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Partitioning the structural features that underlie expansin-like and elicitor activities of cerato-platanin

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Partitioning the structural features that underlie expansin-like and elicitor activities of Cerato-platanin

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Abstract

Cerato-platanin family (CPF) proteins are produced by fungi and elicit defences when applied to plants, behaving as PAMPs/MAMPs. CPF proteins share structural similarity to plant and bacterial expansins, and have been demonstrated, in some cases, to possess expansin-like loosening activity on cellulose. This is the case of cerato-platanin (CP), the founder of the CPF, which shows both eliciting and cellulose-loosening activities, raising the question as to whether the expansin-like activity may be responsible for defense activation. To pinpoint structural and thermodynamic features underlying eliciting and expansin-like activity of CP, we carried out site-directed mutagenesis targeting separately net charge (N84D mutation), conformational stability (V63A mutation), or conserved position previously shown to affect expansin-like activity in CP (D77A mutation), and characterized wild-type protein and its variants. Removing or adding negative charges on the protein surface led to reducing or increasing, respectively, the expansin-like activity. The activity was instead not affected by mutations affecting protein fold and stability. In contrast, all the mutants showed reduced capacity to elicit defences in plants. We conclude that the expansin-like activity of CP depends on net charge and ability to (weakly) bind cellulose, whereas the eliciting activity on plants does not depend on the cellulose-loosening activity.

Key words: Cerato-platanin; expansin-like activity; PAMPs; cellulose-binding; *E.Coli* SHuffle

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1. Introduction

Plants protect themselves from potential pathogenic microorganisms through a complex interplay of signaling pathways and defense responses, which include mitogen-activated protein kinase (MAPK) cascades, transcription of defence related genes, production of reactive oxygen species (ROS) and nitric oxide, and the synthesis of defensive compounds such as phytoalexins [1,2].

To detect microorganisms, the plant immune system exploits microbial molecular patterns commonly known as pathogen- or microbe-associated molecular patterns (PAMPs or MAMPs) [3,4]. PAMPs/MAMPs are evolutionarily stable macromolecules in microbes, such as for instance fungal chitin, β -glucans or bacterial flagellin. Furthermore, other plant cell wall molecules, released during the interaction between the plant and the pathogen, such as galactoglucomannan or oligogalacturonides (DAMPs, damage-associated molecular patterns) may trigger the plant immune system activation [5]. As a result, PAMPs/DAMPs allow the plant to switch from a developmental status to a defensive mode, able to reject potentially harmful microbes [6].

The cerato-platanin family (CPF) members are cysteine-rich proteins secreted and localized on fungal cell walls [7]. These proteins are involved in fungal development and in the interactions between fungi and host plants, functioning as PAMPs and inducing canonical defence mechanisms such as the production of ROS and phytoalexins or the transcription of defence related genes [8–10].

Researchers worldwide have identified more than 100 genes belonging to CPF proteins in fungi over the last 20 years [7]. Nevertheless, a thorough structural and functional characterization at the protein level has been performed up to date only for a few members of the family [7]. Among the most studied proteins we may mention cerato-platanin (CP) from *Ceratocystis platani*, the founder member of the CPF [11,12], MgSM1 from *Magnaporthe oryzae* [13], BcSpl1 from *Botrytis cinerea* [14] and FgCPPs from *F. graminearum* [15] among others.

CP shows a globular fold with two α -helices and six β -strands organized in a peculiar double $\psi\beta$ -barrel (DPBB) fold that is highly conserved in all the CPF members and it is also present in endoglucanases, in plant and bacterial expansins, and in the defense protein barwin [16]. Studies on the interaction between CP and its substrates highlight the presence of a flat and shallow groove in the CP fold which is placed on a side of the β -barrel and is rich in polar and aromatic residues that are able to interact with carbohydrates such as N-acetylglucosamine, thus explaining the ability of CP to bind cell wall fungal chitin [17]. The structural similarity of CP with the D1 domain of bacterial and plant expansins is of remarkable interest to unravel the role of CP in plant interaction. Expansins are proteins of about 26 kDa consisting of two domains: a C-terminal domain (D2) forming an Ig-like β -sandwich and an N-terminal domain (D1) which forms a double $\psi\beta$ -barrel fold similar to the CP domain [18]. Expansin-like proteins have been found in several fungi and bacteria where they are able to fragment cellulose aggregates with no evidence of lytic activity [18]. Intriguingly, like expansins, CP displays cellulose-loosening activity and seems to exert various roles in fungal growth and developmental processes of the fungus [19].

In a previous structure/function correlation study performed with the use of the D77A mutant of CP [17], we hypothesized that the ability to induce defences in plants and the expansin-like activity were closely related

1 features of CP. The D77A mutant was designed by considering that D77 is a highly conserved residue both in
2 the CP family and among expansin-like proteins and it lies on the hydrophilic groove in the region of the barrel
3 involved in cellulose loosening [16,20]. The lack of both eliciting and expansin-like activities exhibited by the
4 D77A mutant suggested a correlation between activity on cellulose and protein perception by the plant immune
5 system [17]. Notwithstanding that, the mutation D77A was affected in both its net charge and its
6 conformational stability, making impossible to identify the predominant effect responsible for the loss of
7 activity.
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11 In order to pinpoint the structural and thermodynamic features underlying eliciting and expansin-like activity
12 of CP, in the present study we exploited a mutagenesis approach by which we rationally designed single point
13 mutations in CP targeting separately net charge, conformational stability, or evolutionarily conserved
14 positions. Accordingly, we first produced the V63A mutation: valine is located in the hydrophobic core of CP
15 and the mutation to alanine was expected to alter protein stability while leaving charge unaffected. In addition,
16 we produced the mutant N84D basing on the observation that endoglucanases, lytic transglycosylases and
17 other proteins sharing a similar fold with CP have an aspartate residue at position 84, whereas asparagine is
18 present in CP in a conserved position [16]. Moreover, in the D1 domain of the expansin BsEXLX1, the residues
19 D71, Y73 and D82 are required for creep activity and paper weakening (Georgelis et al., 2015) and are
20 structurally related to the aromatic and hydrophilic residues forming the shallow groove probably involved in
21 substrate recognition in CP. Therefore, we mutated asparagine 84 to aspartic acid (N84D), with the aim of
22 determining whether both the position and the net charge are responsible for the expansin-like activity in CP.
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34 **2. Materials and Methods**

35 **2.1 Site-directed mutagenesis of CP and mutant purification.**

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38 XL10 Gold *E. coli* competent cells (Agilent Technologies, Santa Clara, CA, USA) were used for site-directed
39 mutagenesis using the *cp*-pGEX-2T plasmid from *E. coli* SHuffleT7 cells [17]. Mutagenesis was carried out
40 using the QuikChange Lightning Site-Directed Mutagenesis Kit (Agilent, Santa Clara, CA, USA), following
41 the instructions from the manufacturer. The following primers, synthesized by Eurofins Genomics (Ebersberg,
42 Germany) were used:
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48 V63A:

49 5'-CCGTTGGGAATCGTGGCCTTCCAGCAAGTAC-3'

50 5'-GTACTTGCTGGAAGGCCACGATTCCCAACGG-3'

51 N84D:

52 5'-GCAGTAGGATTCACGTCGAATCCACCTCTGCCA-3'

53 5'-TGGCAGAGGTGGATTTCGACGTGAATCCTACTGC-3'

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55 The correct nucleotide sequences were checked by DNA sequencing (Eurofins Genomics, Ebersberg,
56 Germany) and the resulting plasmids were transformed separately into *E. coli* SHuffleT7 cells (New England
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1 Biolabs, Ipswich, MA, USA)[17]. Large-scale expression and purification of mutants and wild type (wt) CP
2 were performed according to the protocol reported in Luti et al., [17].
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4 **2.2 Structural characterization of mutants and interaction with sugars**

5 *Circular dichroism spectra.* CD spectra were recorded in the far-UV region from 190 to 260 nm, using a
6 JASCO J-810 spectropolarimeter (Tokyo, Japan). The proteins were dissolved in water to a final concentration
7 of 6.25 μM . The spectra were acquired using a 1 mm path length quartz cell, a band-width of 1 nm, a response
8 time of 0.5 s, a data pitch of 0.5 nm and a scanning speed of 100 nm min^{-1} . Each spectrum was the average of
9 six consecutive scans followed by subtraction of the blank spectra (water). The observed signal –in mdeg– was
10 converted into mean residue molar ellipticity (expressed in $\text{deg cm}^2/\text{dmol}$), according to the formula $[\theta] = \text{CD}$
11 $/ [(c \text{ m}^{-1}) 10 d n]$, where CD is the observed signal, c is the protein concentration in g/L, m is the molar mass
12 in g mol^{-1} , d is the path length in cm and n is the number of amino acid residues.
13

14 *Differential Scanning Fluorimetry.* Differential scanning fluorimetry (DSF) was performed using a 7500 Real-
15 Time PCR machine (Applied Biosystems, Foster City, CA, USA) by selecting the ROS filter. A master mix
16 solution was prepared with 0.133 mg mL^{-1} of protein and 1:500 of SYPRO Orange (ThermoFisher Scientific,
17 Waltham, MA, USA) in 50 mM Sodium Acetate pH 5.0. 15 μL of this mixture were added to 5 μL of buffer
18 to compare wtCP with mutants, and 5 μL of different concentration of sugar polymers in a MicroAmp Fast
19 Optical 96-Well Reaction Plate (# 4346906), sealed with MicroAmp Optical Adhesive Film (all from
20 ThermoFisher Scientific, Waltham, MA, USA) in a final volume of 20 μL for each measurement. All samples
21 were prepared in duplicate. The samples were heated with a ramp speed that provides a 2 min pause at 35 $^{\circ}\text{C}$,
22 followed by a ramp to 95 $^{\circ}\text{C}$ over 90 min. The Sypro orange fluorescence was plotted versus temperature
23 (melting curve) and the resulting plot converted into its first order derivative, which provided the melting
24 temperature (T_m) as a positive peak.
25

26 *Equilibrium denaturation experiment.* For each of the variants tested, we prepared 20 samples containing the
27 protein at a concentration of 0.1 mg mL^{-1} and guanidine hydrochloride (GndHCl) at molar concentrations
28 ranging from 0 to 6 M in a 50 mM sodium acetate buffer at pH 5.0. We incubated the samples for 30 minutes
29 at 25 $^{\circ}\text{C}$ and acquired tryptophan emission spectra from 300 to 500 nm using a Synergy H1 hybrid multimode
30 reader (BioTek, Winooski, VT) and an excitation wavelength of 280 nm. We analysed data by calculating the
31 centre of mass (COM) of each spectrum as $\text{COM} = (\sum_i F_i) / (\sum_i \nu_i F_i)$, where F_i is the fluorescence emitted at
32 a wavenumber of ν_i . The values obtained were plotted vs GndHCl concentration and analysed with the equation
33 edited by Santoro & Bolen [22], to yield the difference in free energy between folded and unfolded states in
34 the absence of denaturant ($\Delta G_{\text{U-F}}^{\text{H}_2\text{O}}$), the dependence of $\Delta G_{\text{U-F}}$ on denaturant concentration (m), and the
35 concentration of mid-denaturation (C_m). We normalized data to the fraction of folded protein and plotted
36 results.
37

38 *Stopped-flow unfolding kinetics.* We followed unfolding of the CP variants in real-time in the presence of
39 Ficoll or Carboxymethyl cellulose (CMC). For this experiment, we exploited a Bio-Logic SFM-3 stopped-
40 flow device (Claix, France) equipped with an FC-08 (0.8 mm) cuvette and coupled to a fluorescence detection
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1 system. Briefly, we prepared the protein at an initial concentration of 0.3 mg mL⁻¹ in a 50 mM sodium acetate
2 buffer at pH 5.0. Unfolding was triggered by a 10-fold dilution of the protein into solutions containing GndHCl
3 and different amounts of either polysaccharide. Final conditions were 0.03 mg mL⁻¹ protein in 4.2 M GndHCl
4 with a polysaccharide concentration ranging from 0 to 36 mg mL⁻¹, 50 mM sodium acetate buffer at pH 5.0
5 25 °C. Excitation wavelength was 280 nm. We measured the total fluorescence emitted above 320 nm thanks to
6 a band-pass filter. The traces obtained were fitted to a single exponential function of the form $F(t) = m^*t + q$
7 + $\exp(-k_{\text{obs}}^*t)$, where k_{obs} represents the kinetic rate constant observed and $m^*t + q$ represents the plateau
8 signal, which slightly decreases with time due to photobleaching. We then plotted k_{obs} Versus the concentration
9 of the polysaccharide.

10 *Dynamic light scattering (DLS) measurements.* Samples were prepared to a final protein concentration of 80
11 μM in 50 mM Sodium Acetate buffer, pH 5, 25 °C. We used cellohexaose (Megazyme, Chicago, IL, USA) at
12 a final concentration of 200 μM as a cellulose substrate. Before measurements, the protein samples were
13 filtered using Anotop filters, with a 20 nm cut-off. (Whatman, Little Chalfont, UK). DLS measurements were
14 performed using a Zetasizer Nano S device from Malvern Instruments (Malvern, Worcestershire, UK)
15 thermostated with a Peltier system in a low-volume 10 × 4 mm disposable cells. The values of refractive index
16 and viscosity set on the instrument were determined using the software provided by the instrument, based on
17 the information of buffer and temperature provided by the user. The size distributions shown in Figure S2
18 represent the average of three consecutive measurements.

19 *Molecular docking.* The docking calculations were performed on PC crystal structure (pdb 2KQA). PC mutants
20 were generated by using UCSF Chimera [23] and the Dunbrack rotamer library [24]. The coordinates of
21 cellohexose were generated by using the GlyCaNS toolbox
22 (<http://haddock.chem.uu.nl/enmr/services/GLYCANS/>). The CNS topologies and parameters needed for the
23 subsequent docking stage were generated by using the software ACPYPE [25]. All the docking calculations
24 were performed by using the HADDOCK 2.2 software [26,27]. The simulations were guided by selecting
25 residues known or inferred to be involved in the interaction with sugars as “active” (i.e. involved in the
26 interaction between PC and the ligand polysaccharide), while the solvent accessible surface neighbours of
27 active residues were selected as “passive”. See Table S1 for a list of active and passive residues. For each
28 docking, 1000 structures were generated for initial rigid docking, and the top scoring 200 complexes underwent
29 further flexible minimization/refinement and then energy minimization in explicit water. Docked structures
30 were then clustered according to the fraction of common contacts (FCC) algorithm [28] with a 0.75 FCC cut-
31 off. For each docking, the best structure was selected considering the lowest scoring structure of the most
32 populated cluster.

33 The cellohexose binding affinity for each docking calculation was calculated by using the K_{DEEP} server [29].
34 Structural analysis and molecular graphics were performed by using the software UCSF Chimera [23].
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53 **2.3 Mutant activity on cellulose**

1 *Filter paper weakening.* The ability of wtCP and mutants to weaken filter paper was examined on Whatman
2 no. 1 filter paper (GE Healthcare, Chicago, IL, USA) as previously described [17-19] with some modifications.
3 Filter paper was cut into 0.6 cm diameter discs of about 4 mg each, and two discs were incubated in 1 mL of
4 50 mM sodium acetate buffer (pH 5.0) containing the proteins at 10 μ M concentration. Buffer alone without
5 proteins was used as negative control. The experiments were performed in 2 mL tubes, sealed with parafilm
6 and incubated at 38°C for 48 h, onto an HLC thermomixer at 750 rpm. At the end of incubation discs were
7 removed and the absorbance at 500 nm was measured. Each measurement was taken immediately after shaking
8 the suspension in the measuring cuvette, and three different measurements were acquired for every suspension.
9
10 *Synergistic activity with cellulase.* In order to assay the synergistic activity of wtCP or mutants and cellulase
11 we treated one disc of no. 1 filter paper as reported above. After 24 h of incubation, about 0.01U of cellulases
12 (Sigma Aldrich C2730, Saint Louis, MO, USA) were added to each tube and 30 μ L of solutions were recovered
13 after 3, 6, 24, 48, 72 and 144 h to perform the reducing sugars quantification through the dinitrosalicylic acid
14 method (DNS). Briefly, 100 μ L of DNS were added to 30 μ L of sample and heated at 95°C for 5 minutes and
15 the absorbance at 540 nm evaluated. A solution of glucose (2 mg mL⁻¹) was used to build a calibration curve.
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25 **2.4 Eliciting activity of the mutants**

26 *Phytoalexin production.* The ability to induce the synthesis of phytoalexin-like molecules from the leaves of
27 the non-host *Arabidopsis thaliana* (Col. 0) was verified as previously described [17-19]. Briefly, five drops of
28 10 μ L each, containing 150 μ M wtCP or mutants were applied on the lower surface of *A. thaliana* leaves and
29 placed into a moist chamber under constant light at 22°C. Controls with water were performed for each sample
30 and, after 24 h of incubation, the droplets were recovered. Fluorescence was recorded with a PerkinElmer LS-
31 55 spectrofluorimeter (Waltham, MA) using excitation and emission wavelengths at 320 nm and 386 nm
32 respectively, with 5nm for both excitation and emission. Results were expressed as fluorescence intensity/drop.
33 Six independent measurements were averaged.
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40 *H₂O₂ production.* *A. thaliana* leaves were treated with wtCP and mutants, and H₂O₂ was visualized *in situ*
41 according to Luti et al., [17]. Briefly, 10 μ L drops of a 150 μ M solution of each protein were applied on leaves
42 and incubated for 24h in a moist chamber at 22°C. After incubation, droplets were removed and H₂O₂
43 production was estimated by adding the specific probe 2'-7' dichlorodihydrofluorescein diacetate (DCFH₂-
44 DA; SigmaAldrich, Saint Louis, MO, USA). Leaves were then incubated in a 20 mM sodium phosphate buffer
45 at pH 6.8 containing 10 μ M DCFH₂-DA, at room temperature for 1 h. After staining, the samples were washed
46 twice with fresh buffer to remove the excess of fluorophore and mounted in buffer on microscopic slides.
47 Green fluorescence was then observed, using an excitation wavelength of 460nm and emission wavelength at
48 512nm, under a confocal Leica TCS SP5 scanning microscope (Leica, Mannheim, Germany).
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57 **3. Results**

58 **3.1 Features of the mutants and rationale**

1 The present study was performed with three CP mutants. D77A, in which aspartate 77 is replaced by alanine,
2 was available from a previous work [17]. Aspartate 77 is crucial for the expansin-like activity and is located
3 in the same position occupied by D82 in expansins [17]. The mutants V63A and N84D were obtained with the
4 present study. Valine 63 is located in the hydrophobic core of the protein and its substitution to alanine (V63A)
5 was expected to destabilize the protein while leaving charge unaltered. The mutant N84D, in which asparagine
6 84 was replaced by aspartic acid was produced with the dual purpose of increasing net charge in the CP region
7 involved in substrate interaction and determining its role in CP expansin-like activity (Figure 1;Table 1).
8 Mutations of V63 and N84 were checked by agarose gel electrophoresis and sequencing, while purity of
9 purified proteins was checked by SDS-PAGE (Fig S1).
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17 **3.2 The mutation V63A induces dramatic structural changes on CP**

18 In order to assess the impact of the mutations on the fold of CP, we investigated the secondary structure of the
19 mutants in comparison with the wild-type CP (wtCP) protein, by means of far-UV CD spectra (Fig. 2A). The
20 results of this analysis revealed that N84D and D77A showed far-UV spectra superimposable to that of the
21 wtCP consisting of two negative bands at ca. 230 and 210 nm. Thus, these variants shared the same secondary
22 structure as the wtCP, confirming that the mutations did not alter the topology of the protein. By contrast,
23 V63A exhibited a different CD spectrum (Fig 2A), with the band at 230 nm less intense and the other peak
24 more evident and skewed to the left. The spectrum indicates that V63A variant possesses a higher fraction of
25 residues populating an unfolded conformation, thus suggesting that the substitution of valine with alanine
26 within the hydrophobic core of the protein affected dramatically the protein fold.
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28 Subsequently, the impact of mutations on the thermal resistance of CP was assessed by DSF, which allowed
29 rapid and sensitive evaluation of the melting temperature (T_m) [30,31] using the fluorescent probe SYPRO
30 Orange. The DSF curves obtained for all the variants revealed a single transition with a T_m of 68.5 ± 0.5 °C (Fig
31 2B) for the wtCP, whereas the mutants V63A, D77A, N84D showed values of 60.5 ± 0.5 °C, 56.5 ± 0.5 °C and
32 67.5 ± 0.5 °C, respectively (Table 2). Thus, the substitution with alanine in position 63 caused a dramatic effect
33 on thermal resistance, confirming that this residue plays a pivotal role in maintaining structure integrity.
34 Intriguingly, the mutation D77A noticeably affected T_m . This result is in accordance with previous results
35 obtained on the thermodynamic stability of this variant [17] and corroborates our CD analysis as this mutation
36 does not alter the protein fold but destabilizes the protein [17]. By contrast, N84D showed a T_m very similar to
37 the value measured for the wtCP, suggesting that the negative charge acquired upon mutation did not
38 destabilize the protein fold.
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40 Thermodynamic stability of our CP variants was determined by means of guanidine hydrochloride-induced
41 denaturation. For wtCP and D77A the experiment had been carried out previously [17] and data were reported
42 in Fig. 2C and Table 2 only for comparison. The N84D variant resulted to be only slightly destabilized, as
43 indicated by the free energy change upon denaturation ($\Delta G_{U-F}^{H_2O}$), the dependence of ΔG_{U-F} on denaturant
44 concentration (m), and the concentration of mid-denaturation (C_m) values which were very similar to the wtCP.
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Conversely, the V63A appeared to be strongly destabilized, with $\Delta G_{U-F}^{\text{H}_2\text{O}}$, m and C_m lower than wtCP (Table 2). Interestingly, V63A, albeit improperly folded, exhibited higher thermal stability but a lower thermodynamic unfolding free energy than D77A. Of notice, this variant showed a strong deviation of the m value in the equilibrium denaturation. This parameter was reported to correlate with the change in accessibility to solvent area upon denaturation (ΔASA), and both m and ΔASA are known to correlate with heat capacity change (ΔC_p) between folded and unfolded state (Myers et al., 1996). Normally, m and ΔASA undergo only minor changes within a set of point mutants of the same protein. The strong difference observed in the m obtained for V63A, together with the asymmetry between heat resistance and thermodynamic stability, may therefore suggest that the folded state of this variant is less compact compared to wtCP and this corroborates our CD analysis.

3.3 Expansin-like activity of CP depends on net charge and not on structural stability

We tested the expansin-like activity of mutants by using the filter paper weakening method (Figure 3A). While the mutation D77A led to a complete loss of expansin-like activity [17], the other mutations characterized here led to unchanged or slightly increased activity. In fact, V63A did not affect differently cellulose fragmentation when compared to wtCP. Indeed, the turbidity values were of 0.50 ± 0.06 and 0.53 ± 0.04 for wtCP and V63A, respectively. Interestingly, the N84D mutant produced more paper fragmentation than wtCP. The increase was already apparent after 6h of incubation and became statistically significant at 24 h when the N84D showed a turbidity value of 0.65 ± 0.08 ($P\leq 0.05$).

The role of these single mutations on the expansin-like activity was also determined by assaying the synergistic effect with cellulase (Figure 3B). While glucose released by cellulase alone at 144h of incubation resulted to be $35 \pm 8 \mu\text{g}$, pre-incubation of paper with wtCP increased glucose production up to $471 \pm 40 \mu\text{g}$. The same experiment was repeated with N84D and V63A, obtaining an amount of glucose released of $587 \pm 49 \mu\text{g}$ and $464 \pm 47 \mu\text{g}$, respectively. In summary, V63A showed similar expansin-like activity to the native protein, despite this variant exhibited major structural modifications, while N84D showed better performances when compared to wtCP ($P\leq 0.05$).

3.4 The eliciting activity is highly sensitive to mutations

CP was reported to induce plant defence responses on *Arabidopsis* leaves [7]. Consequently, we tested the ability of the CP variants to induce the synthesis of phytoalexins (which belong to the class of plant defence secondary metabolites) and ROS (Reactive Oxygen Species, produced by plants upon infection or stress). *Arabidopsis* leaves were treated with drops containing wtCP or mutants at the concentration of $150 \mu\text{M}$. Phytoalexins were assayed on the recovered drops by fluorescence spectroscopy. The production of ROS was detected on leaf tissues by means of confocal microscopy. As expected, wtCP was able to induce the production of phytoalexins with a fluorescence of 24.7 ± 5.0 a.u. By contrast, all the mutants showed a weak induction of phytoalexins, with observed values of 1.9 ± 0.4 , 3.4 ± 0.4 and 2.1 ± 0.7 a.u. for N84D, V63A and D77A, respectively (Figure 4A).

1 We obtained similar results when testing the ability of either CP or its mutants to trigger the production of
2 ROS (Figure 4B). Representative confocal microscopy images revealed the ROS burst induced by wtCP, as
3 indicated by the increase in green fluorescence of 280% compared to control leaves. In contrast, fluorescence
4 emission was completely absent in N84D- and D77A-treated leaves, thus indicating a complete loss of eliciting
5 activity. Interestingly, V63A was able to induce a small yet significant increase in ROS production, as
6 suggested by the increase in fluorescence of 30% compared to control.
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10 11 12 **3.5 Molecular modelling of the interaction between CP and cellulose**

13 We investigated the structural determinants of cellulose interaction with CP in both the wild type and in the
14 N84D and D77A forms by using *in silico* molecular docking computations. The V63A mutant was not taken
15 into consideration in this part of the project because this residue is located in the core of the protein. To mimic
16 cellulose, we made use of a linear hexamer composed of D- β -glucopyranose moieties linked with β (1 \rightarrow 4)
17 bonds (cellohexose). For analysis we used the monomeric form of the protein because, differently from EXLX1
18 expansin-like protein (Georgelis et al., 2012), CP does not dimerize during cellohexose interactions, as
19 demonstrated by DLS experiments (Figure S2). In fact, the apparent hydrodynamic diameter of the protein
20 alone or in combination with cellohexose was the same (4 nm) indicating that, differently from canonical
21 expansin-like proteins, CP remains in monomeric form during polysaccharide interactions.
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23 For the calculations, we applied a knowledge-based docking approach. In this technique, the ligand is guided
24 to the correct binding site by using a set of constraints derived from experimental data or bioinformatics
25 predictions. In the present study, the docking was guided by selecting as “actives” the surface residues mutated
26 in this work (N84 and D77 of the native CP and the respective mutated residues D84 and A77) and other
27 residues inferred to be involved in the CP/cellulose interaction (namely S4, Y5, D6, Y7, Y9, and R101; [16-
28 17]). The docking algorithm rewards the formation of close interactions between the ligand and the set of active
29 residues. A second set of residues (referred to as “passive residues”) consisting of the residues found on the
30 surface of the protein and in close contact with the “active residues” was also included in the calculation. The
31 docking algorithm also considers favourably the presence of passive residues in the protein-ligand interface
32 (see Table S1 for a complete list of the active and passive residues).
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34 The results of the docking computations are reported in Figure 5. In the case of wtCP, the cellohexose bound
35 in a wide cleft on the protein surface in correspondence of the active residues used to guide the calculation
36 (Figure 5A). In particular, D6 and R101 clamp on both sides the polysaccharide chain, stabilizing the
37 orientation of the ligand. We observed the same binding mechanism in the computations involving mutated
38 CP. Furthermore, D77 was directly involved in the binding through a hydrogen bond with one cellohexose's
39 hydroxyl group. In the case of the docking performed with the N84D mutant (Figure 5B), the introduction of
40 a negative charge in the binding region did not considerably change the general binding pose of cellohexose
41 on the CP surface. However, in wtCP, N84 is H-bonded with D6 and did not form a direct interaction with
42 cellohexose, while in the N84D mutant, D84 was directly involved in the interaction with the ligand. Finally,
43 in the case of the D77A mutant (Figure 5C), the model polysaccharide bound in a slightly different pose. The
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cellohexose appeared to be shifted along the axis of the molecule of about two glucose subunits, even if the general orientation of the ligand was the same with respect to the other variants. The replacement of a negatively charged residue with a hydrophobic residue of smaller size resulted in the lack of the D77-cellohexose H-bond observed in the previous docking.

The binding affinity of cellohexose to CP was calculated by using a machine-learning approach for predicting binding affinities using 3D-convolutional neural networks implemented in the K_{DEEP} server [29] and the results are reported in Table 3. The results obtained indicate that the introduction of a negatively charged residue (N84D) in the region binding cellulose slightly increased the binding affinity through the formation of a direct hydrogen bond with the substrate. In contrast, the elimination of a negatively charged residue (D77A) resulted to some extent into a reduction of the binding affinity, possibly caused by the loss of one hydrogen bond.

3.6 The interaction between CP and cellulose analogues recapitulates expansin-like activity

In order to further shed light on the interactions between CP and cellulose, we exploited carboxymethyl cellulose (CMC), as it shows higher solubility when compared to cellohexose and it has been successfully employed in this type of experiments [15,32].

The interactions between ligands and proteins result commonly into an increase in the ability of the latter species to resist denaturation, be it thermal or chemical. This effect can be exploited to detect binding, by relating the observed denaturation shift to affinity. Thus, we attempted to detect binding by measuring thermodynamic stability and thermal resistance in the absence and presence of polysaccharides. The results indicate that, in both cases, the differences observed were not enough to detect binding (Fig. 6A and 6B). In fact, $\Delta G_{U-F}^{\text{H}_2\text{O}}$ obtained was equal to $36.8 \pm 1.5 \text{ KJ mol}^{-1}$ in the presence of 36 mg mL^{-1} CMC; this value is within experimental error from the value obtained in the absence of polysaccharides, $35.2 \pm 0.5 \text{ KJ mol}^{-1}$.

We obtained similar results also when gauging thermal resistance. The T_m in the presence of CMC (69.0 ± 0.5 °C) resulted not significantly higher than the T_m obtained in the absence of the molecule (68.5 ± 0.5 °C).

Lastly, we attempted to detect binding by following unfolding in real time. The rationale behind this approach is that chemical denaturation might become slower if the release of the ligand from the folded state becomes the rate-limiting step. Here, we followed tryptophan fluorescence in real time during unfolding of CP, triggered by a rapid dilution of the protein into solutions containing GndHCl. We then repeated the experiment in the presence of CMC (Figure 6C) and measured the unfolding rate constant as a function of the CMC concentration (Figure 6D). Figure 6C shows two representative traces, normalized to the fraction of folded protein, obtained in the absence and presence of 36 mg mL^{-1} CMC. This figure illustrates that the presence of CMC renders CP unfolding slower. The unfolding rates of CP are 0.190 ± 0.019 and $0.095 \pm 0.019 \text{ s}^{-1}$ in the absence and presence of 36 mg mL^{-1} CMC, respectively. Figure 6D shows the ratio between the rate constant (k_{obs}) observed at a given [CMC] and the value in the absence of CMC (k_0). The results of this experiment (Fig. 6D) confirm that unfolding of CP becomes slower upon increasing concentrations of CMC. This effect becomes visible at concentrations higher than 10 mg mL^{-1} and reaches a plateau after 20 mg mL^{-1} , suggesting that after this concentration CP binding sites become saturated. We analysed data with an arbitrary sigmoidal function,

1 obtaining a concentration of half binding $C_{1/2}$ of $18.1 \pm 0.5 \text{ mg mL}^{-1}$. Although this trend indicates an interaction
2 between CP and CMC, CMC solutions are dense and viscous, and this may slow down unfolding.
3 Consequently, we repeated the same experiment with Ficoll 400 (36 mg mL^{-1}). The decrease of the unfolding
4 rate was much less marked in the presence of Ficoll, thus corroborating our idea that the effect observed with
5 CMC was due to a specific interaction, rather than to a solvent effect. We then repeated the experiment with
6 all three mutants. Consistently, D77A showed lower affinity for CMC than wtCP. The ability to slow down
7 unfolding became visible at concentrations higher than 20 mg mL^{-1} and did not reach a plateau in our
8 experimental setting. While fitting the data to a sigmoidal curve became unreliable in the absence of a plateau,
9 an estimate of $C_{1/2}$ yielded a value $> 30 \text{ mg mL}^{-1}$. Intriguingly, N84D showed higher affinity for CMC than
10 wtCP, with an estimated $C_{1/2}$ of $12.0 \pm 0.5 \text{ mg mL}^{-1}$. The V63A variant had an affinity for CMC similar to that
11 observed for wtCP, with a $C_{1/2}$ of $19.1 \pm 0.5 \text{ mg mL}^{-1}$. Taken together, these results indicate that the interaction
12 between CMC and CP is very labile and weak, being unable to induce significant thermal/thermodynamic
13 stabilization. However, the interaction can be indirectly detected and correlates with its expansin-like activity.
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23 4. Discussion

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27 Members of the CPF can trigger plant defence responses and generally induce resistance against microbial
28 pathogens [7]. The molecular mechanism of their action is still unknown and a plant receptor for these proteins
29 has not been identified yet. Interestingly, it has been recently reported that ScCP1 from *Sclerotinia*
30 *Sclerotiorum* could interact with the pathogenesis-related protein 1 (PR1) in the plant cell apoplast [33].
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34 Several CPF members described until now are able to interact with polysaccharides, such as chitin and
35 cellulose [34]. These features reinforce the idea of an involvement of CPF in host-pathogen interactions. In
36 fact, CP and other proteins of the family, thanks to their well-established expansin-like activity on cellulose,
37 may drive the hydrolysis of plant cell walls or facilitate the entrance of the hyphae during mechanical
38 penetration [19,35].
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42 In this study, we investigated the relationship between expansin-like activity and eliciting activity of CP by
43 exploiting a small set of rationally designed mutants. Our results indicate, for the first time, a partitioning
44 between the structural features allowing the protein to carry out expansin-like and elicitor activities. To
45 summarize, our results indicate that negative charges on the protein surface dominate the expansin-like
46 activity. Removing these charges strongly inhibits their activity, whereas mutations that destabilize fold and
47 stability of CP leave expansin-like activity unaltered. In contrast, the eliciting activity is generally highly
48 sensitive to mutations.
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54 The study was born from the observation that the D77A mutant of CP lacked both expansin-like and eliciting
55 activities and showed an alteration in conformational stability probably due to the loss of a negative charge
56 [17]. To better understand the structure/function relationship we designed two more mutants, V63A and N84D.
57 Firstly, they were designed because both V63 and N84 are conserved residues among CPF members; secondly,
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1 because of their structural role: valine 63 is in the hydrophobic core of the protein; asparagine 84 lies in the
2 hydrophilic shallow groove probably responsible for the interaction with cellulosic substrates [7,16].

3 The V63A mutant showed a different CD spectrum compared to the wild type protein, and a T_m lower by 8
4 degrees. Furthermore, the equilibrium unfolding experiments confirmed its lower stability, suggesting that the
5 folded state of this variant is less compact compared to the wild type, and this corroborates our CD analysis.
6 By contrast, the N84D mutant showed similar structural properties to wtCP, indicating that this mutation
7 affects neither folding nor stability. Interestingly, the D77A mutant, which is properly folded, was less stable
8 than the V63A mutant. In summary, the three mutants showed different structural and thermodynamic
9 properties: N84D has no effect on structure and stability despite the acquisition of a negatively charged residue;
10 D77A does not alter the structure but strongly destabilizes the protein; V63A causes dramatic effects on
11 folding, thermal resistance and thermodynamic stability.
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13 The biological activity of the mutants was determined both as eliciting activity and ability to loosen cellulose.
14 While all three mutants weakly induced defence responses, both V63A and N84D fully retained their expansin-
15 like activity, indicating that expansin-like and elicitor activities of CP are governed by different structural
16 properties. Overall, the results indicate that the ability to induce defence responses in plants seems to be more
17 susceptible to structural changes in the protein than the cellulose-weakening activity, which in turn seems to
18 be a very strong feature dependent on the net charge of the protein.

19 The cellulose-weakening activity of CP can be measured as release of paper fragments in solution as previously
20 demonstrated [19]. Our results showed for V63A a similar activity to wtCP, supporting previous data which
21 indicated that structural destabilization upon mutation does not alter the activity on cellulose. By contrast, the
22 N84D mutant presented an increased expansin-like activity with respect to wtCP. At the used concentration
23 ($0.01\text{U}\cdot\text{mL}^{-1}$) this mutant showed both enhanced paper fragmentation and improved synergistic effect when
24 compared to all the other variants. Previous results indicating the lack of synergistic effect with cellulases
25 were probably due to the high cellulose load in the test ($0.2\text{-}1.0\text{ U}\cdot\text{mL}^{-1}$) [19]; in fact, higher cellulase
26 concentration hydrolyse filter paper quickly masking the protein contribution. Taken together, our data suggest
27 that the expansin-like activity of CP is not affected by structural modifications, but is influenced by the charge
28 of those residues located in the putative oligosaccharide binding region corresponding to S4, Y5, D6, Y9, D77,
29 N84 and R101. In support of our findings, the loss of function observed in D77A can be attributed to the
30 removal of the carboxyl group, rather than to the structural destabilization caused by the substitution to alanine
31 as further suggested by the molecular modelling simulations. Therefore, these experiments show that expansin-
32 like activity of CP does not depend on the stability or on the structure of the protein, but it rather depends on
33 the net charge.
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35 Like expansins, CPF members do not show any enzymatic activity but some of them can show loosening
36 ability on cellulosic materials [35]. In our previous study, CP appeared to be able to act on cellulose without
37 stable binding, as revealed by SDS-PAGE followed by Coomassie blue staining [19]. The proposed CP-
38 cellulose interaction was thus in accordance with the absence of a D2 binding domain typical of plant and
39 bacterial expansins [19]. In this study, we used CMC, a model substrate to detect binding of expansins and
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1 expansin-like proteins with cellulose [15,32] and analysed tryptophan fluorescence in real time during
2 unfolding of CP in order to detect binding ability. By doing so we were able to detect a weak interaction
3 occurring between CMC and the “active residues” of the hydrophilic groove of CP. In accordance with this
4 result, the suppression of a charge in the D77A mutant led to a decrease in CMC affinity, while the N84D
5 mutant showed a greater affinity for CMC than wtCP.
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8 In conclusion, we may assert that CP weakly binds cellulosic substrates while performing its non-lytic
9 loosening, in accordance with other single domain expansin-like proteins not belonging to CPF such as LOOS1
10 from the basidiomycete *Bjerkandera adusta*. LOOS1 is a DPBB folded protein which possesses expansin-like
11 activity and binding capacity on its unique domain [35], corroborating the idea that this structure may be
12 sufficient to produce expansin-like activity.
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14 Since expansins can weaken cellulosic substrates, their possible application in the production of biofuel, as
15 enhancers of enzymatic hydrolysis, has received great attention in the last decade [36]. Interestingly, we
16 produced the N84D mutant that showed higher activity than wtCP because of an additional negative charge
17 located in the putative region involved in sugar interaction. This mutant is stable and binds cellulose weakly,
18 competing in a bland way with cellulase for accessing the substrate. This variant represents a promising
19 candidate as enhancer of enzymatic hydrolysis of cellulose and an interesting subject for future studies aimed
20 to improve biofuel production methods from cellulosic sources.
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35 **Declaration of competing interest**

36 None
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40 **Author Contribution**

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42 **Simone Luti**: Conceptualization; Data curation; Investigation; Writing - original draft. **Francesco Bemporad**:
43 Investigation; Methodology; Writing, review & editing. **Vivoli Vega M**: Investigation; Validation. **Leri M**:
44 Investigation; Validation. **Musiani F**: Investigation; Formal analysis. **Bacelli I**: Visualization; Writing,
45 review & editing. **Pazzagli L**: Conceptualization; Funding acquisition; Supervision; Writing, review & editing.
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60 Figure captions

1 **Fig.1: CP structure.** Structure of CP (PDB ID: 2KQA) with the residues V63, D77 and N84, which
2 have been mutated and used in this study. All images generated using PyMOL version 1.8.
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6 **Fig. 2: Structural characterisation of CP.** (a) Far-UV CD spectra of 6.25 μM wtCP (black), V63A
7 (red), D77A (pink) and N84D (blue). (b) DSF measurements of wtCP and mutants at 0.133 mg/mL
8 concentration, using Sypro orange as environmentally sensitive dye. The fluorescence was plotted
9 *versus* temperature (melting curve) and the resulting plot converted into its first derivative with wtCP,
10 V63A, D77A and N84D. (c) Equilibrium denaturation experiment of 0.04 mg mL⁻¹ wtCP and mutants
11 in the presence of GndHCl concentrations ranging from 0 to 5.6 M.
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21 **Fig. 3: Expansin-like activity of CP.** (a) Weakening activity of CP and mutants on filter paper
22 measured as paper fragments released in suspension after different time of incubation. Absorbance at
23 500 nm was measured on a Ultraspec 2000 (Pharmacia biotech) spectrophotometer. Data are the mean
24 \pm s.d. of values obtained by two experiments performed in triplicate (* $P \leq 0.05$ respect to wtCP:
25 Statistics: t-test). (b) Synergistic activity of CP and mutants with 0.01U of cellulase measured as
26 reducing sugars production. Data are the mean \pm s.d. of values obtained by three experiments (*
27 $P \leq 0.05$ respect to wtCP: Statistics: t-test).
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34 **Fig. 4: Eliciting activities of CP on *Arabidopsis* leaves.** (a) Phytoalexins production. Fluorescence
35 emitted by 150 μM of wtCP, N84D, V63A and D77A was assayed after 24h of incubation. Data are
36 the mean \pm s.d. of values obtained by three independent experiments (* $P \leq 0.05$ respect to wtCP;
37 Statistics: t-test). (b) ROS production. Representative image of *A. thaliana* leaves treated for 24h with
38 H₂O (as control) or with 150 μM of proteins. H₂O₂ was visualized in situ by the fluorescent probe
39 DCFH2-DA.
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50 **Fig. 5: Results of the docking calculations.** Molecular surface of CP in the case of the calculations
51 made using the wtCP (a) or in the case of the N84D (b) or the D77A (c) mutants. The protein surface
52 has been coloured in cyan and yellow for the active and passive residues used to guide the docking
53 calculations, respectively. Position 77 and 84 have been coloured in dark green and orange,
54 respectively. Protein's and cellobiose's atoms have been represented as "balls-and-sticks" coloured
55 accordingly to the atom type. CP residues forming a H-bond interaction with cellobiose have been
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1 indicated in each panel (grey labels correspond to residues forming a H-bond interaction through the
2 backbone).

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6 **Fig. 6: CP-cellulose interactions.** (a) Equilibrium denaturation experiment of 0.04 mg mL⁻¹ wtCP
7 and CMC at 36 mg mL⁻¹. (b) DSF measurements of 0.133 mg mL⁻¹ wtCP with or without
8 carboxymethyl cellulose. The fluorescence was plotted *versus* temperature (melting curve) and the
9 resulting plot converted into its first derivative. (c) Stopped flow representative traces, normalized to
10 the fraction of folded protein, obtained in the absence and presence of 36 mg mL⁻¹ CMC. (d) Ratio
11 between the rate constant (k_{obs}) observed at a given [CMC] and the value in the absence of CMC (k_0)
12 for wtCP and variants.
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23 Supplemental figure captions

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27 **Fig. S1: Proteins purification.** 4-20% SDS-PAGE of the pure proteins. Lane St: Prestained Protein
28 All Blue marker (Bio-rad); Lane 1: wtCP; Lane 2: N84D; Lane 3: V63A; Lane 4: D77A. 5 μ g of each
29 protein were applied; gel was stained with Coomassie Brilliant Blue.
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36 **Fig. S2: DLS experiments.** Particle size distributions of 80 μ g mL⁻¹ wtCP alone (solid line) or with
37 200 μ g mL⁻¹ of cellohexaose (dashed line). Experiment was performed in 50 mM Sodium Acetate
38 buffer, pH 5 at 25 °C.
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Table 1. Chosen mutations and expected outcome on protein.

Chosen mutation	Mutation Effect	Expected Outcome
D77A	Carboxyl group removing in key aminoacid	Loss of expansin-like activity
V63A	Destabilization of hydrophobic core of the protein	Altered protein stability
N84D	Increasing the surface negative charge of the protein	Increased expansin-like activity

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Table 2. Thermal resistance and thermodynamic stability properties of proteins

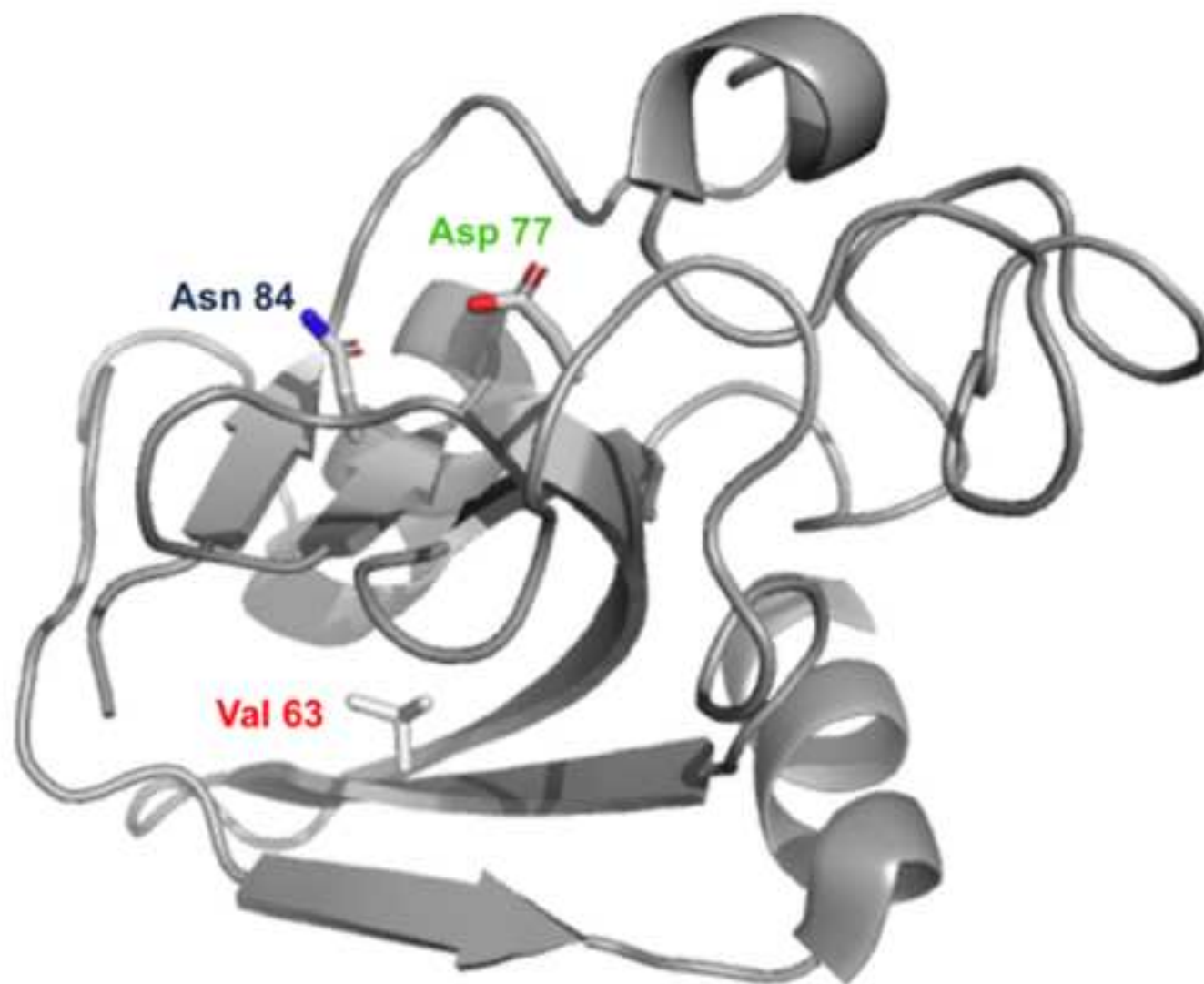
Protein	T _m (°C)	$\Delta G_{U-F}^{H_2O}$ (KJ mol ⁻¹)	<i>m</i> (KJ mol ⁻¹ M ⁻¹)	C _m (M)
wild type	68.5±0.5	35.2±0.5*	14.7±0.3*	2.4±0.2*
V63A	60.5±0.5	14.3±0.5	7.4±0.3	1.9±0.2
D77A	56.5±0.5	21.2 ± 0.4*	14.7 ± 0.3*	1.4 ± 0.2*
N84D	67.5±0.5	32.7±0.5	14.7±0.3	2.2±0.2

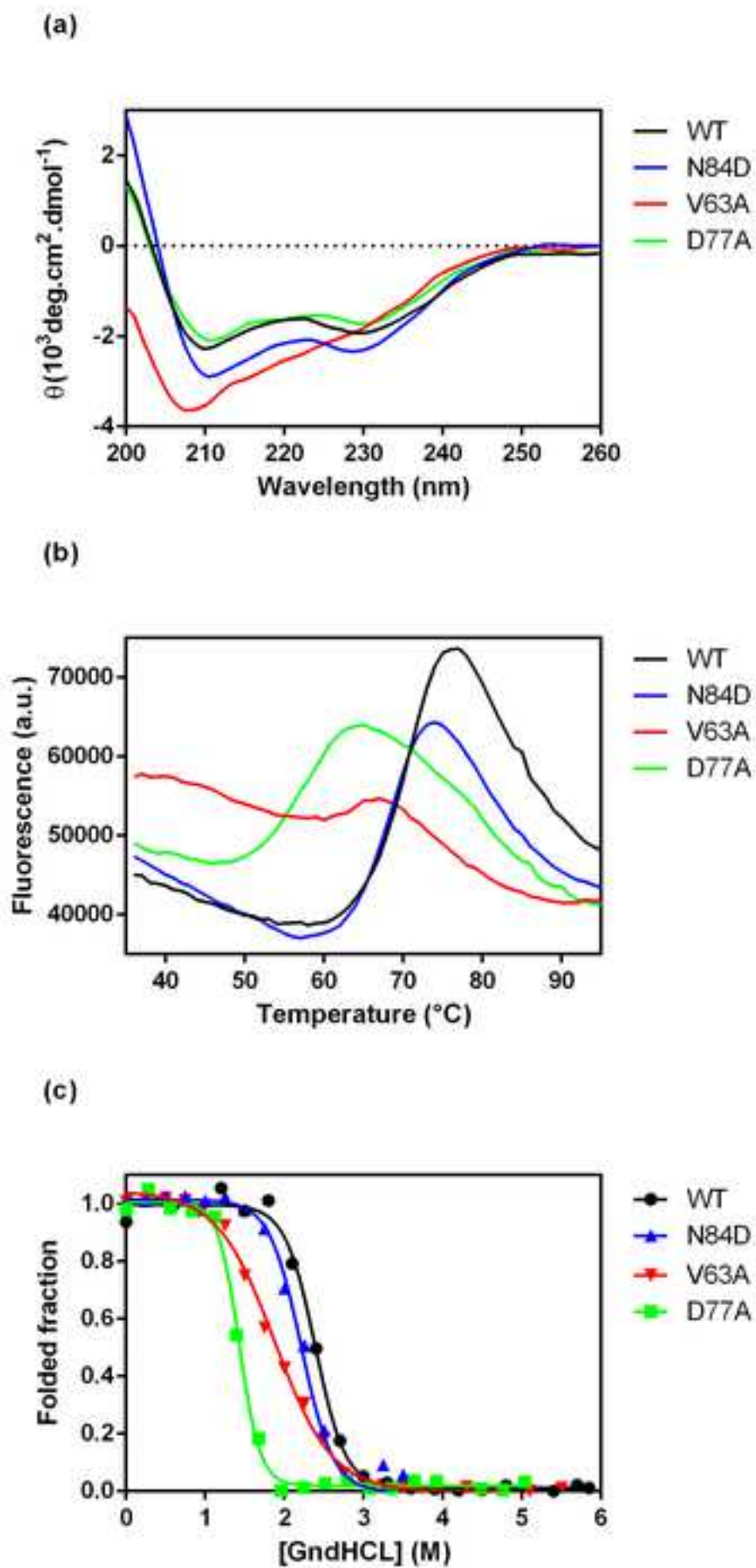
* data reported from Luti et al., 2017.

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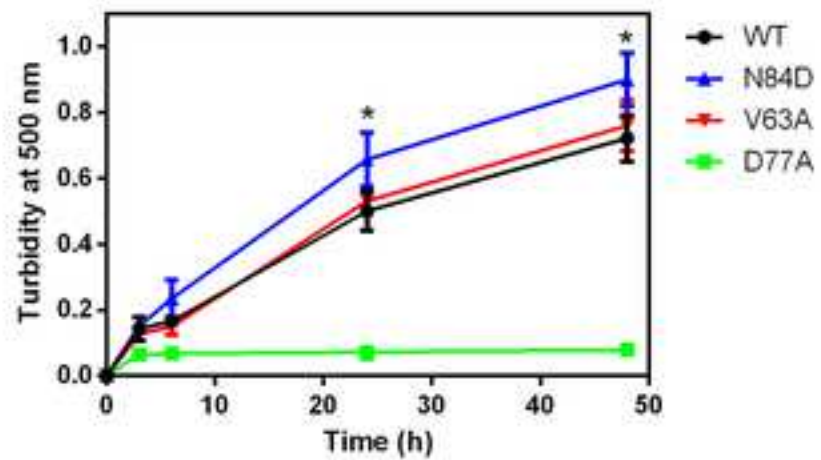
Table 3. Binding affinities calculated through the K_{DEEP} server.

Protein	pK_d	ΔG (kcal mol⁻¹)
wt	5.78	-7.81
N84D	6.03	-8.15
D77A	5.31	-7.16

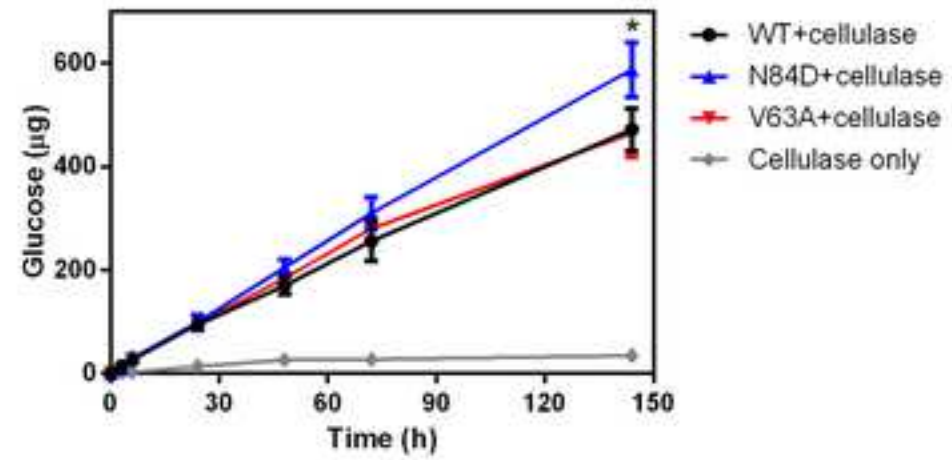




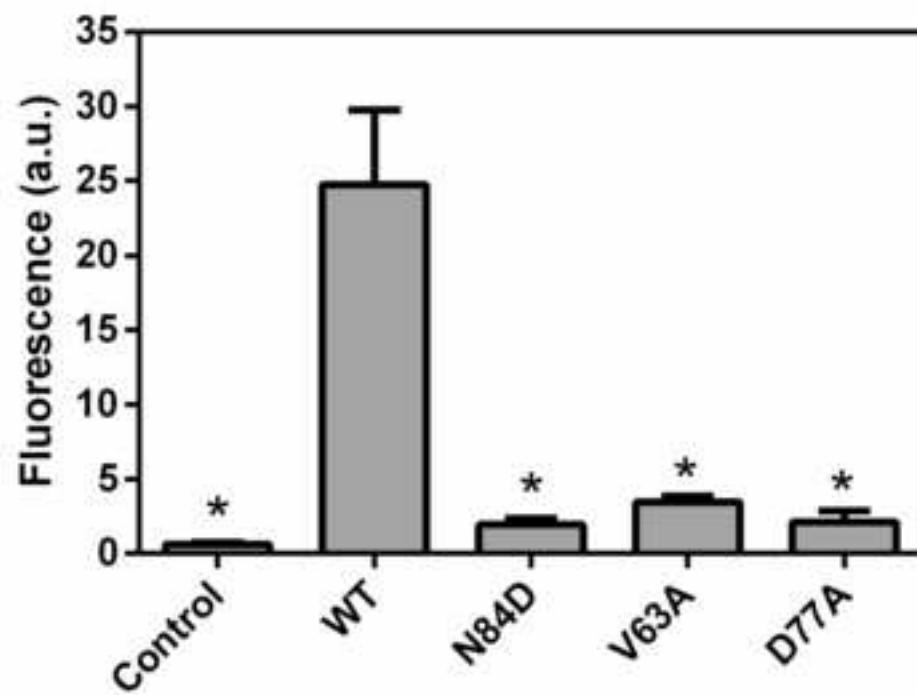
(a)



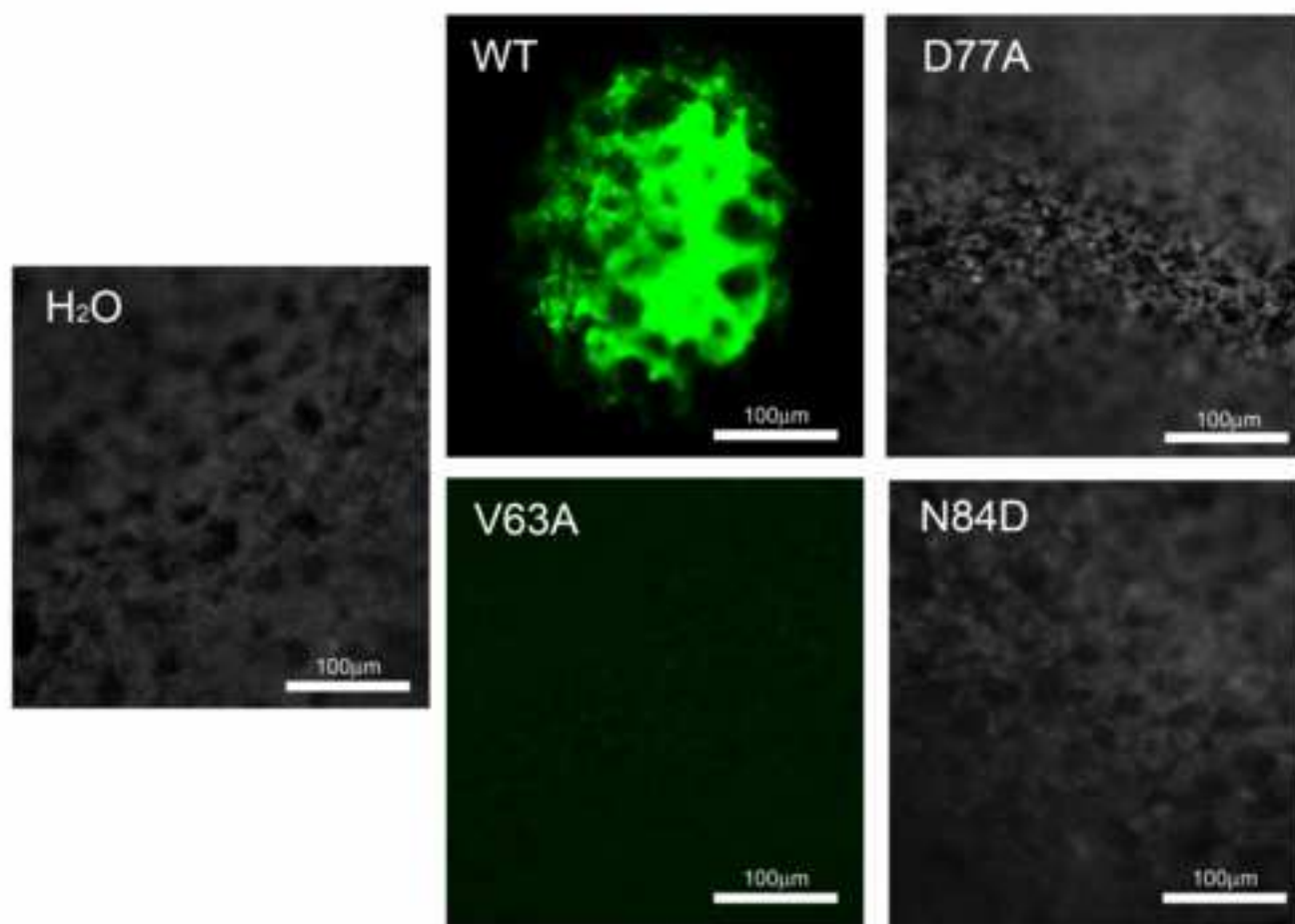
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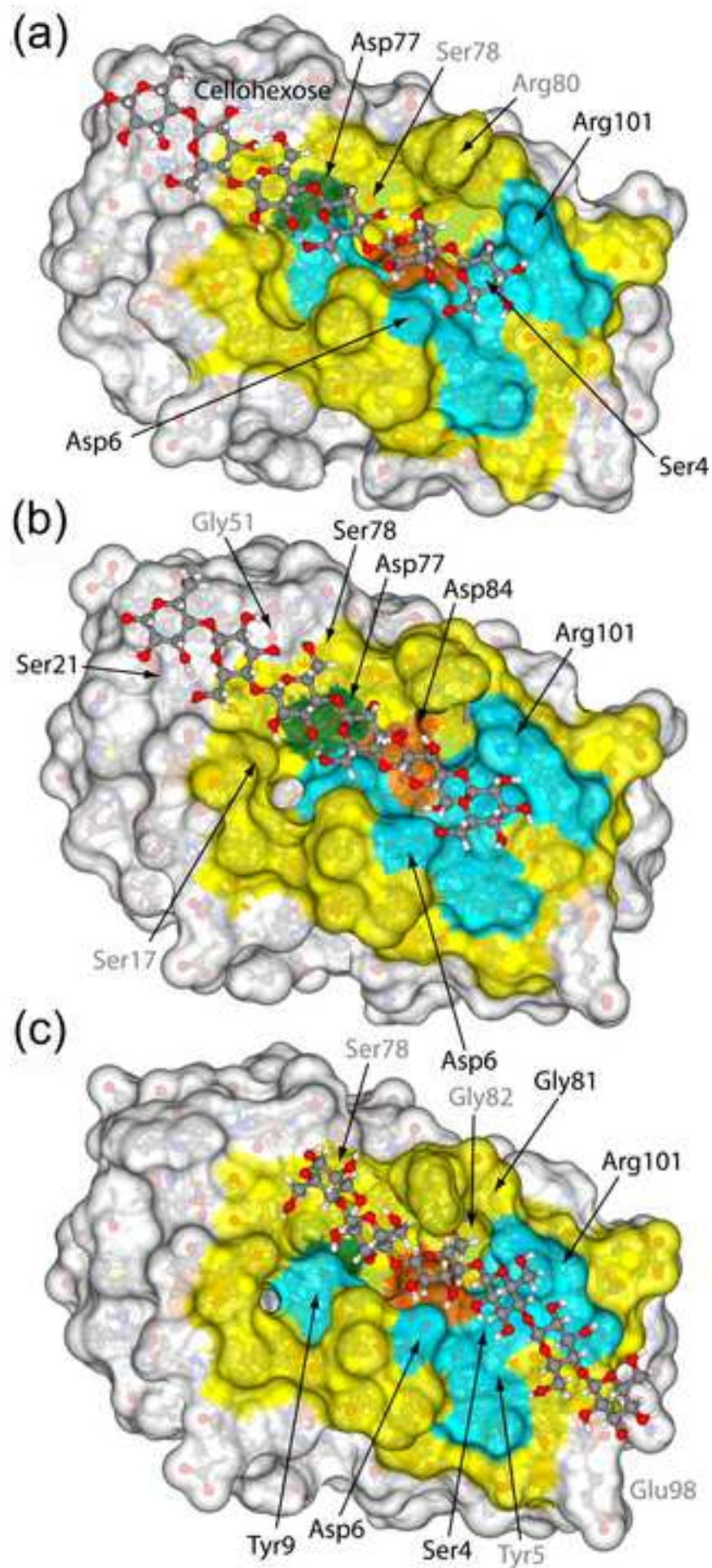


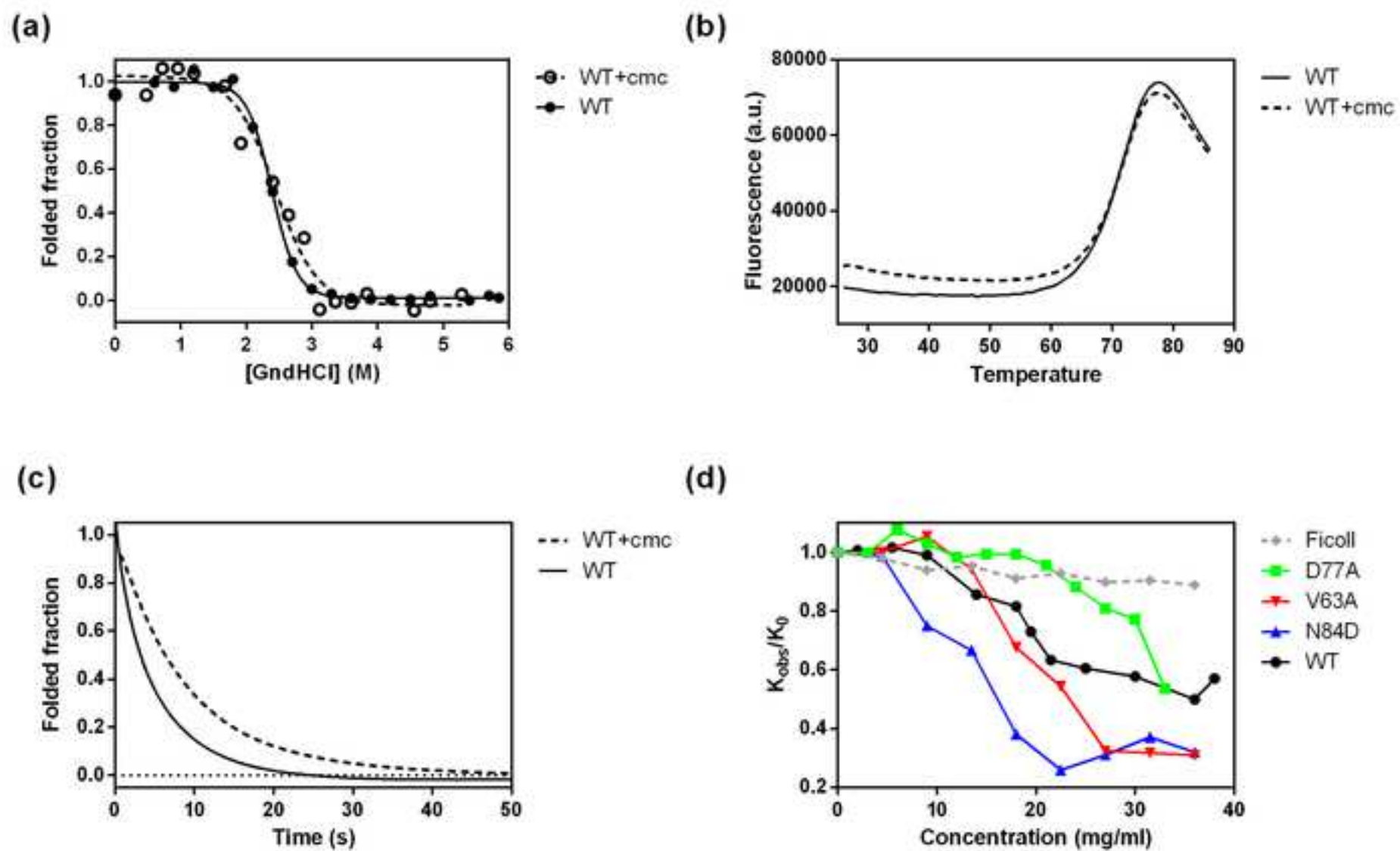
(a)



(b)







Author Statement

Simone Luti: Conceptualization; Data curation; Investigation; Writing - original draft. **Francesco Bemporad:** Investigation; Methodology; Writing, review & editing. **Vivoli Vega M:** Investigation; Validation. **Leri M:** Investigation; Validation. **Musiani F:** Investigation; Formal analysis. **Bacelli I:** Visualization; Writing, review & editing. **Pazzagli L:** Conceptualization; Funding acquisition; Supervision; Writing, review & editing.