The sea urchin metallothionein system: Comparative evaluation of the SpMTA and SpMTB metal-binding preferences

Mireia Tomas*, Jordi Domènech, Mercè Capdevila, Roger Bofill and Silvia Atrian

Article info

Abstract

Metallothioneins (MTs) constitute a superfamily of ubiquitous metal-binding proteins of low molecular weight and high Cys content. They are involved in metal homeostasis and detoxification, amongst other proposed biological functions. Two MT isoforms (SpMTA and SpMTB) have been reported in the echinoderm Strongylocentrotus purpuratus (sea urchin), both containing 20 Cys residues and presenting extremely similar sequences, although showing distinct tissue- and ontogenic expression patterns. Although exhaustive information is available for the Cd(II)-SpMTA complex, this including the full resolution of its 3D structure, no data has been reported concerning either SpMTA Zn(II) and Cu(I) binding properties, or the characterization of SpMTB at protein level. In this work, both the SpMTA and SpMTB isoforms, as well as their separate α and β domains, have been ruminantly synthesized in the presence of Zn(II), Cd(II) or Cu(II), and the corresponding metal complexes have been analyzed using electrospray mass spectrometry, and CD, ICP-AES and UV-vis spectroscopies. The results clearly show a better performance of isoform A when binding Zn(II) and Cd(II), and of isoform B when coordinating Cu(I). Thus, our results confirm the differential metal binding preference of SpMTA and SpMTB, which, together with the reported induction pattern of the respective genes, highlight how also in Echinodermata the MT polymorphism may be linked to the evolution of different physiological roles.

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1. Introduction

Metallothioneins (MTs) are a superfamily of universal and ubiquitous low molecular weight proteins that bind essential and toxic metal ions through their abundant Cys residues, forming multinuclear metal–thiolate clusters. They are involved in many crucial biological processes, such as metal homeostasis and detoxification, or oxidative stress protection, among others [1]. Although their protein sequences are dramatically heterogeneous and consequently their classification is not a trivial affair, their metal-binding behavior converges to either Zn-thioneins, which show a binding preference for divalent metal ions, or Cu-thioneins, with an optimized coordination of monovalent metal ions [2]. Beyond this simplistic dual model, the comprehensive consideration of the metal-binding features of a huge number of MT peptides later led to the proposal of a continuous gradation between both extreme behaviors [3]. Unfortunately, the sequence determinants of the metal specificity of MTs are still unknown [4], although it should be considered a key element for a better understanding of the evolutive differentiation [5] and physiological function of these peculiar metalloproteins [6]. Polymorphism is one of the most significant features of MTs, since at least all the eukaryote organisms studied until now exhibit MT systems composed of almost similar paralog forms [5]. Ideally, each isoform would correspond to a definite physiological function, but it is plainly evident that in most cases ontogenic, tissue- and/or functional differentiation has not been achieved, so that no clear correlation exists between multiple MT isoforms and their in vivo performance.

In this scenario, and with the aim of gathering data on the most prominent model organisms, we undertook the characterization of the metal-binding preferences of the purple sea urchin (Strongylocentrotus purpuratus, echinoderm) MT isoforms. Echinoderms constitute a very interesting phylum since, as the unique invertebrate deuterostomes, they are evolutive close to the Chordates, which are also deuterostomes (for a recent review on Echinodermata MTs, see [7]). At genomic level, seven different homologous MT genes have been reported in S. purpuratus, of which only three produce biologically relevant isofoms: SpMTA and the pair SpMTB1–SpMTB2, the latter encoding proteins with identical sequence [8–10]. Significantly,
SpMTA and SpMTB exhibit different quantitative tissular and temporal expression patterns. Hence, SpMTA is transcribed in ectodermal tissues, while SpMTB is expressed in both the ectodermal and endodermal structures. Both genes are constitutively expressed, and under physiological conditions SpMTA mRNA levels reach up to 10 times those of SpMTB. However, under metal (zinc or cadmium) overload, SpMTB expression is further induced, so that SpMTB achieves protein levels similar to those of SpMTA [8,10].

SpMTA is a 64-amino acid peptide including 20 Cys, while the SpMTB sequence is two amino acids longer, with fully conserved Cys at the N-terminal moiety, and attracted great interest, since the Cys–Cys motifs typical of the C-terminal (β domain) of the mammalian MTs (Fig. 1) [15], cDNAs encoding for its separate moieties were obtained by PCR reactions that added restriction sites to the respective 5′-ends of the coding sequences, using the purified cDNA of the EST clone as template and the following primers: for the N-terminal fragment or α domain (comprising residues 1–36 of the full polypeptide), 5′-CCGGATCCTAGCATGCA-3′ (upstream) and 5′-GCGCCACCTAGCATGCA-3′ as downstream primer; and for the C-terminal fragment or β domain (residues 37–64 of the full polypeptide), 5′-ACACAGGGATCCTAGCATGCAAACAGCTGGTCGC-3′ (upstream) and 5′-AAACCTGGTCGCTAGCATGCAACAGCTGGTCGC-3′ (downstream). The cDNA of the SpMTB isoform was obtained from the Sea Urchin Genome Project Library Collection, in the California Institute of Technology (Caltech), and corresponded to the EST clone PMC6237956 (seq ID NCBI 214576). The correspondant SpMTA [12–14], which culminated with the determination of the number and position of the binding domains in different MTs [11].

2. Materials and methods

2.1. Cloning of MT cDNA constructs

All the metal-MT complexes investigated in the present study were recombinantly synthesized in Escherichia coli through cloning of their coding sequences into the pGEX-4T1 plasmid (GE Healthcare, Little Chalfont, UK) to yield primary glutathione S-transferase (GST)-MT fusions from which the corresponding metal-MT complexes were subsequently purified [16,17]. The SpMTA cDNA was kindly provided by Prof. Kagi and subsequently cloned into the pGEX-4T1 BamH1/SalI restriction sites added by PCR amplification, using 5′-CCGGATCCTAGCATGCA-3′ (upstream primer and 5′-GCGCCACCTAGCATGCA-3′ as downstream primer). According to the reported SpMTA domain boundaries [15], cDNAs encoding for its separate moieties were obtained by PCR reactions that added BamH1/Xhol restriction sites with the following oligonucleotides: for the N-terminal fragment or α domain (comprising residues 1–36 of the full polypeptide), 5′-CCGGATCCTAGCATGCAAACAGCTGGTCGC-3′ (upstream) and 5′-AAACCTGGTCGCTAGCATGCAACAGCTGGTCGC-3′ (downstream); and for the C-terminal fragment or β domain (residues 37–64 of the full polypeptide), 5′-ACACAGGGATCCTAGCATGCAAACAGCTGGTCGC-3’ (upstream) and 5′-AAACCTGGTCGCTAGCATGCAACAGCTGGTCGC-3′ (downstream). The cDNA of the SpMTB isoform was obtained from the Sea Urchin Genome Project Library Collection, in the California Institute of Technology (Caltech), and corresponded to the EST clone PMC6237956 (seq ID NCBI 214576). The corresponding ORF was amplified by a PCR reaction that added BamH1/Xhol restriction sites to the respective 5′ and 3′ ends of the coding sequence, using the purified cDNA of the EST clone as template and the following primers: 5′-CCGGATCCTAGCATGCAAACAGCTGGTCGC-3′ (upstream) and 5′-AAACCTGGTCGCTAGCATGCAACAGCTGGTCGC-3′ (downstream). Based on the reported domain boundaries of SpMTA [15], cDNAs encoding for the separate SpMTB

![Fig. 1](image-url)
moieties were obtained by PCR reactions with the following oligonucleotides: for the N-terminal fragment or α domain (encompassing residues 1–37 of the full polypeptide), 5′-CCCCGATCCATGGTGCTCAAGTGTGTCG-3′ (upstream) and 5′-AAACTTCAGTCATCCAGCCTGGAGTTCCACC-3′ (downstream); and for the C-terminal fragment or β domain (residues 38–66 of the full polypeptide), 5′-ACACACGGCATAATGCCTAAATGGCGGCATGCGC-3′ (upstream) and 5′-AACCCCTTCAGCTGACATGGACAGTCTCCCCCTC-3′ (downstream). All the PCR reactions consisted of 35-cycle amplifications, performed with 1.25 U of GoTaq DNA polymerase (Promega, Madison, USA), 0.25 mM dNTPs and 0.24 μM of the required primers at 2 mM MgCl2 (final concentration), in a final volume of 100 μL, under the following cycle conditions: 30 s at 94 °C (denaturation), 30 s at 58 °C (hybridization) and 30 s at 72 °C (elongation). An initial denaturation step where samples were heated at 94 °C for 2 min ensured that the target DNA was completely denatured, and elongation conditions were maintained for 7 min after the 35 cycles. The final products were analyzed by agarose gel electrophoresis/GeRed Nucleic Acid Gel Stain (Biotium, Hayward, USA) staining; the band with the expected size was excised and subcloned into the pGEX-4T1 vector.

2.2. Recombinant synthesis and purification of metal-MT complexes

The SpMTA-GST and SpMTB-GST fusion polypeptides and their respective α and β domains were biosynthesized in S. L cultures of transformed protease-deficient E. coli BL21 cells. Expression was induced with isopropyl β-D-thiogalactopyranoside (IPTG) and cultures were supplemented with 500 μM CuSO4, 300 μM ZnCl2 or 300 μM CdCl2 (final concentrations) and were allowed to grow for a further 3 h. In the case of Cd(II)-enriched cultures, both normal and low aeration conditions (N.A. and L.A., respectively) were assayed according to the procedure described elsewhere [18]. A total protein extract was prepared from these cells as previously described [16]. Metal complexes were recovered from the MT-GST fusion constructs by thrombin cleavage and batch-affinity chromatography using Glutathione-Sepharose 4B (General Electric HC). The metal complexes were finally purified through FPLC in a Superdex75 column (General Electric HC) equilibrated with 50 mM Tris–HCl, pH 7.0. Selected fractions were confirmed by 15% SDS–PAGE and kept at –80 °C until further use. All procedures were performed using Ar (pure grade 5.6) saturated buffers, and all syntheses were performed at least twice to ensure reproducibility. Further details on the purification procedure can be found in [16]. As a consequence of the cloning requirements, the dipeptide Gly–Ser was present at the N-terminus of all polypeptides; however, this had previously been shown not to alter the MT metal-binding capacities [17].

2.3. In vitro Cd- and Cu-binding studies

The titration of all Zn-MT complexes with Cd(II) or Cu(I) at pH 7 were carried out following the methodology previously described [19,20], using CdCl2 or [Cu(CH3CN)4]ClO4 solutions, respectively. The acidification/reneutralization experiments were also performed by adapting the procedure reported in [21]. Essentially, 10–20 μM preparations of the Cd-peptides were acidified from pH 7.0 to pH 1.0 to 2.0 with 1 × 10−3 M HCl. CD and UV–vis spectra were recorded at several pH values both immediately after acid addition and 10 min later, always with identical results. Finally, the samples were kept at pH 1.0–2.0 for 20 min and were then reneutralized with 1 × 10−3 M NaOH, and CD and UV–vis spectra were recorded at several pH values. In some cases, several molar equivalents of an aqueous solution of Na2S were added to the reneutralized MT forms, with the aim of reproducing the original Cd-MT CD fingerprints. All results were corrected for dilution effects, and during all experiments strict oxygen-free conditions were kept by saturating all solutions with Ar.

2.4. Protein quantification and spectroscopic analyses

The S, Zn, Cd and Cu content of the Zn-, Cd- and Cu-MT preparations was analyzed by means of Inductively Coupled Plasma Atomic Emission Spectroscopy (ICP-AES) in a Polyscan 61E (Thermo Jarrell Ash) spectrometer, measuring S at 182.040 nm, Zn at 213.856 nm, Cd at 228.802 and Cu at 324.803 nm. Samples were treated as in [22], but were alternatively incubated in 1 M HCl at 65 °C for 5 min prior to measurements, in order to eliminate possible traces of labile sulfides, as otherwise described [23]. Protein concentrations were calculated from the acid ICP-AES sulfur measurement, assuming that all S atoms were contributed by the MT peptide. A Jasco spectropolarimeter (Model J-715) interfaced to a computer (J700 software) was used...
Table 1

| MT     | Theoretical MW | Experimental MW |
|--------|----------------|-----------------|
| SpMTA  | 6532.0         | 6534.9 ± 0.8    |
| αSpMTA | 3874.0         | 3873.9 ± 0.4    |
| βSpMTA | 2820.2         | 2819.6 ± 0.5    |
| SpMTB  | 6700.7         | 6698.6 ± 0.4    |
| αSpMTB | 3913.5         | 3912.4 ± 0.1    |
| βSpMTB | 2949.3         | 2948.6 ± 0.4    |

for CD measurements at a constant temperature of 25 °C maintained by a Peltier PTC-351S apparatus. Electronic absorption measurements were performed on an HP-8453 Diode array UV–vis spectrophotometer. All spectra were recorded with 1 cm capped quartz cuvettes, corrected for the dilution effects and processed using the GRAMS 32 Software.

2.5. Mass spectrometry

MW determinations were performed by electrospray ionization time-of-flight mass spectrometry (ESI-TOF MS) on a Micro Tof-Q instrument (Bruker) interfaced with a Series 1100 HPLC Agilent pump, equipped with an autosampler, all of which were controlled by the Compass Software. Calibration was attained with 0.2 g NaI dissolved in 100 mL of a 1:1 H2O:isopropanol mixture. Samples containing MT complexes with divalent metal ions were analyzed under the following conditions: 20 µL of protein solution injected through a PEEK (polyether heteroketone) column (1.5 m × 0.18 mm i.d.), at 40 µL min⁻¹; capillary counter-electrode voltage 5 kV; desolvation temperature 90–110 °C; dry gas 6 L min⁻¹; spectra collection range 800–2000 m/z. The carrier buffer was a 5:95 mixture of acetonitrile:ammonium acetate/ammonia (15 mM, pH 7.0). Alternatively, the Cu-MT samples were analyzed as follows: 20 µL of protein solution injected at 30 µL min⁻¹; capillary counter-electrode voltage 3.5 kV; lens counter-electrode voltage 4 kV; dry temperature 80 °C; dry gas 6 L min⁻¹. Here, the carrier was a 10:90 mixture of acetonitrile:ammonium acetate/ammonia (15 mM, pH 7.0). For analysis of all recombinant MT molecular masses, 20 µL of the corresponding Zn–MT samples were injected under the same conditions described before, but using a 5:95 mixture of acetonitrile:formic acid pH 2.5 as liquid carrier, which caused the complete demetalation of the peptides. The same conditions were then used to remove Zn(II) ions from mixed-metal Zn, Cu-MT species in order to quantify their total Cu content.

3. Results and discussion

3.1. Integrity and identity of the recombinant polypeptides

The cDNAs coding for the SpMTA and SpMTB isoforms as well as for their separate domains were cloned into the pGEX-4T1 plasmid for peptide synthesis. DNA sequencing confirmed that all these constructs included no artifactual nucleotide substitutions, and that the respective coding sequences were cloned in the correct frame after the GST encoding fragment. Recombinant syntheses yielded MT peptides of which the identity, purity and integrity was confirmed by ESI-MS of the respective apoforms, obtained by acidification at pH 2.5 of the corresponding Zn-MT complexes. Hence, in each synthesis a unique peak was detected, in which the MW was consistent with the calculated MW of the respective recombinant MT peptide, including N-terminal Gly–Ser residues derived from the GST-fusion construct (Table 1).

3.2. SpMTA exhibits better Zn(II) and Cd(II) binding abilities than SpMTB

Both SpMTA and SpMTB yield a major Zn7-MT species when recombinantly synthesized as Zn(II)-complexes (Table 2, Fig. 2(A) and (D)), although the occurrence of several minor undermetalated, partially oxidized species is markedly significant for SpMTB, where all the species ranging from Zn6 to Zn3 are produced (Fig. 2(D)). This result already suggests a lower ZnII-binding ability for SpMTB compared to SpMTA. The CD fingerprints of both Zn-preparations are similar and characteristic of Zn-MT complexes [23], and which are also very similar to the CD spectrum previously reported for recombinant Cd7–SpMTA [12,13].
very minor Cd$_{2,1}$ species was identified in the Cd-SpMTA prepara-
tion (Table 2, Fig. 4(A)), its relative abundance is practically negli-
gible if compared with the minor forms detected in Cd-SpMTB (Fig. 4(D)). In this case, an unexpected major Cd$_8$ species was produced,
which Cd$_2$- and Cd$_7$S$_2$-SpMTB are worth noting (Table 2, Fig. 4(D)).
The CD fingerprint of this preparation differs from that of Cd-SpMTB [14].
Moreover, this spectrum can be also obtained by Cd/SpMTB replacement
after the addition of 4 Cd(II) equivalents to Zn$_2$-SpMTA at pH 7 (Fig. 6). In contrast to these results, the β-SpMTA and β-SpMTB domains yield patently different results when synthesized in Zn(II)-
and Cd(II)-enriched media. Hence, both isoforms are recovered as major Zn$_3$ complexes, although the partially undermetalated species detectable for SpMTA do not appear for SpMTB (Table 2, Fig. 2(C) and (F)). Also, as determined from the difference between conventional and acid ICP-AES measurements (data not shown), the presence of inorganic sulfide in the recombinant Zn-β-SpMTB preparations may account for the red-shift observed for Zn-β-SpMTB CD spectrum in comparison to Zn-β-SpMTA (Fig. 3(C)). Even more pronounced differences arise for the Cd(II)-binding capacity of the separate β domains, as β-SpMTA yields a major, canonical Cd$_3$ species accompanied by minor Zn-containing complexes, while β-SpMTB predominantly folds into Cd$_3$S$_2$-β-SpMTB, thus with complete absence of Zn (Fig. 4(C) and (F), respectively, and Table 2). Moreover, the Cd$_3$S$_2$-β-SpMTB sulfide containing complex exhibits an acute red-shift in its CD spectrum when compared to Cd$_3$-β-SpMTA (Fig. 5(C)).

At this point, it becomes evident that the Cd-binding abilities of SpMTB are poorer than those of SpMTA [3], and that this feature is most likely due to a patent lower Cd(II)-binding ability of β-SpMTB if compared to β-SpMTA, as corroborated by the fact that the respective α domains exhibit no major differences. In order to better understand the peculiarities of the Cd-SpMTA and Cd-SpMTB complexes, and to support the higher Cd(II)-binding preference of β-SpMTA compared to β-SpMTB, additional experimental results were considered. In the first place, it is worth noting that the CD spectra of the Zn/Cd replacement processes in both β domains evolved following a distinct pattern,
Fig. 4. Representative charge states for the ESI-MS spectra recorded at pH 7.0 of recombinant Cd-SpMTA (A), Cd-αSpMTA (B), Cd-βSpMTA (C), Cd-SpMTB (D), Cd-αSpMTB (E) and Cd-βSpMTB (F). The observed species are collected in Table 2.

Table 3
Analytical characterization of recombinant SpMTA and its independent constitutive fragments synthesized in Cu-enriched media under normal (N.A.) and low (L.A.) aeration conditions.

| MT          | Concentration and metal/MT ratioa | pH | Species (% abundance) | ESI-MSb | Theoretical MW | Experimental MW |
|-------------|----------------------------------|----|-----------------------|---------|----------------|-----------------|
| SpMTA (N.A.) | 0.5 × 10⁻⁴ M                     | pH 7.0 | M₄-SpMTA (100) | 6782.7–6786.0 | 6771.0 ± 0.1 |
|             | 5.5 Cu                           |     | M₄-SpMTA (80)       | 7032.8–7039.5 | 7029.7 ± 1.2 |
|             | 3.0 Zn                           |     | M₃-SpMTA (40)       | 6845.2–6849.4 | 6838.0 ± 1.1 |
| SpMTA (L.A.) | 0.5 × 10⁻⁴ M                     | pH 7.0 | M₄-SpMTA (100) | 7032.8 | 7033.3 ± 2.2 |
|             | 8.7 Cu                           |     | Cu₄-SpMTA (20)      | 7032.8 | 7030.6 ± 0.8 |
|             | 1.7 Zn                           |     | Cu₄-SpMTA (20)      | 7032.8 | 7034.9 ± 0.1 |
| αSpMTA (N.A.) | 0.3 × 10⁻⁴ M                    | pH 7.0 | Oxidized dimeric species | – | – |
|              | 6.0 Cu                           | pH 2.5 | N.D.                 | – | – |
|              | 0.1 Zn                           |     | N.D.                 | – | – |
| αSpMTA (L.A.) | 0.2 × 10⁻⁴ M                    | pH 7.0 | N.D.                 | – | – |
|              | 4.2 Cu                           | pH 2.5 | Oxidized dimeric species | – | – |
| βSpMTA (N.A.) | 0.7 × 10⁻⁴ M                    | pH 7.0 | M₁₂-βSpMTA (100)    | 3132.9–3137.1 | 3129.6 ± 0.1 |
|              | 5.1 Cu                           |     | M₁₁-βSpMTA (30)     | 3195.5–3200.5 | 3195.6 ± 0.3 |
|              | 0.2 Zn                           |     | M₁₀-βSpMTA (20)     | 3070.4–3073.7 | 3067.6 ± 2.1 |
| βSpMTA (L.A.) | 0.05 × 10⁻⁴ M                   | pH 7.0 | M₁₂-βSpMTA (100)    | 3132.9–3137.1 | 3131.6 ± 1.6 |
|              | 3.4 Cu                           |     | M₁₀-βSpMTA (40)     | 3070.4–3073.7 | 3066.8 ± 1.9 |
|              | 0.7 Zn                           | pH 2.5 | N.D.                 | – | – |

a MT concentration and metal/MT ratio calculated from acid ICP-AES results.
b Experimental and theoretical molecular weights corresponding to the Cu-peptides. Species shown in bold correspond to the major components of the preparations. In the case of Zn, Cu mixed-metal species, the indicated theoretical molecular weights correspond to the homometallic Cuₓ and Znₓ species (MW_{CuₙMT} – MW_{ZnₙMT}) and the metal-to-protein stoichiometries deduced at pH 7.0 are indicated as Mₓ (M = Zn + Cu). Cu contents at pH 2.5 were calculated from the mass difference between holo- and apoproteins. N.D.: not detected.
especially after the 3rd Cd(II) equivalent added (Fig. 7(A)–(D)). But most interestingly, while the CD fingerprint of the recombinant Cd-

βMT isoform, the initial CD fingerprint can only

reneutralization process followed

conversion of the derivative-shaped band into a Gaussian band (Fig. 7(G)). Taken together, these

results reveal that the decrease of the divalent metal ion-preference (Zn(II) or Cd(II)) of an MT peptide entails the

tire SpMTB protein, which behaves as a less proficient Cd(II)-binding

species (MWCu-MT–MWZn-MT), and the metal-to-protein stoichiometries deduced at pH 7.0 are indicated as M_i (M = Zn + Cu). Cu contents at pH 2.5 were calculated from the mass difference between holo- and apoproteins. N.D.: not detected.

Table 4

Analytical characterization of recombinant SpMTB and its independent constitutive fragments synthesized in Cu-enriched media under normal (N.A.) and low (L.A.) aeration conditions.

| MT        | Concentration and metal/MT ratio | pH 7.0 | Species (% abundance)  | ESI-MS<sup>3</sup> | Theoretical MW | Experimental MW |
|-----------|----------------------------------|--------|------------------------|--------------------|---------------|-----------------|
| SpMTB (N.A.) | 0.9 × 10⁻⁴ M 15.4 Cu 0.2 Zn | pH 7.0 | M₃-βSpMTB (100) 4163.7–4161.7 55.6 3443.9 | 4163.7 4159.2 1.1 | 3443.3 3435.2 0.7 | 4163.7 4158.6 0.6 |
|           | 5.5 Cu 1.9 Zn                  |        | M₄-βSpMTB (40) 3262.0–3266.3 31.0 3449.7 | 3262.2 3259.7 1.0 | 3449.3 3440.9 1.2 | 3262.2 3260.2 0.4 |
|           | 0.3 × 10⁻⁴ M 10.0 Cu 0.2 Zn    |        | M₅-βSpMTB (100) 3387.1–3393.0 32.0 3449.7 | 3387.1 3385.5 0.9 | 3449.7 3448.5 1.0 | 3387.1 3385.5 0.1 |
| aSpMTB (N.A.) | 0.6 × 10⁻⁴ M 5.5 Cu 0.5 Zn | pH 7.0 | M₄-aSpMTB (100) 4163.7–4161.7 55.6 3443.9 | 4163.7 4159.2 1.1 | 3443.3 3435.2 0.7 | 4163.7 4158.6 0.6 |
|           |                                |        | M₄-aSpMTB (90) 3913.5 | 3903.3 3.1 | 3913.5 | 3903.3 3.1 |
|           |                                |        | M₄-aSpMTB (60) 3324.6–3329.6 33.2 3449.7 | 3324.6 3326.2 0.8 | 3449.7 3448.5 1.0 | 3324.6 3326.2 0.8 |
|           |                                |        | M₄-aSpMTB (80) 3387.1–3393.0 32.0 3449.7 | 3387.1 3385.5 0.9 | 3449.7 3448.5 1.0 | 3387.1 3385.5 0.1 |
| aSpMTB (L.A.) | 0.3 × 10⁻⁴ M 7.0 Cu 0.0 Zn | pH 2.5 | M₄-aSpMTB (100) 4163.7 | 4157.6 0.5 | 4157.6 0.5 |
|           |                                |        | M₄-aSpMTB (50) 3426.2 | 3421.9 0.9 | 3421.9 0.9 |
|           |                                |        | M₄-aSpMTB (90) 3913.5 | 3903.3 3.1 | 3913.5 | 3903.3 3.1 |
| βSpMTB (N.A.) | 0.3 × 10⁻⁴ M 3.3 Cu 0.2 Zn | pH 7.0 | M₃-βSpMTB (100) 3262.0–3266.3 31.0 3449.7 | 3262.2 3259.7 1.0 | 3449.3 3440.9 1.2 | 3262.2 3260.2 0.4 |
|           |                                |        | M₃-βSpMTB (80) 3387.1–3393.0 32.0 3449.7 | 3387.1 3385.5 0.9 | 3449.7 3448.5 1.0 | 3387.1 3385.5 0.1 |
|           |                                |        | M₃-βSpMTB (60) 3199.5 | 3195.3 0.6 | 3195.3 0.6 |
|           |                                |        | M₃-βSpMTB (80) 3262.0 | 3257.9 1.0 | 3257.9 1.0 |
|           |                                |        | M₃-βSpMTB (100) 3387.1 | 3385.5 0.1 | 3385.5 0.1 |
| βSpMTB (L.A.) | N.D.                          | pH 7.0 and 2.5 | N.D. | N.D. | N.D. | N.D. |

<sup>*</sup> MT concentration and metal/MT ratio calculated from acid ICP-AES results.

<sup>1</sup> Experimental and theoretical molecular weights corresponding to the Cu-peptides. Species shown in bold correspond to the major components of the preparations. In the case of Zn,Cu mixed-metal species, the indicated theoretical molecular weights correspond to the homometallic Cu species (MW_{Cu-MT–MW_{Zn-MT}}), and the metal-to-protein stoichiometries deduced at pH 7.0 are indicated as M_i (M = Zn + Cu). Cu contents at pH 2.5 were calculated from the mass difference between holo- and apoproteins. N.D.: not detected.

3.3. Cu-binding abilities: SpMTB exhibits a stronger Cu-thionein character than SpMTA

Two different types of preparations were obtained for SpMTA when synthesized in Cu-supplemented E. coli cells, depending on the aeration conditions of the corresponding bacterial cultures (cf. Table 3). Although in both cases SpMTA was recovered as a mixture of Zn,Cu heteronuclear species, the average Zn content of the preparation was always higher in the normally aerated culture, with concomitant average lower Cu content (5.5 Cu:3.0 Zn), and with the opposite results for the low aeration conditions (8.7 Cu:1.7 Zn). In concordance with these ICP-AES results, in the former case the most abundant detected species by ESI-MS was M₄-βSpMTA, followed by a collection of higher nuclearity complexes (M₅–M₁₀, Fig. 8(A)), which should be mainly consisting of Cu₈Zn₉ complexes (1 < x < 5), since ESI-MS data at pH 2.5 identified a highly predominant Cu₈ cluster (Fig. 8(B)). Conversely, when SpMTA was synthesized under low aeration, the most abundant species in solution switched to M₈ (M = Zn + Cu), readily followed by M₉–M₁₁ complexes (Fig. 8(C)), whose acid ESI-MS analyses revealed to be built by Cu₈ and Cu₉ cores (Fig. 8(D)) plus some added Zn(II) ions. The homologous SpMTB was also synthesized under both culture conditions. Under normal aeration, this isoform yielded preparations similar to those of SpMTA when produced under low aeration, i.e. a predominant M₈ complex accompanied by M₉ and M₁₀ species that acid ESI-MS identified as predominant Cu₈Zn₇ complexes (Table 4, Fig. 8(E) and (F)). With respect to the recombinant
4. Conclusions

From the set of data reported in this work, a differential metal ion binding preference is envisaged for SpMTA and SpMTB, since different results attribute a more pronounced Cu-thionein character to SpMTB than to SpMTA, and vice versa for the Zn(II)- and Cd(II)-thionein character. In summary, and according to the criteria proposed to evaluate the preference of a given MT peptide either for monovalent or divalent metal ion coordination [3], it is worth noting that (i) SpMTA is able to fold into almost unique and sulfide-free Zn(II)/Cd(II)-complexes, while for these metal ions SpMTB yields preparations containing a large variety of species, and with a significant sulfide content; and (ii) SpMTB is able to yield, at standard bacterial cell copper concentrations (i.e. normal aeration of the E. coli cultures), Cu-complexes that SpMTA is only able to produce in higher-copper environments. Dissection of the metal-binding abilities of both isoforms into their domains suggests that while the β domain would be the only one responsible for the diminished divalent metal binding capacity of SpMTB, both domains would contribute to the enhanced capacity of SpMTB for Cu(I) binding.

The differential Zn/Cu binding performance of SpMTA and SpMTB suggests that both isoforms exert at least partially distinct complementary functions in the physiology of sea urchins. For divalent metal ions, SpMTA may be mostly responsible for housekeeping Zn homeostasis, while SpMTB would only come into play under Zn or Cd overload emergency situations. This hypothesis is supported by the fact that observed for divalent metal ions. Hence, αSpMTA yielded, under all assay conditions, mixtures of Zn, Cu heterometallic complexes with highly oxidized and/or poorly metalated species, together with dimeric complexes (Table 3), which exhibit almost flat CD fingerprints (Fig. 9(C) and (D)). Contrarily, αSpMTB invariably yielded well defined results: at regular oxygenation Cu5-, Cu4Zn1-, Cu4 and Cu3-αSpMTB were the predominant species (Fig. 10(A) and (D), Table 4), and at low aeration conditions a mixture of homometallic Cu complexes with major Cu4 and Cu5 stoichiometries was detected (Fig. 10(F), Table 4). These results were consistent with well-defined CD spectra, typical of Cu-MT chromophores, despite the fact that they exhibited rather low intensity (Fig. 9(C) and (D)). Contrarily to the α domains, no significant differences could be detected for the recombinantly synthesized β peptides in copper-supplemented media. At normal aeration both isoforms yield a mixture of species with close average Zn/MT (0.2) and Cu/M (3.3–5.1) ratios (Tables 3 and 4) and very similar CD spectra (Fig. 9(E)), where M is the major species present (Fig. 10(B) and (C)). ESI-MS at pH 2.5 (Fig. 10(E)) confirmed the almost quasi homometallic nature of the Cu-βSpMTB preparations, and comparison with data at pH 7.0 allowed to deduce that the M5 major species corresponds to a mixture of Cu5 and Cu4Zn1 complexes. At low aeration, very diluted samples were obtained and consequently very weak CD spectra were recorded (Fig. 9(F)). However, βSpMTA ESI-MS analysis at pH 7.0 (Fig. 10(G)) shows a heterometallic species distribution similar to that observed at normal oxygenation, with a major M3 form (Table 3). In view of the results obtained in the recombinant syntheses of the βSpMT fragments, and in order to have a more precise picture of their differential behavior when coordinating Cu(I), Zn/Cu titrations were performed and followed through their CD spectra evolution. In the case of βSpMTA, the maximum degree of folding was achieved for 3–4 Cu(I) equivalents added to the Zn(II)-βSpMTA preparation, with little further variation (Fig. 11(A)–(C)). Contrasting, βSpMTB reaches its maximum folding after 6 Cu(I) equivalents added (Fig. 11(D)–(F)), which is in concordance with a greater Cu(I) binding capacity. Finally, when comparing these results with those of the recombinant complexes, it becomes obvious that the CD spectrum of SpMTA is fully reproduced after 4 Cu(I) equivalents added (Fig. 11(G)), whereas for βSpMTB, the highest CD spectra resemblance is attained after 7 Cu(I) equivalents added (Fig. 11(H)).
that SpMTA is constitutively expressed at a higher rate than SpMTB at low Zn concentrations, while increased divalent metal ion concentrations induce SpMTB expression up to similar values than SpMTA [8], analogously to what has been described for the nematode CeMT1 and CeMT2 isoforms [27]. Interestingly, the existence of multiple MT isoforms involved in different physiological roles has been very recently confirmed in another sea urchin species, Paracentrotus lividus, in which its five different MT isogenes respond distinctively to Cd transcriptional activation, thus showing a certain degree of evolutionary functional differentiation [28]. Unfortunately, no information is available for the copper physiology in sea urchin, or for copper requirements during its development stages, or for the response of both SpMT isogenes to copper overload. It would not be far-fetched to assume that SpMTB, given its higher Cu-thionein character, could better perform some function related to Cu metabolism in this group of organisms. Overall, and as for other taxa in which MT peptides have
Fig. 8. Representative charge states for the ESI-MS spectra recorded at pH 7.0 and pH 2.5 of recombinant SpMTA (A, B) and SpMTB (E, F) synthesized in Cu-supplemented E. coli media at normal aeration (N.A.) conditions, and of recombinant Cu-SpMTA (C, D) obtained at low aeration (L.A.) conditions. The observed species are collected in Tables 3 and 4.

Fig. 9. CD spectra corresponding to the entire MT (A, B) and the separate constitutive α (C, D) and β (E, F) domains of SpMTA (black) and SpMTB (red) recombinantly synthesized in Cu-supplemented media at normal aeration (N.A.) and low aeration (L.A.) conditions (left and right, respectively). (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)
Fig. 10. Representative charge states for the ESI-MS spectra of recombinant αSpMTB (A, D, F), βSpMTA (B, G) and βSpMTB (C, E) synthesized in Cu-supplemented E. coli media at normal (N.A.) and low (L.A.) aeration conditions. M-MT species where M = Zn + Cu are assigned in spectra recorded at pH 7.0 (A–C, G), while Cu-MT species are assigned when recorded at pH 2.5 (D–F). The observed species are collected in Tables 3 and 4.

Fig. 11. CD spectra recorded during the titration of Zn-βSpMTA (A–C) and Zn-βSpMTB (D–F) with Cu(I) at pH 7. Also, a comparison with the CD spectrum of the corresponding recombinant Cu-βMT samples obtained at normal aeration conditions (in red) is shown (G, H). (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)
been previously characterized, more or less pronounced differential metal ion binding properties would underlie the observed polymorphism, even when the involved polypeptides share a 85% sequence identity, as is the case for S. purpuratus MTs.

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