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- 1 Exploring the structural and functional role of a strictly
- <sup>2</sup> conserved aspartate residue of M42 aminopeptidase

# 3 catalytic site

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## 13 Abstract

14 The M42 aminopeptidases are a family of dinuclear aminopeptidases widely distributed in Prokaryotes. They have been proposed to be associated to the proteasome, achieving the 15 complete destruction of peptides. Their most peculiar characteristic is their quaternary 16 17 structure, a tetrahedron-shaped particle made of twelve subunits. The catalytic site of M42 aminopeptidases is defined by seven conserved residues. Five of them are involved in metal 18 ion binding which is important to maintain both the activity and the oligomeric state. The sixth 19 conserved residue is the catalytic glutamate acting as a general base deprotonating the water 20 molecule during peptide bond hydrolysis. The seventh residue is an aspartate whose function 21 22 remains poorly understood. This aspartate residue, however, must have a critical role as it is 23 strictly conserved in all MH clan enzymes. In addition, it takes part to some kind of catalytic 24 triad with the histidine residue and the metal ion of the M2 binding site. We assess its role in TmPep1050, an M42 aminopeptidase of *Thermotoga maritima*, through a mutational 25 approach. Asp-62 was substituted with alanine, asparagine, or glutamate residue. The three 26 27 Asp-62 substitutions completely abolished TmPep1050 activity and impeded dodecamer 28 formation. They also interfered with metal ion binding as only one cobalt ion is bound per 29 subunit instead of two. The structural data showed that the Asp62Ala substitution has an 30 impact on the active site folds becoming similar to TmPep1050 dimer. We propose a structural 31 role for Asp-62, helping to stabilize a crucial loop in the active site. In its absence, the loop will 32 remain unstructured failing to position correctly the catalytic base and a metal ion ligand of the33 M1 site.

# 34 **1. Introduction**

The M42 aminopeptidases are dinuclear enzymes proposed to achieve peptide degradation 35 downstream the proteasome in Bacteria and Archaea [1, 2]. In Pyrococcus horikoshii, four 36 37 M42 aminopeptidases (PhTET1, PhTET2, PhTET3, and PhTET4) have been described, each having a different, but complementary, substrate specificity[3-6]. PhTET2 and PhTET3 have 38 39 even been shown to form heterocomplexes, suggesting that peptidasome particles may 40 exist[7, 8]. The M42 aminopeptidases are characterized by a genuine guaternary structure 41 made of twelve subunits organized spatially as a tetrahedron[1, 3–6, 9–13]. The association 42 of the twelve subunits is often described as the assembly of six dimers, each dimer being located at a tetrahedron edge[1]. Four gates are found at the middle of the tetrahedron faces, 43 leading to a wide inner chamber. The gate size (between 12 Å and 20 Å) probably restricts 44 45 the access to the inner chamber to unfolded peptides only[1, 10, 13, 14]. The twelve catalytic sites are oriented inward the chamber, compartmentalizing the activity. The amino acids, 46 generated after peptide hydrolysis, exit the catalytic sites through four channels located at the 47 tetrahedron vertexes. The oligomerization of M42 aminopeptidases is controlled by their metal 48 ion cofactors, as shown for PhTET2, PhTET3, PfTET3 of Pyrococcus furiosus, and 49 50 TmPep1050 of Thermotoga maritima[1, 12, 13, 15, 16]. Indeed, active dodecamers disassemble into inactive dimers upon the depletion of metal ions. The disassembly is 51 52 although reversible as adding metal ions to dimers restores the dodecamer formation and activity. 53

54 The M42 family, ubiquitous to Bacteria and Archaea[17], belongs to the MH clan alongside the M18, M20, and M28[18]. All MH clan enzymes share a common  $\alpha/\beta$  globular 55 fold for their catalytic domain, the archetypal model being the aminopeptidase Ap1 of Vibrio 56 proteolyticus[19]. The typical subunit of M42 aminopeptidases possesses a PDZ-like 57 dimerization domain in addition to the catalytic domain[1]. Seven residues define the catalytic 58 site of MH clan enzymes[18, 20]. Five of them are involved in the binding of two metal ion 59 cofactors. The first metal ion binding site, M1, consists of three conserved residues, a histidine 60 and two aspartates. One of the aspartates is shared with the second metal ion binding site, 61 M2. Two other residues define the M2 site: a conserved histidine and a glutamate/aspartate. 62 These five residues impose the tetrahedral coordination geometry to the two bound metal ions. 63 The implication of both metal ions in the catalytic mechanism has been thoroughly studied in 64 65 V. proteolyticus aminopeptidase Ap1[21-23]. Indeed, they interact with the free N-terminal 66 amine and the first carbonyl of the peptide substrate. The M2 has been proposed to assist the

water molecule deprotonation. The hydroxide is then transferred to the M1 for peptide bond hydrolysis. In M42 aminopeptidases, the two metal ions are also involved in the dimerdodecamer transition. We previously showed that the M1 strictly controls the oligomerization of TmPep1050 while M2 could have a stabilizer role[13]. Intriguingly, their roles were swapped in PfTET3[12], suggesting that other factors may influence oligomerization.

72 In addition to the five metal ion ligands, two other residues, a glutamate and an aspartate, define MH clan enzymes. These two residues are not involved directly in metal ion 73 binding but are strictly conserved in the whole clan. The glutamate residue acts as a general 74 base deprotonating the water molecule during the peptide bond hydrolysis[24]. It may also 75 play a role in decreasing the Lewis acidity of the M2. As proposed for V. proteolyticus 76 aminopeptidase Ap1, the catalytic base Glu-151 is within hydrogen bond distance to interact 77 78 with His-97[21, 25]. The aspartate residue is not directly involved in hydrolysis but may have 79 a role in decreasing the Lewis acidity of the M2[26]. In V. proteolyticus aminopeptidase Ap1, Asp-99 has been proposed to form some kind of catalytic triad (hereafter mentioned as Asp-80 His-Me) with His-97 and Zn<sup>2+</sup> in the M2 binding site[21, 23, 25, 27]. Indeed, a strong hydrogen 81 82 bond links the O<sup>81</sup> atom of Asp-99 with the N<sup>8</sup> atom of His-97, forcing the imidazole ring to be deprotonated. As a result, the Lewis acidity of Zn<sup>2+</sup> would be decreased sufficiently to allow 83 84 the transfer of the hydroxide to the M1 site, which is closer to the peptide bond to be hydrolyzed. 85

Still, how the aspartate residue participates in peptide bond hydrolysis remains 86 87 intricate. Its role has only been hypothesized for *V. proteolyticus* aminopeptidase Ap1. To our 88 knowledge, nothing has been well established about its implication in metal ion binding or 89 active site stabilization for MH clan enzymes. In this work, we propose to study the Asp-His-Me triad of TmPep1050, an M42 aminopeptidase from *T. maritima*. TmPep1050 is a leucyl-90 91 aminopeptidase whose activity depends on its oligomeric states: inactive dimer and active 92 dodecamer[13]. The transition between dimers and dodecamers is regulated by its metal ion cofactor, Co<sup>2+</sup>. We present the characterization of three TmPep1050 variants for which Asp-93 62, equivalent to Asp-99 of V. proteolyticus aminopeptidase Ap1, was substituted with alanine. 94 asparagine, or glutamate residue. All substitutions completely abolished the activity and 95 dodecamer formation of TmPep1050. The Asp-62 variants were still able to bind one cobalt 96 ion per subunit. As dodecamer formation relies on the M1 site, we propose that the M2 site 97 remained functional despite the substitution. The structure of TmPep1050<sub>D62A</sub> was solved and, 98 99 by comparing with the dimer and dodecamer structures, a structural role was inferred for Asp-100 62.

# 101 2. Materials and Methods

#### 102 **2.1. Mutagenesis of** *Tm***\_1050**

Site-directed mutagenesis was performed following the single-primer reactions in parallel protocol (SPRINP)[28] using the pCEC43 as template vector. This vector allows the production of TmPep1050 under the regulation of P<sub>BAD</sub> promoter in *E. coli*[13]. The primers used to generate the Asp-62 variants of TmPep1050 are described in Table S1. Phusion DNA polymerase (ThermoFisher Scientific) was used for DNA polymerization reactions. Cloning was carried out in the *E. coli* MC1061 strain[29]. The genetic constructs were checked by sequencing (Genetic Service Facility, University of Antwerp).

#### 110 2.2. Production and purification

The Asp-62 variants of TmPep1050 were produced and purified to homogeneity according to 111 the protocol described elsewhere [13]. It consists of four steps: (i) a heat treatment of crude 112 cell extract at 70°C for 15 min., (ii) an anion-exchange chromatography using Source 15Q 113 resin (GE Healthcare Life Sciences), (iii) an hydrophobic interaction chromatography using 114 Source 15Phe resin (GE Healthcare Life Sciences), and (iv) a size-exclusion chromatography 115 using a Superdex200 (GE Healthcare Life Sciences, XK16/20 column of 120-ml volume). 116 During the last step of purification, an elution peak was observed at 95 ml for the three Asp-117 62 variants. The purified Asp-62 variants were concentrated to 250 µM using an Amicon Ultra-118 119 15 ultrafiltration unit with 30-kDa cutoff (Merck Millipore). Protein concentration was estimated 120 by measuring the absorbance at 280 nm and applying the mass extinction coefficient of 18,910 121 M<sup>-1</sup> cm<sup>-1</sup>. For molecular weight determination, 50 µM of purified protein was loaded on a Superdex200 (GE Healthcare Life Sciences, XK16/20 column of 120-ml volume) using 50 mM 122 MOPS, 0.5 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 1 mM CoCl<sub>2</sub>, pH 7.2 as running buffer. The size exclusion column 123 was calibrated using Gel-filtration standard (Biorad) and High-molecular-weight gel filtration 124 calibration kit (GE Healthcare Life Sciences) under the same running conditions. 125

#### 126 2.3. Mass spectrometry

Sample preparation and native MS analysis were performed as previously described[13]. TmPep1050<sub>D62A</sub> was conditioned in 20 mM Ammonium acetate, pH 7.2 using Zeba 7-kDa desalting columns (Thermo Fisher Scientific). Prior to MS analysis, the sample was diluted to  $5 \mu$ M in 20 mM Ammonium acetate, pH 7.2. The spectrum was recorded in positive ion mode on a traveling-wave ion mobility Q-TOF instrument (Synapt G2 HDMS, Waters).

#### 132 2.4. Enzymatic and cobalt binding assays

133 The specific activity of Asp-62 variants was assayed using L-leucine-*p*-nitroanilide (L-Leu-134 *p*NA, Bachem AG) as substrate. 10  $\mu$ I of 1  $\mu$ M enzyme was added to 990  $\mu$ I of reaction mix 135 containing 2.5 mM L-Leu-pNA in 50 mM MOPS, 10 % methanol, 1 mM CoCl<sub>2</sub>, pH 7.2. The 136 reaction was performed at 75°C during 1 h and stopped by adding 1 ml of 20% acetic acid. 137 Released *p*-nitroaniline was quantified by measuring the absorbance at 410 nm. Prior to cobalt binding assays, apo-enzyme was prepared as followed for the three Asp-62 variants. 200 µl 138 of 250 µM enzyme was diluted in 1.8 ml of 50 mM MOPS, 0.5 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 10 mM 1,10-139 phenanthroline, pH 7.2. The sample was then dialyzed four times against 200 ml of 50 mM 140 MOPS, 0.5 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, pH 7.2. After the dialysis, the sample was concentrated to 100 µl 141 using an Amicon Ultra-15 ultrafiltration unit with 30-kDa cutoff (Merck Millipore). Protein 142 concentration was determined as described above. To study the metal binding, apo-enzyme 143 was diluted to 20 µM in 50 mM MOPS, 0.5 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, pH 7.2 and CoCl<sub>2</sub> at a concentration 144 ranging from 0 to 1280 µM. The reaction volume was 140 µl and the samples were incubated 145 at 50°C for 24 h. The thermostability of the samples was determined by Thermal Shift Assay 146 using SyproOrange (Thermofisher Scientific) as described previously[13]. Bound Co<sup>2+</sup> was 147 quantified using Amplex UltraRed fluorescent probe (ThermoFisher Scientific) as described 148 previously[13]. 149

#### 150 **2.5. X-ray crystallography**

TmPep1050<sub>D62A</sub> was crystallized in 0.1 M citric acid, 5% PEG3350, pH 5.2 using the hanging 151 drop diffusion method. Drops containing 2 µl of 250 µM TmPep1050<sub>D62A</sub> and 2 µl of 152 crystallization reagent were set up in EasyXtal Tool plates (Qiagen). Fully grown crystals were 153 obtained after a week at 292 K. Crystal shape and size were improved by microseeding. 154 Diffraction data were collected on the FIP-BM30a beamline at the European Synchrotron 155 Research Facility (Grenoble, France)[30, 31]. Data collection and refinement statistics are 156 157 presented in Table 1. XDS program package[32] was used for indexing diffraction data. The 158 phase was determined by molecular replacement using Phaser[33] with the monomer A 159 coordinates of TmPep1050 dodecamer structure (PDB code 4P6Y). The model was build 160 using phenix.autobuild in PHENIX software package[34]. The model was iteratively refined manually with Coot[35] and automatically with *phenix.refine*. Model stereochemical quality 161 was assessed using MolProbity[36]. Protein structures were analyzed with PDBe Pisa[37], 162 Arpeggio[38], and PyMOL Molecular Graphics System version 2.2 (Schrödinger,LLC). 163

# **3. Results and discussion**

#### **3.1.** The role of Asp-62 in the activity and oligomerization of TmPep1050

Asp-62, equivalent to Asp-99 of *V. proteolyticus* aminopeptidase Ap1, was substituted with alanine, asparagine, or glutamate residue. The resulting variants were named TmPep1050<sub>D62A</sub>, TmPep1050<sub>D62N</sub>, and TmPep1050<sub>D62E</sub>, respectively. The variants were recombinantly produced in *E. coli* and purified to homogeneity. Their molecular weights were 170 determined by size-exclusion chromatography (Figure 1). The substitution of Asp-62 had a 171 dramatic impact on the TmPep1050 oligomerization state as only dimers were observed even 172 in the presence of Co<sup>2+</sup>. The oligomerization state of TmPep1050<sub>D62A</sub> was confirmed by native MS showing dimers mainly (Figure 2). MS experiments also highlighted a minor tetrameric 173 174 form. Minor intermediate oligomers (tetramer, hexamer, and octamer) were reported previously for the wild-type enzyme[13]. These oligomers were inferred as intermediate forms 175 occurring during the dimer-dodecamer transition. For TmPep1050<sub>D62A</sub>, only tetramers were 176 observed as intermediates, supporting the impaired oligomerization of the Asp-62 substitution. 177 178 The three variants displayed less than 0.04% of relative activity on L-Leu-pNA compared to the dodecameric wild-type enzyme. 179

The loss of activity subsequent to Asp-62 substitution was expected according to the 180 putative function of Asp-99 in V. proteolyticus Ap1. The negative charge and the side chain 181 length are critical for the activity. Intriguingly, Asp-62 substitution prevents the self-assembly 182 of dimers into dodecamers. Such an observation contrasts with the known role of the two metal 183 184 ion binding sites in TmPep1050 oligomerization. Indeed, we previously demonstrated that the 185 M1 binding site strictly controls the oligomerization as the His307Ala substitution prevented 186 the self-assembly of dimers[13]. The substitution of His-60, a residue of the M2 binding site, perturbed the oligomerization to a lesser extent as both dimers and dodecamers were 187 observed. As Asp-62 interacts with His-60, we could have expected that the Asp-62 188 substitution would have a rather small impact on oligomerization like His-60 substitution. 189 Unexpectedly, the Asp-62 variants behaved as if the M1 binding site had been impaired as 190 191 only dimers were observed. Therefore, Asp-62 substitution could somehow interfere with the 192 metal ion binding in both M1 and M2 binding sites.

#### 193 **3.2.** Asp-62 substitution partly impaired the Co<sup>2+</sup> binding

194 To show the potential impact of Asp-62 substitution on Co<sup>2+</sup> binding, thermal stability of Asp-62 variants were measured by thermal shift assay in presence of various concentrations of 195 Co<sup>2+</sup>. Using such a method, we previously showed that TmPep1050 dimers depleted of cobalt 196 were more prone to thermal denaturation than TmPep1050 dodecamers, with melting 197 temperatures of 91°C and 97°C, respectively. The addition of Co<sup>2+</sup> restored the thermal 198 stability of TmPep1050 dimers[13]. Prior to the binding assays, the Asp-62 variants were 199 treated with 1,10-phenanthroline to remove any bound  $Co^{2+}$ . Then 20  $\mu$ M of each apo-enzyme 200 was incubated in presence of Co<sup>2+</sup> at a concentration ranging from 0 to 1280 µM for 24 h. The 201 melting temperatures were subsequently determined to monitor an eventual cobalt binding. 202 203 Apo-TmPep1050<sub>D62N</sub> had a melting temperature of 89.1  $\pm$  0.4°C, close to that of the wild-type 204 dimer. Apo-TmPep1050<sub>D62A</sub> and apo-TmPep1050<sub>D62E</sub> had a melting temperature of 86 ± 0.1°C 205 and 83.2 ± 0.2°C, respectively, indicating that the Asp62Ala and Asp62Glu substitutions

destabilized partly the structure. As the thermal stability increased with the cobalt concentration (Figure 3), the three variants were found to bind cobalt ions. Of note, none of them exhibited a thermostability comparable to the dodecamer upon Co<sup>2+</sup> binding. Indeed, the maximal melting temperature of TmPep1050<sub>D62N</sub>, TmPep1050<sub>D62E</sub>, and TmPep1050<sub>D62A</sub> was 95.1°C, 91.1°C, and 90.7°C, respectively, for a Co<sup>2+</sup>-to-protein molar ratio of 50.

Bound Co<sup>2+</sup> was guantified to define how many metal ion binding sites are occupied in 211 each variant. 20 µM of apo-enzyme was incubated with varying Co<sup>2+</sup> concentration ranging 212 from 0 to 1280 µM. The concentration of bound Co<sup>2+</sup> was determined at the equilibrium using 213 Amplex Ultra Red fluorescent probe for quantification. The results are showed in Figure 4. 214 TmPep1050<sub>D62A</sub>, TmPep1050<sub>D62N</sub>, and TmPep1050<sub>D62E</sub> were able to bind Co<sup>2+</sup> with an 215 apparent association constant of 70  $\pm$  12, 86  $\pm$  2  $\mu$ M, and 87  $\pm$  13  $\mu$ M, respectively. Only one 216 cobalt ion was bound per monomer (maximal bound Co<sup>2+</sup> being 1.3), suggesting that one of 217 the two metal binding sites is impaired in the Asp-62 variants. In comparison, wild-type 218 TmPep1050 could bind two cobalt ions with an apparent association constant of 50 µM[13]. 219 220 For the Asp-62 variants, the cobalt quantification did not allow to determine whether both M1 221 and M2 sites were partially occupied or one of them remained vacant. The second hypothesis 222 is the likeliest based on (i) the dimeric state of Aps-62 variants and (ii) the M1 site controlling 223 the dimer-dodecamer transition. Thus, the M2 site was inferred to be functional even when Asp-62 was substituted. Our results contrast with what has been reported for the human 224 copper/zinc superoxide dismutase. This enzyme displays an Asp-His-Me triad analogous to 225 that of the MH clan enzymes. The substitution of the aspartate residue with an asparagine or 226 227 alanine residue completely abolished both the activity and the metal ion binding[39].

# 3.3. The structure of TmPep1050<sub>D62A</sub> reveals how Asp-62 maintains the structural fold of the active site

To understand how Asp-62 could influence Co<sup>2+</sup> binding in the M1 site, we solved the structure 230 of TmPep1050<sub>D62A</sub> at 1.5 Å. The crystallization condition was 0.1 M citric acid, 5% PEG3350, 231 pH 5.2. Unfortunately, X-ray fluorescence scanning of crystals did not detect any trace of 232 metal ion, probably due to cobalt chelation by citric acid[40]. According to PDBe Pisa, the 233 crystal structure confirmed the dimeric state of TmPep1050<sub>D62A</sub>. When compared to 234 235 TmPep1050 dodecamer, the subunit structure of TmPep1050<sub>D62A</sub> presents the same structural 236 dissimilarities as TmPep1050 dimer (Figure 5A)[13]. Two segments, Lys-229 - Ala-235 and 237 Lys-247 – Ser-254, diverge greatly between both structures. For instance, the backbone is 238 displaced by 3 Å and 11 Å for Lys-232 and Arg-249, respectively. Both segments were 239 described as critical in the oligomerization as they are involved in the interaction between dimers. The Asp-62 substitution also affects the catalytic pocket fold as the  $\alpha$ 8 and  $\alpha$ 10 helices 240

are too flexible to be modelled. Furthermore, the Gln-196 – Val-202 loop is displaced outwards
 the catalytic site, explaining the lack of activity of TmPep1050<sub>D62A</sub>.

Because of Gln-196 - Val-202 loop displacement, Glu-197 and Glu-198 are not 243 correctly positioned to fulfill their function (Figure 5B). The former is the catalytic base, 244 245 assisting the deprotonation of a water molecule during the first step of the peptide bond hydrolysis. The latter is involved in the binding of a cobalt ion at the M1 site, along with His-246 247 307 and Asp-168. In the TmPep1050<sub>D62A</sub> structure, the His-307 side chain adopts a m-70° 248 rotamer, compared to the **m**170° rotamer observed in the dodecamer structure. Consequently, the imidazole ring is tilted by about 28° between both structures. Asp-168 belongs to both M1 249 and M2 sites as it bridges the two cobalt ions. The Asp-62 substitution did not have any effect 250 251 on the Asp-168 position. Regarding the M2 site, the side chains of Asp-220 and His-60 are oriented differently in TmPep1050<sub>D62A</sub> compared to dodecameric TmPep1050. The Asp-220 252 side chain adopts an **m**-20° rotamer instead of a **t**0° rotamer found in the dodecamer structure. 253 Due to side chain orientation change, the carboxylate center is displaced by 2.4 Å and 254 255 interacts with the carbonyl of Ile-221. In the monomer A of TmPep1050<sub>D62A</sub>, the imidazole ring of His-60 is correctly positioned with a p-80° rotamer but is rotated by 180°. Consequently, N<sup>81</sup> 256 and N<sup>2</sup> of His-60 are found interacting with two water molecules which are not observed in 257 the dodecamer structure. In the monomer B, His-60 side chain adopts two alternate rotamers, 258 m90° and p-80°, with an occupation of 0.57 and 0.43, respectively. 259

260 Excepting the alternate rotamers of His-60, the TmPep1050 dimer structure shows the 261 same dissimilarities in the active site as TmPep1050<sub>D62A</sub>. Despite being structurally similar, TmPep1050 dimers are still able to bind two cobalt ions and revert to dodecamers while 262 TmPep1050<sub>D62A</sub> cannot. Hence the Asp-62 substitution must have a structural role for 263 stabilizing the metal-bound form. In the dodecamer structure, O<sup>81</sup> and O<sup>82</sup> of Asp-62 are in 264 tight H-bond interaction with the backbone nitrogen of Glu-197 and Glu-198, respectively 265 (Figure 5B). Thus, Asp62Ala substitution probably prevents this interaction and, consequently, 266 267 could not stabilize the GIn-196 – Val-202 loop. It has a dramatic impact on the M1 site as Glu-198 cannot be correctly positioned to allow the binding of a cobalt ion. The side chain length 268 and charge of residue 62 are critical since TmPep1050<sub>D62N</sub> and TmPep1050<sub>D62E</sub> are also 269 270 inactive dimers. The position of the Asp-62 carboxylate is probably imposed by H-bond 271 interactions with Ser-15 and His-60. In the TmPep1050<sub>D62A</sub> structure, Ser-15 hydroxyl is rotated by 107.2° compared to wild-type dimer and dodecamer. In addition, charge repulsion 272 273 may occur between Glu-18 and Asp-62. Placing a glutamate residue at position 62 would not 274 allow the correct alignment of the carboxylate with Ser-15 and His-60. In the case of an asparagine residue at position 62, the amide group would not sustain the charge repulsion 275

from Glu-18. Furthermore, the amine group would clash with either the Ser-15 or His-60 dueto short H-H distance.

## 278 **4. Conclusion**

The carboxylate-histidine-metal ion triad is commonly encountered in metalloenzymes. 279 Christianson and Alexander reported that at least 35% of histidine residues bound to a metal 280 ion interacts with either a glutamate or an aspartate residue [26]. The supposed role of the 281 carboxylate is to modulate the basicity of the neighbor histidine residue, affecting the Lewis 282 283 acidity of bound metal ion. It may also have an impact of metal ion binding affinity and 284 nucleophilicity of the water molecule bound to the metal ion[41, 42]. The MH clan enzymes 285 display such a carboxylate-histidine-metal ion triad at the M2 site. Both the aspartate and 286 histidine residues of the triad are strictly conserved in the whole clan. A function of the aspartate residue has been postulated for the MH clan archetypal enzyme, the V. proteolyticus 287 aminopeptidase Ap1[21-23, 25, 27]. Asp-99 has been described to decrease Lewis acidity of 288 the M2 site Zn<sup>2+</sup> via a strong H-bond between Asp-99 and His-97. Consequently, the 289 interaction between the nucleophilic hydroxide and the M2 site Zn<sup>2+</sup> is disfavored and may 290 transitorily alter the coordination geometry of the metal ion. 291

292 In this work, we showed that the aspartate residue of the Asp-His-Me triad could have 293 another function for the M42 aminopeptidases. Using TmPep1050 as a case of study, Asp-62, equivalent to Asp-99 of V. proteolyticus aminopeptidase Ap1, was substituted with alanine, 294 asparagine, or glutamate residue. This substitution completely abolished the activity, as 295 expected, but also interfered in dodecamer formation. We suppose that the Asp-62 296 297 substitution affects the M1 binding site as the oligomerization of TmPep1050 strictly relies on 298 the M1. The structure of TmPep1050<sub>D62A</sub> presented the same structural dissimilarities as 299 observed for the wild-type dimer. Notably, the GIn-196 – Val-202 loop is disordered, preventing 300 the correct positioning of Glu-197 and Glu-198 (the catalytic base and a ligand of the M1 site). Asp-62 has probably a structural role for stabilizing this active site loop since its carboxylate 301 function is in close interaction with the backbone nitrogen of Glu-197 and Glu-198. 302 Nevertheless, we could not ascertain the structural role of Asp-62 to all MH clan enzymes, as 303 304 no structure of equivalent variant has been reported in other families. It would be interesting to know whether substituting the aspartate residue would destabilize the active site of a 305 306 monomeric enzyme like V. proteolyticus aminopeptidase Ap1. If the destabilization is also 307 observed for such an enzyme, it will support the structural role of this strictly conserved aspartate residue thorough the whole MH clan. If not, it will indicate that the destabilization of 308 the active site as observed for TmPep1050<sub>D62A</sub> could be an adaptation to oligomerization. 309

- Nevertheless, both hypotheses agree with the dependency of M42 aminopeptidase
- 311 oligomerization on the active site fold.

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|   | TmPep1050 <sub>D62A</sub> |
|---|---------------------------|
| Data collection                                 |                           |
| Temperature (K)                                 | 100                       |
| Radiation source                                | ESRF BM30A                |
| Wavelength                                      | 0.9797                    |
| Detector  | ADSC QUANTUM 315r         |
| Oscillation range (°)                           | 0.5                       |
| Exposure time (s)                               |                           |
| Space group                                     | C 2 21 1                  |
| Unit cell parameters                            |                           |
| α, β, γ (°)                                     | 90.00, 90.00, 90.00       |
| a, b, c (Å)                                     | 42.18, 113.96, 267.23     |
| Resolution                                      | 48.00 – 1.50 (1.58-1.50)  |
| Unique reflections                              | 103,939                   |
| R <sub>merge</sub> (%)                          | 0.09                      |
| Redundancy                                      | 5.8                       |
| <i <b="">σ&gt;</i>                              | 14.80 (3.34)              |
| Completeness (%)                                | 99.7 (89.4)               |
| CC1/2 (%)                                       | 99.8 (87.6)               |
| Refinement                                      |                           |
| Resolution                                      | 48.00 – 1.50              |
| Reflections                                     | 103,177                   |
| R <sub>free</sub> set test count                | 5,158                     |
| Rwork/Rfree                                     | 0.183/0.215               |
| Protein molecules per ASU                       | 2                         |
| V <sub>M</sub> (Å <sup>3</sup> /Da)             | 2.27                      |
| Solvent content (%)                             | 45.79                     |
| Protein/solvent atoms                           | 4,950/820                 |
| r.m.s.d. bond lengths (Å)                       | 0.36                      |
| r.m.s.d. bond angles (°)                        | 0.55                      |
| Average B-factors (Å <sup>2</sup> )             | 20.0                      |
| Favored/disallowed Ramachandran $\phi/\psi$ (%) | 97.71/0.00                |
| PDB code  | 5L6Z                      |

**Table 1** Data collection and refinement statistics. Values in parentheses are for the highest-

432 resolution shell.



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Figure 1 Size exclusion chromatography of TmPep1050<sub>D62A</sub> (plain line), TmPep1050<sub>D62N</sub> (dot line), and TmPep1050<sub>D62E</sub> (dashed line). An elution peak was observed at 95 ml for the three variants using a Superdex 200 column (V = 120 ml). *Inset:* The column was calibrated using albumin (Ab), conalbumin (C), aldolase (Ad), ferritin (F), and thyroglobulin (T) as standards. The correlation between the logarithm of the relative mass and the elution volume is linear, with a R<sup>2</sup> of 0.91. The 95% confidence intervals are represented as dots.





- The sample was conditioned in 20 mM Ammonium acetate and diluted to 5 μM prior to native
- 443 MS analysis.

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Figure 3 Thermal shift assay showing  $Co^{2+}$  binding effect on Asp-62 variant thermostability. The thermostability of TmPep1050<sub>D62A</sub> (circle), TmPep1050<sub>D62N</sub> (square), and TmPep1050<sub>D62E</sub> (triangle) was determined by measuring T<sub>m</sub> at each Co<sup>2+</sup> concentration. From our previous study, the thermostability of wild-type TmPep1050 dimer is presented as rhombi[13]. Error bars represent standard error with n = 3.



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Figure 4 Co<sup>2+</sup> binding of apo-TmPep1050<sub>D62A</sub> (circle), apo-TmPep1050<sub>D62N</sub> (square), and apo-TmPep1050<sub>D62E</sub> (triangle). The results are expressed as the number of bound Co<sup>2+</sup> per molecule of enzyme (mol. enz.) in response to a varying concentration of Co<sup>2+</sup> ranging from 0 to 1280  $\mu$ M. Error bars represent standard error with n = 3.

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Figure 5 The structure of TmPep1050<sub>D62A</sub>. (A) Structural alignment of TmPep1050<sub>D62A</sub> subunit (PDB code 5L6Z, light grey) with TmPep1050 dodecamer subunit (PDB code 6NW5, green), and TmPep1050 dimer subunit (PDB code 5NE6, light blue). The structural dissimilarities are indicated with arrows. Zn<sup>2+</sup> and Co<sup>2+</sup> are represented as grey and pink spheres, respectively. (B) Close-up view of the active sites of TmPep1050<sub>D62A</sub> (light grey), TmPep1050 dodecamer (green) and dimer (light blue). The water molecules interacting with His-60 in the TmPep1050<sub>D62A</sub> structure are showed as red spheres.