at the proximity of the metal binding site (Cys-5 and Cys-22). The amino acids involved in the β sheet formation are not unequivocally determined (see Table 5) suggesting the dynamic
nature of the system. Interestingly, the distance between the two sheets is constant, despite the structured part starts at Ser-3 or at Phe-6.

Table 5. Schematic representation of amino acid residues involved in β sheet formation.

<table>
<thead>
<tr>
<th>Structure</th>
<th>Amino acid residues involved in 1st strand</th>
<th>Amino acid residues involved in 2nd strand</th>
</tr>
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<tbody>
<tr>
<td>A</td>
<td>5 6</td>
<td>18 19</td>
</tr>
<tr>
<td>B</td>
<td>4 5</td>
<td>17 18</td>
</tr>
<tr>
<td>C</td>
<td>4 5 6 7 8 9 10</td>
<td>18 19 20 21 22</td>
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<tr>
<td>E</td>
<td>6 7</td>
<td>19 20 21 22</td>
</tr>
<tr>
<td>F</td>
<td>3 4 5 6</td>
<td>16 17 18</td>
</tr>
<tr>
<td>G</td>
<td></td>
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</tr>
</tbody>
</table>

Discussion

Knowledge of metal binding properties of BRI2 and the products of its proteolytic cleavage is limited to one report related to metallochemistry of ABri, the amyloidogenic product of furin cleaved homologue of BRI2. There are no similar studies for the wild type peptide (BRI2-23) that, contrary to its pathogenic homologue, does not aggregate and even effectively inhibits Aβ aggregation.

Our major interest was to examine the interaction of BRI2-23 with soft metal ions, by using Hg(II) and Ag(I) as Cu(I) probes, to assess the impact of metal sequestration on peptide structure and metal binding affinity at conditions where aggregation is disfavored. Hg(II) is known for very high affinity for thiolate containing ligands. Hg(II) binding to low molecular weight compound like cysteine, penicillamine, glutathione typically yields extremely stable
complex species. On the other hand, the selectivity of metal ion binding to peptides containing two or more cysteinyl residues is strongly dependent on the distance between the thiolates involved in metal ion sequestration,\(^{65}\) over and above their conformational orientation.\(^{57,66}\) Design of peptides that predispose binding thiolates into metal preferred orientation yielded exceptionally stable complex species able to compete with low molecular weight compounds in Hg(II) sequestration.\(^{57}\)

As demonstrated by the CD and NMR analyses apo BRI2 23 preferentially adopts random conformation, which do not favors the intramolecular mercury(II) binding to Cys-5 and Cys-22, separated by 16 amino acid residues. Nonetheless, BRI2-23 has the ability to form Hg(II) mononuclear intramolecular species with a HgS\(_2\) coordination geometry, as indicated by the analysis of ESI MS data (Figure 3). The isotopic profile of the detectable species perfectly matches the simulated mononuclear intramolecular HgBRI2-23 complex. Furthermore, although PAC spectroscopy does not discriminate between the intra- or inter-molecular metal site, it confirms the occurrence of HgS\(_2\) coordination geometry in solution at low pH (Figure 7). Similarly, the large chemical shift changes observed for Cys-5, Cys-22 and residues nearby, indicates that these residues are coordinated to Hg(II) in the predominant metal complex (Figure 10). Interestingly, nearly exclusive intramolecular binding is observed at metal ion concentrations not exceeding 0.5 molar equivalents with respect to the peptide, as reflected by the NMR and potentiometric analysis (Figures 6 and 8). NMR analysis also revealed the presence of two metal-bound species with form (X) still predominant in the equilibrium. Relatively low concentration of minor species (Y) does not allow for its comprehensive structural analysis.

As indicated by the large broadening of NMR signals and by the ITC experiments, further increase of M:L ratios results in the evolution of metal-bridged oligomeric species (e.g.
Hg\textsubscript{m}(BRI2 23\textsubscript{m}) that may coexist in solution with HgBRI2-23 macrochelates. This behavior epitomizes previously detected pH dependent scrambling of ABri.\textsuperscript{67}

Our titration experiments indicate that Hg(II) interact with BRI2-23 in a concentration dependent manner, yielding primarily a mixture of intramolecular macrochelate and structurally undefined species that with metal concentration exceeding 0.7 Hg(II) equivalents are converted into polymeric species of unknown structure and stoichiometry. Aggregates giving response in ThT assays are formed, indicating that the presence of 0.7 Hg(II) or Ag(I) equivalents does lead to aggregation (Figure 2).

The three dimensional structure of mononuclear BRI2-23 complex was unapproachable directly from NMR data analysis. However, NMR provided reliable constraints to molecular dynamics analysis. Nearly all MD trajectories present predominantly \(\beta\) sheet arrangement of peptide structure upon metal binding in contrast to completely random structure of free peptide (Figure 11). As indicated in Table 5, the residues involved in \(\beta\)-sheet structure are not well defined. This behavior is consistent with a relative flexibility of the \(\beta\)-sheet rearrangement and it is in agreement with the lack of precise NOEs constraints unequivocally leading to the NMR structure of the metal complex. However, as supported by the chemical shift analysis (Figure 9) and by the MD data (Figure 11), metal ion binding has critical impact on molecular architecture of the peptide and enforces its refolding to yield species of predominant \(\beta\) pleated sheet conformation, hence more prone to aggregation. Although these species are not exactly the same in terms of proposed molecular architecture of the peptide backbone, most of them resemble either parallel or antiparallel \(\beta\)-sheet rich structures. The suggested conformational shift is additionally supported by changes of far-UV profile of CD spectra demonstrating transition towards \(\beta\)-sheet arrangement of peptide backbone (Figure 4).
Formation of oligomeric species observed at higher metal concentration may either proceed through subsequent Hg(II) and peptide incorporation that yields expanded metal bridged precipitate or alternatively represents the required core for BRI2-23 aggregation. The time dependence and molecular mechanism of aggregation remains to be elucidated. In consequence, metal ion sequestration may eradicate a key attribute of BRI2-23, its Aβ aggregation inhibitory activity and/or even convert the peptide into an aggregation promoter.

Acknowledgements

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Electronic supplementary information (ESI) available:

Mass spectrometry analysis of ternary system; Fig. 1S, ESI-MS spectra of Hg(II) complex of BRI2-23 peptide at pH 11 in ammonium hydroxide (0.4M). $c_{BRI2-23} = 1 \times 10^{-4}$M; Hg(II)/BRI2-23/Cys ratio 1:1:1; MeOH/H$_2$O = 1:2. Isothermal titration calorimetry studies; Fig. 2S, Isotopic profile of ternary Hg(II) complex species of BRI2-23 and cysteine at pH 11 in ammonium hydroxide (0.4M). [BRI2-23] $1 \times 10^{-4}$M; Hg(II)/BRI2-23/Cys ratio 1:1:1; MeOH/H$_2$O = 1:2; Isothermal titration calorimetry; Fig. 3S, Total measured heat associated with titration of BRI peptide with Hg(II), and the binding isotherm derived from the enthalpy
of each injection in the function of molar equivalents of Hg(II); Fig. 4S, 199mHg PAC data for BRI2-23 under the indicated experimental conditions; Fig. 5S. 2D 1H-1H TOCSY spectra (NH-Hα region) of Bri2-23 0.5mM in H2O:D2O 90:10 at 298K and pH 3.0 in presence of A) 0.9 equivalents of Ag(I) ions; B) 0.9 equivalents of Hg(II) ions; Fig. 6S, 2D 1H-1H TOCSY spectra of Bri2-23 0.5mM in H2O:D2O 90:10 at 298K and pH 3.0, in presence of 0.5 equivalents of Hg(II) ions, in presence of 0.9 equivalents of Hg(II) ions; Fig. 7S, Superimposed selected structures obtained by NMR DYANA calculation; Fig. 8S. RMS fluctuation calculated for each residues of structures E and F.

References


