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Amphipathic helical peptides hamper protein-protein interactions of the intrinsically disordered chromatin nuclear protein 1 (NUPR1)

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Running title: Peptide binding to an IDP

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Abbreviations: ANS, 8-anilino-1-naphthalene sulfonic acid; CD, Circular dichroism; C-RING1B, C-terminal region of the Polycomb RING protein 1; IDP, intrinsically disordered protein; ITC, isothermal titration calorimetry; MD, molecular dynamics; MSL1, male specific lethal protein; NUPR1, nuclear protein 1; NMR, nuclear magnetic resonance; PDAC, pancreatic ductal adenocarcinoma; PLA, protein ligation assay; PPI, protein-protein interaction; SAM, sterile alpha motif; SLiM, short lineal motif; TFP, trifluoperazine.

ABSTRACT

Background: NUPR1 is a multifunctional intrinsically disordered protein (IDP) involved, among other functions, in chromatin remodelling, and development of pancreatic ductal adenocarcinoma (PDAC). It interacts with several biomolecules through hydrophobic patches around residues Ala33 and Thr68. The drug trifluoperazine (TFP), which hampers PDAC development in xenografted mice, also binds to those regions. Because of the large size of the hot-spot interface of NUPR1, small molecules could not be adequate to modulate its functions.

Methods: We explored how amphipathic helical-designed peptides were capable of interacting with wild-type NUPR1 and the Thr68Gln mutant, inhibiting the interaction with NUPR1 protein partners. We used *in vitro* biophysical techniques (fluorescence, circular dichroism (CD), nuclear magnetic resonance (NMR) and isothermal titration calorimetry (ITC)), *in silico* studies (docking and molecular dynamics (MD)), and *in cellulo* protein ligation assays (PLAs) to study the interaction.

Results: Peptide dissociation constants towards wild-type NUPR1 were $\sim 3 \mu\text{M}$, whereas no interaction was observed with the Thr68Gln mutant. Peptides interacted with wild-type NUPR1 residues around Ala33 and residues at the C terminus, as shown by NMR. The computational results clarified the main determinants of the interactions, providing a mechanism for the ligand-capture that explains why peptide binding was not observed for Thr68Gln mutant. Finally, the *in cellulo* assays indicated that two out of four peptides inhibited the interaction of NUPR1 with the C-terminal region of the Polycomb RING protein 1 (C-RING1B).

Conclusions: Designed peptides can be used as lead compounds to inhibit NUPR1 interactions.

General significance: Peptides may be exploited as drugs to target IDPs.

Keywords: Cancer, calorimetry, drug design, molecular dynamics, NMR, peptides.

1. INTRODUCTION

Protein-protein interactions (PPIs) are critical for molecular communications and processes such as cell division, transcription, signal transduction and programmed cell death. PPIs are also key in the development of various diseases and pathological conditions, such as neurodegeneration and cancer (1,2). PPIs can occur between two structured proteins (or domains); two IDPs; a well-folded protein and an IDP; a folded domain and a short peptide; or between two peptides (3). In those interactions, the secondary structural element α -helix appears in 62 % of all interfaces (4,5). Therefore, drugs inhibiting specific PPIs involving α -helices could have an important therapeutic potential. However, the use of small molecules targeting PPIs is difficult due to two reasons. First, interface regions in PPIs are usually large (700-3000 Å²) (6), with many polar and hydrophobic interactions. And second, the interfaces are typically shallow, with lack of a clear pocket to allow binding of a small molecule (7). Thus, the chemical space of small molecules deviates from that of PPI inhibitors, perhaps explaining the low hit rates of small drugs in PPI inhibition (7,8). Peptides and peptidomimetics (i.e., modified peptides) can be used to target PPIs; in fact, 15-40 % of all possible PPIs involve a linear peptide (9,10). Peptides can have several advantages against small molecules in targeting a protein-protein interface; the most important one is that they can be rationally designed on the basis of the protein sequences of their binding partners. Peptidomimetics further combine the folding properties of peptides, with the lack of proteolytic and metabolic degradation (11,12). In a structure-based approach to drug-design, peptide binding regions derived from protein-protein interfaces can be used as a starting point for the design of PPI inhibitors. However, if the aim is to target an isolated IDP, which is basically an ensemble of rapidly inter-converting conformational states, the design of peptides or peptidomimetics capable of hampering its biomolecular interactions is even more challenging.

NUPR1 is an 82-residue-long (8 kDa), highly basic, monomeric IDP that is over-expressed during the acute phase of pancreatitis (13). It does not have a stable secondary and tertiary structure

1 in any region of its sequence, as occurs in most of other IDPs (14-16). NUPR1 is believed to function
2 during transcription, and it is an essential element in the stress cell response and cell-cycle
3 regulation, but its exact functions are not fully understood (17,18). It plays key roles in pancreatic
4 tumorigenesis, acting downstream of the Kras^{G12D} oncogene mutation, which is critical for pancreatic
5 carcinogenesis (19). Furthermore, NUPR1 is involved in apoptosis by forming a complex with
6 prothymosin α (20), another IDP, as well as implicated in DNA binding (21) and repair (22), through
7 interaction with the male specific lethal protein (MSL1), and in the association to proteins of the
8 Polycomb group (23). All these interactions provide the rationale to consider NUPR1 inhibition as a
9 clinically feasible strategy to expand the PDAC drug arsenal.
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12 We have shown that binding to all those biomolecules occurs through two distant regions of
13 the NUPR1 sequence (20, 22-24): (i) the hydrophobic patch around Ala33, containing the two
14 aromatic rings of the protein, Tyr30 and Tyr36; and, (ii) the region around Thr68, which is a
15 polypeptide patch with relatively high hydrophobicity. The Ala33Gln and Thr68Gln mutants have
16 both shown impaired binding to NUPR1 protein partners, therefore pinpointing the importance of
17 those regions (23). As tested in PDAC-derived cell-based assays (24), both regions also intervene in
18 the binding of small molecules capable of inducing cell-growth arrest and senescence, reduced cell
19 migration, and decreased chemoresistance, thus mimicking NUPR1-deficiency. In fact, one of those
20 drugs, TFP, completely abolishes tumor development *in vivo* on xenografted PDAC-derived cells in
21 mice (24). Therefore, the same hot-spot NUPR1 regions involved in binding to its natural
22 biomolecular partners are also implicated in the binding to small compounds with potential
23 therapeutic action. Given the importance of α -helices in PPIs (4,5), in this work, we investigated the
24 use of synthetic peptides, with an improved tendency to form amphipathic α -helices in solution, to
25 target the same hydrophobic hot-spot regions of NUPR1, and acting as inhibitors of the PPIs of
26 NUPR1. Bioactive peptides of different origins have shown anti-cancer properties ((25) and
27 references therein), and the possibility of using them in the treatment of tumors, targeting IDPs, is
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1 intriguing. It is important to pinpoint that this is the first report of the design of peptides that can bind
2 NUPR1. Therefore, we benchmarked the use of helical-designed peptides to target an IDP and
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4 hamper its protein-protein interactions, and we studied the peptide effects against NUPR1 *in vitro*, *in*
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6 *silico*, and *in cellulo*.
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9 To test this hypothesis, we carried out an *in vitro* characterization of the binding between a
10 series of peptides derived from the wild-type sequence Ac-VKNWMTETLLVQ-NH₂. This sequence,
11 which was originally obtained from the capsid protein of HIV-1, is highly amphipathic, having an
12 intrinsic tendency to adopt helical conformations and to self-associate (26). Three other peptides
13 have been designed *in silico* to have a better helicity than the wild-type sequence (27,28), without
14 altering their amphipathic features (26). The four peptides were assayed through spectroscopic
15 (namely, fluorescence, CD and NMR) and calorimetric (ITC) techniques against NUPR1. In
16 addition, *in silico* studies were carried out to reveal the key determinants in their binding to NUPR1.
17 Finally, *in cellulo* assays were carried out by means of PLAs. Our hypothesis-driven experiments
18 showed that there was binding between wild-type NUPR1 and the helical-designed peptides both *in*
19 *vitro* and *in cellulo*, with a dissociation constant of ~ 3 μM. The peptide binding region of NUPR1,
20 as mapped by NMR spectra, involved: (i) the hydrophobic patch at the 30s region of the protein; and,
21 (ii) other residues located at both protein termini. The ITC control experiments with the NUPR1
22 Thr68Gln mutant further suggested that this residue was important for peptide binding. Blind *in*
23 *silico* studies showed that binding occurred through the same NUPR1 hot-spot region detected in the
24 NMR studies, providing an anchoring to a specific region of the peptide surface, and suggesting a
25 rationale for the experimental lack of binding of the Thr68Gln mutant. The PLAs indicated that the
26 interactions of wild-type NUPR1 with C-RING1B were completely abolished *in cellulo* in the
27 presence of two out of the four peptides; the activity of the other two could not be assessed, likely
28 due to cytotoxicity or peptide inability to penetrate the cells. Our findings suggest that NUPR1
29 interactions with its natural partners can be hampered by the use of designed amphipathic α-helical
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peptides, based exclusively on physico-chemical grounds (amphipathy and helicity). Moreover, our results represent a proof-of-concept that peptides can be rationally designed against IDPs to inhibit their PPIs.

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2. MATERIALS AND METHODS

2.1. Materials

Deuterium oxide was obtained from Apollo Scientific (UK). Sodium trimethylsilyl [2,2,3,3-²H₄] propionate (TSP), imidazole, Trizma base, ANS, deuterated acetic acid and its sodium salt, and nickel resin were from Sigma-Aldrich (Spain). Dialysis tubing, with a molecular weight cut-off of 3500 Da, was from Spectrapor (Spectrum Laboratories, Japan). Amicon centrifugal devices with a cut-off molecular weight of 3000 Da were from Millipore. Standard suppliers were used for all other chemicals. Water was deionized and purified on a Millipore system.

2.2. Proteins expression and purification

Wild-type NUPR1 and the Thr68Gln mutant were produced and purified in Luria-Bertani (LB) media as described (21-23) in C41 cells (29). For ¹⁵N-labelled wild-type NUPR1 the same cell growth protocol as in LB was used, with ¹⁵NH₄Cl (1 gr per litre of media) as the sole source of nitrogen, with M9 medium, a vitamin mix and a mixture of metal ions to improve bacterial growth. The yield of the protein in such minimal medium was roughly 2/3 of that obtained in LB. After purification (either from LB or in minimal media), the protein was extensively dialyzed against water. Given the poor expression of the Thr68Gln mutant in rich media (23), we did not attempt to label it. The mutation at the other hot-spot region of NUPR1, Ala33Gln, resulted in lack of protein expression (23). Protein concentration was determined from the absorbance of the two tyrosines in the amino acid sequence (30).

2.3. Design and synthesis of the peptides

Our initial design for the peptide sequence was Ac-VKNWMTETLLVQ-NH₂. This peptide is a part of the capsid protein of HIV-1 (residues Val181-Gln192 of the whole sequence); in the intact protein, this patch forms an α -helix through which the capsid protein dimerizes (31,32). The predicted percentage of helical structure for this peptide, according to AGADIR (27,28), was 4.6 %, and that obtained experimentally from CD measurements was 6.9 % (26). All the peptides had a

1 tendency to self-associate with dimerization constants in the order of 4 μM (26). The other three
2 peptides (Table 1) were designed, by using AGADIR, with the aim to improve the helicity of the
3 wild-type peptide. The increase of the predicted helicity in the designed peptides was probably due to
4 stabilization through a stacking interaction between the two aromatic rings of the natural Trp4 and
5 the mutated Tyr8 (located on the same side of the helix), and a capping effect at the C terminus of
6 the helix. The experimentally determined helicity was always lower than that predicted. Peptide
7 concentrations were determined from the absorbance of aromatic residues (30).
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10 All the peptides were synthesized by GenScript (New Jersey, USA) with a purity larger than 95
11 %, as tested by chromatography and mass spectrometry.
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14 2.4. Fluorescence

15 Fluorescence spectra were collected on a Cary Varian spectrofluorimeter (Agilent, USA),
16 interfaced with a Peltier, at 25 °C. Experiments were carried out at pH 6.8 (20 mM phosphate
17 buffer). The samples were prepared the day before and left overnight at 5 °C. Before experiments
18 samples were left for 1 h at 25 °C. NUPR1 concentration (either wild-type or Thr68Gln mutant) was
19 15 μM , and the corresponding peptide concentration was 20 μM (in protomer units). For experiments
20 in the presence of ANS, a final concentration of 100 μM of the probe was added; peptide and protein
21 concentrations were the same as in the experiments where the intrinsic fluorescence was monitored.
22 A 1-cm-pathlength quartz cell (Hellma) was used.
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25 (a) *Steady-state spectra (intrinsic and ANS)*- To follow the intrinsic fluorescence of tyrosine
26 and/or tryptophan, samples were excited both at 280 and 295. The slit widths were 5 nm for both the
27 excitation and emission lights. The spectra were recorded between 300 and 400 nm. The signal was
28 acquired for 1 s and the increment of wavelength was set to 1 nm. Blank corrections were made in all
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1 The same experimental set-up was used for experiments in the presence of ANS, but the
2 excitation wavelength was 370 nm and emission light was collected from 400 to 600 nm. Only
3 experiments with the wild-type NUPR1 were carried out with this probe.
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7 (b) *Thermal denaturations*- Thermal denaturations were carried out in samples containing ANS
8 and wild-type NUPR1, by following the fluorescence at 480 nm, after excitation at 370 nm. The scan
9 rate was 60 °C/h, data were collected every 0.2 °C and the average time was 1 s.
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14 (c) *Binding experiments*- For the titration between the peptides and wild-type NUPR1,
15 increasing amounts of NUPR1, in the range 0-10 μM, were added to a solution with a fixed
16 concentration of the corresponding peptide (2 μM, in protomer units). The experimental set-up was
17 that described above, except that excitation wavelength was only 295 nm (to avoid subtraction of the
18 increasing amount of NUPR1 used during the titration); in that way, we were only monitoring the
19 Trp4 of the peptides during the experiment. The experiments were acquired at 25 °C in phosphate
20 buffer, 50 mM (pH 7.4). The samples were prepared the day before and left overnight at 5 °C; before
21 measurements, the samples were incubated 1 hour at 25 °C. The fluorescence values of a blank
22 solution containing only the corresponding peptide were subtracted for each point. The dissociation
23 constant of the peptide/NUPR1 complex, K_D , was calculated by fitting the plot of the observed
24 fluorescence change of Trp4 *versus* added NUPR1 to (33,34):
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$$41 \quad F = F_0 + \frac{\Delta F_{\max}}{2[P-i]_T} \left[([P-i]_T + [NUPR1]_T + K_d) - \left(\left([P-i]_T + [NUPR1]_T + K_d \right)^2 - 4[P-i]_T[NUPR1]_T \right)^{1/2} \right] \quad (1),$$

42 where F is the measured fluorescence at any particular concentration of NUPR1 after subtraction of
43 the blank; ΔF_{\max} is the maximal change in the fluorescence of the P-i peptide when all of NUPR1 is
44 forming the complex compared to the fluorescence of isolated P-i peptide; F_0 is the fluorescence
45 intensity when no NUPR1 was added; $[P-i]_T$ is the constant, total concentration of the corresponding
46 peptide (2 μM, in protomer units); and $[NUPR1]_T$ is that of NUPR1, which is varied during the
47 titration. The titration with each peptide was repeated twice. At all used concentrations, the
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absorbance of NUPR1 was kept lower than 0.2 units of absorbance (at 280 nm) to avoid inner-filter effects, during fluorescence excitation (35).

2.5. CD spectroscopy

CD spectra were collected on a Jasco J815 spectropolarimeter (Jasco, Japan) fitted with a thermostated cell holder, and interfaced with a Peltier unit. The instrument was periodically calibrated with (+) 10-camphorsulphonic acid. The wild-type NUPR1 and the peptide concentrations were 15 μ M and 20 μ M (in protomer units), respectively for the experiments of complex formation.

(a) *Steady-state spectra*: Experiments were acquired at pH 6.8 (20 mM phosphate buffer) with a scan speed of 50 nm/min, a response time of 2 s, a band width of 1 nm, and averaged over six scans at 25 °C. All spectra were corrected by subtracting the corresponding baseline. Samples were prepared the day before and left overnight at 5 °C.

(b) *Thermal denaturations*- Thermal denaturations were carried out with a band width of 1 nm, a response of 8 s and a scan rate of 60 °C/h. Data were collected every 0.2 °C, following the raw ellipticity at 222 nm.

2.6. NMR spectroscopy

The NMR experiments were acquired on a Bruker Avance DRX-500 spectrometer equipped with a triple resonance probe and *z*-pulse field gradients. Experiments were acquired at 25 °C and pH 4.5 (acetate buffer, 50 mM), by adding the corresponding amount (50 μ L) of a stock solution of 0.5 M deuterated acid in D₂O; probe temperature was calibrated with a methanol NMR standard (36). We carried out the experiments at that pH to allow for the detection of the largest possible number of cross-peaks, since at pH 7.0 only 10 residues are observed (23). The pH of the samples was measured before and after the experiments with an ultra-thin electrode (Sigma-Aldrich); for all the peptides during the titrations the pH at both measurements was 4.5 ± 0.2 .

The peaks in the 2D ¹H,¹⁵N-HSQC (37) NMR spectra of NUPR1 were identified by using previously determined assignments at pH 4.5 (22). The HSQC was acquired by using sensitivity

1 enhancement with echo-antiecho-TPPI and gradient selection (36), with the carrier proton frequency
2 at 8.00 ppm, and a spectral width in the ^1H dimension of 6.00 ppm. With this spectral width and the
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4 phase cycling used in the corresponding sequence from the Bruker library (hsqcetgp) the water signal
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6 was not observed. A concentration of wild-type ^{15}N -labelled NUPR1 of 190 μM was used, and the
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8 corresponding amount of peptide (in the concentration range from 200 to 400 μM , in protomer units)
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10 was added. Experiments to determine which residues were first affected upon addition of the
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12 corresponding peptide were carried out at specific peptide concentrations (indicated in Table 2). 2D
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14 $^1\text{H}, ^{15}\text{N}$ -HSQC spectra were also acquired at 115 μM of NUPR1 with increasing concentrations of
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16 each of the peptides in the range 7 to 50 μM (in protomer units). In this case, the spectra were
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18 acquired with the TPPI method (36) with the carrier proton frequency at the water frequency, which
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20 was removed by presaturation. In these titration experiments, a fresh sample of ^{15}N -labelled NUPR1
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22 was prepared for each peptide concentration.
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29 The spectra at each particular peptide concentration were typically acquired with 2 K complex
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31 points in the ^1H dimension, 128 (or 200 for the titration experiments) complex points in the ^{15}N
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33 dimension, with 64 scans per increment (or 128 scans for the titration experiments). Typical spectral
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35 widths for all experiments were 6000 (^1H) and 1500 (^{15}N) Hz. The resulting matrix of each
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37 experiment was zero-filled to double the number of original points in each dimension and shifted
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39 squared sine-bell apodization functions were applied, prior to Fourier transformation. NMR data
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41 were processed and analyzed using TopSpin 2.1 (Bruker). Signal intensities in the NMR experiments
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43 in the presence of a fixed amount of the peptides were measured by using the same program, by
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45 comparison with the cross-peak intensity of the C-terminal residue, Arg82. For the titration
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47 experiments, intensities were reported as absolute values, after correction for the receiver gain value.
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49 Spectra were referenced with external TSP for ^1H , and for the indirect dimension as described (36).
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54 2.7. ITC

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1 Peptide binding to either wild-type NUPR1 or Thr68Gln mutant was determined with a high
2 sensitivity isothermal titration calorimeter Auto-iTC200 (MicroCal-Malvern Instruments, Malvern
3 UK), with reference power of 10 $\mu\text{cal/s}$, initial delay of 60 s, spacing between injections of 150 s.
4 Protein and peptide concentrations were estimated by using their theoretical absorption extinction
5 coefficients (30).
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11 Protein samples and reference solutions were properly degassed to avoid bubble formation
12 during stirring. Experiments were performed with freshly prepared protein solutions, at 15 $^{\circ}\text{C}$ in 50
13 mM Tris pH 7.4, with 2 % DMSO, to improve peptide solubility. A 20 μM solution of either wild-
14 type or Thr68Gln NUPR1 was titrated into the calorimetric cell with 300-400 μM peptide solution
15 from the injecting syringe. NUPR1 did not show any evidence of aggregation at these concentrations
16 used (nor, at least, up to a concentration of 2 mM, where protein assignment was carried out (22)). A
17 sequence of 19 injections of 2 μL volume was programmed with a stirring speed of 750 rpm. The
18 heat evolved after each ligand injection was obtained from the integral of the calorimetric signal.
19 Control experiments (peptide injected into buffer) were performed under the same experimental
20 conditions. The association constant (K_a) and the enthalpy change (ΔH) of the binding reaction were
21 obtained through non-linear regression of experimental data to a model for a protein with a single
22 ligand binding site. Experiments were performed in replicates and data were analyzed using in-house
23 developed software implemented in Origin 7 (OriginLab, Northampton, MA). We analyzed the
24 titrations including a parameter accounting for the background injection heat effects (which are not
25 much larger than the heat effect due to the interaction), and, therefore, we estimated the dissociation
26 constants and the binding enthalpies with sufficient reliability. The background injection heat effects
27 represent the reference heat in the absence of the interaction along the titration; therefore, the
28 departure from that reference value (in our case, upward or endothermically), whatever the value of
29 the background injection heat effect, reflects the heat effect due to the molecular interaction. Blanks
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or controls injecting titrant into buffer were performed in order to rule out any potential artifacts or the possible effect from titrant self-dissociation.

2.8. *Molecular modeling and dynamics*

The structures of the peptides, including the capping at their termini, were built by using the chemical editors Avogadro (38) and VMD (39). The complete structure of each peptide was refined through MD simulations in water, to obtain a representative ensemble of eight different conformations. The simulation package GROMACS (40) was used in combination with the AMBER ff99SB-ILDN force field (41) and the TIP3P water model (42). For the two charged peptides, P-1 and P-3, a chloride counterion was added to obtain an overall neutral system. The peptides were first equilibrated for 1 ns, and then a production run was carried out for other 2 ns at room temperature and pressure. Simulation conditions, including reference values and coupling times for the thermostat and barostat, modeling of the electrostatics and van der Waals interactions, as well as treatment of atomic distance constraints and periodic boundary conditions, were as described (43,44).

An ensemble of structures of NUPR1, either from wild-type or Thr68Gln mutant, was obtained following a protocol previously reported for IDPs (45, 46), with the protein structure initially built in extended conformation and then collapsed in a long-term run (40 ns) performed in the presence of explicit water molecules. Simulation of peptide/NUPR1 complexes were built starting from unbound NUPR1 with equilibrated gyration radius (2.2 nm), as it was experimentally measured (24). To assess potential artifacts due to overcompaction of the IDP structure, additional control simulations were performed using the TIP4P-D water model (47) which is specifically suited to address such deficiency. Water topology was built in the required GROMACS format by using as a starting model the one of TIP4P/2005 (48) which shares the same geometry, and modifying charges and Lennard-Jones parameters as required (47).

2.9. *Molecular docking*

1 The binding affinity of NUPR1 and each peptide was evaluated by performing a systematic
2 molecular docking. Portions of the NUPR1 sequence containing seven consecutive protein residues
3 were considered, with the rest of the backbone truncated and capped when necessary (by using an
4 acetyl and amide group at the N and C terminus, respectively). The graphical interface AutoDock
5 Tools 1.5.6 (49) was used to prepare the system, and molecular docking was performed by using
6 AutoDock Vina (50) at high exhaustiveness (eight times larger than the default value).
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13 2.10. PLA

14 HeLa cells were seeded in 12-well plates on coverslips and transfected with 2 μ g of DNA
15 (NUPR1-FLAG and RING1B-HA) and 2 μ L of Lipofectamine 3000 Transfection Reagent (Thermo
16 Fisher Scientific) per well. Cells were assayed after 28 h post-transfection. To transport the peptides
17 or FITC-antibody (as control) into HeLa-transfected cells, Pro-JECTTM Protein Transfection Reagent
18 Kit (Thermo Fisher Scientific) was used following manufacturer's recommendations. Briefly,
19 peptide solution (in DMSO) was added to a tube containing a dry film of Pro-JECT reagent and
20 incubated for 5 minutes. Then, Opti-MEM medium was added to bring the final delivery volume to
21 500 μ L. Finally, regular culture media was aspirated from the wells and the delivery mix was
22 transferred to the cells. The cells were incubated for 4 h. After that, cells were washed twice in
23 phosphate buffer solution (PBS), fixed, washed twice again, permeabilized in PBS/0.1% Triton X-
24 100 and saturated with Blocking Solution for 30 min before immune-staining with Duolink[®] by
25 using PLA[®] Technology (Sigma-Aldrich) following the manufacturer's protocol. Slides were
26 processed for *in situ* PLA by using sequentially the Duolink[®] In Situ Detection Reagents Red,
27 Duolink[®] In Situ PLA[®] Probe Anti-Mouse MINUS, and Duolink[®] In Situ PLA[®] Probe Anti-
28 Rabbit PLUS. The following antibodies were used: mouse monoclonal anti-HA (12CA5, Santa Cruz
29 Biotechnology) and rabbit anti-human NUPR1 antibody chemically synthesized (Neosystem, France)
30 following previously described methods (22,23). In these experiments, monitored red fluorescence
31 corresponds to the PLA positive signal, and it indicates that the two molecules belong to the same
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1 protein complex. Blue fluorescence corresponds to nuclei (the so-called DAPI staining). To check
2 the specificity of the PLA signal, negative control experiments omitting one of the primary
3 antibodies were performed. Unspecific binding was optimized by using the proper antibodies and
4 optimizing the transfection times. Protein over-expression during the experiments was used to obtain
5 a clearer and better signal.
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11 Preparations were mounted using Prolong Gold antifade reagent (Invitrogen) and image
12 acquisition was carried out on a Nikon Eclipse 90i fluorescence microscope.
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3. RESULTS

3.1. The design of the peptides

Our initial wild-type sequence for the peptide was Ac-VKNWMTETLLVQ-NH₂. This peptide was derived from the all- α -helical wild-type capsid protein of HIV-1 (residues Val181-Gln192 of the whole sequence) (Table 1); the peptide sequence is highly amphipathic, and it self-associates to favour contact among the different subunits in the viral capsid. Therefore, it could be a good lead peptide to test amphipathy and helicity in the design of peptides capable of targeting NUPR1. The original sequence (hereafter, named P-4) was modified by using AGADIR software (27,28) to improve its helicity; the rationale behind the chosen substitutions of residues has been described previously, and it is briefly described in section 2.3. (26). Experimental measurements on the three derived peptides (the two single mutants P-1 and P-2, and the double mutant P-3), performed by using far-UV CD and measurements of the ellipticity at 222 nm and titration in the presence of organic co-solvents, revealed that the helicity value was lower than that theoretically predicted (26) (Table 1); probably the peptides are populating in aqueous solution several conformations involving helix- or turn-like structures. The peptides have also a self-association tendency with affinities in the order of $\sim 4 \mu\text{M}$ (26). This tendency precluded their NMR characterization in aqueous solution.

3.2. The peptides interacted with wild-type NUPR1 *in vitro*

To test whether the peptides and wild-type NUPR1 interacted *in vitro* we followed a three-part approach. As a control, we used the mutant Thr68Gln, which does not show binding to some of the protein partners of NUPR1 (23). First, we tried fluorescence experiments with the four peptides and with wild-type NUPR1 and the Thr68Gln mutant either by acquiring steady-state spectra (in the absence and in the presence of 8-anilino-1-naphthalene sulfonic acid (ANS)) or by acquiring thermal denaturations of the complexes. We used the steady-state fluorescence spectra to determine whether there were rearrangements in the binding pocket occupied by ANS or, alternatively, in the neighborhood of the sole Trp in the peptides, upon binding of the two biomolecules (NUPR1 only

1 has two tyrosine residues). The use of ANS to monitor binding deserves an explanation, because it
2 usually binds to hydrophobic patches in proteins. That is why, usually, ANS shows low binding (and
3 small fluorescence emission) to a folded protein, and high binding (and large fluorescence emission)
4 to an unfolded one; therefore, it can be used as an extrinsic fluorescent probe to monitor protein
5 unfolding. However, ANS does not always follow that behavior: there are some proteins displaying
6 higher ANS binding (or, at least, larger fluorescence emission) when they have a larger amount of
7 structure, such as prothymosin α , nucleobindin (Calcnuc) and Tim15 (51-53). From our experience,
8 we have also observed ANS binding to IDPs. For example, NS3 protease from hepatitis C virus is
9 partially unstructured in the absence of its zinc metal cofactor; this protein binds ANS when partially
10 unstructured, and then increases ANS binding as it unfolds completely along a temperature
11 denaturation (54-56). The same behavior was observed in NUPR1 when it is bound to small drugs
12 (24). On the other hand, we have also observed that ANS (and other fluorescent probes) sometimes
13 does not work as an extrinsic probe with some folded proteins, contrary to the usual expectation.
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31 Our results show that, when both peptide and NUPR1 were present together in solution, the
32 environment around ANS (Fig. 1 A) and that around Trp4 (Fig. 1 B) of the peptides changed (Fig. 1
33 shows, as a representative example, the data for P-3; data for the rest of the peptides are shown in
34 Fig. S1). As it can be observed, NUPR1 showed variation in the amount of solvent-exposed
35 hydrophobic regions in the presence of the peptides. Experiments with the Thr68Gln mutant and P-3
36 suggest the absence of binding (Fig. S2), following the intrinsic fluorescence of the protein, the
37 peptide and the complex. The finding that there are differences between the spectrum of the intrinsic
38 fluorescence of the complex formed and that obtained from the sum of the spectra of both isolated
39 macromolecules suggests that the environment around Trp4 of the peptides, or that of the two
40 tyrosines of NUPR1, are being affected by binding (Fig. 1 B). A similar conclusion can be drawn
41 from the fluorescence spectrum of ANS: the environment around the probe (and thus solvent-
42 exposed hydrophobic patches) changes in the presence of both molecules (Fig. 1 A).
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2 In addition, we suspected that those protein-peptide interactions, detected by steady-state
3 spectra of intrinsic fluorescence or ANS, may promote some limited, small structural rearrangements
4 in wild-type NUPR1, resulting in a different fluorescence thermal-denaturation pattern compared to
5 that of the unbound protein. To elucidate whether there were changes in the conformational
6 ensemble of wild-type NUPR1 upon peptide binding, we carried out thermal denaturations in the
7 presence of ANS, whose fluorescence changes seem to be larger than those of the intrinsic
8 fluorescence of the peptide/NUPR1 complex; we did not carry out experiments with the peptides and
9 Thr68Gln mutant with ANS because of the absence of binding (Fig. S2). Thermal denaturations in
10 the presence of ANS followed a different pattern for isolated P-3, isolated wild-type NUPR1, and the
11 P-3/NUPR1 complex (Fig. 1 C), but for none of the three solutions a sigmoidal behavior was
12 observed in the unfolding traces. This different behavior for several polypeptide chains or complexes
13 during thermal denaturations, monitored by ANS, has been observed in other proteins. As explained
14 above, ANS (and other fluorescent probes) does not always follow the usual expected behavior when
15 monitoring thermal denaturations of folded proteins by the extrinsic ANS fluorescence (57). From
16 our experience, we have observed ANS binding to IDPs, such as NS3 protease from hepatitis C virus
17 (see above). A similar tendency is observed in NUPR1 (24) when is bound to small drugs. Even if
18 the protein is completely unfolded, it may show some weak ANS binding; on the other hand, the
19 same protein when interacting with a molecule may show a different ANS binding. In addition, both
20 free- and bound- protein may display a different unfolding profile monitored by ANS along the
21 temperature denaturation. This observation can be employed for observing interaction between
22 binding partners, as well as for identifying binding compounds in IDPs or well-folded proteins.
23 Therefore, we could conclude from the fluorescence results that there were interactions between the
24 peptides and wild-type NUPR1, and that these interactions were not present with the Thr68Gln
25 mutant.

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27 The fluorescence changes induced by the peptides in the wild-type NUPR1 spectrum are an
28 indirect piece of evidence of the interaction, as variations in thermal denaturations and steady-state
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1 spectra may be the result of unspecific interactions between the peptides and NUPR1. Then, at a
 2 second stage, we used ITC (58) to determine the dissociation constants, K_{DS} , of the peptides for
 3 NUPR1. The K_{DS} for the peptide/NUPR1 complexes were in the low micromolar range, indicating
 4 that the peptides bound specifically to NUPR1 and they would represent a good starting point for
 5 further affinity optimization to obtain binders with a higher specificity (Table 1, Fig. 2). We further
 6 used ITC to determine the importance of Thr68 on the binding properties of NUPR1. Since we have
 7 already observed that this mutation decreases the affinity of NUPR1 for C-RING1B (23), we decided
 8 to test by ITC whether the peptides were bound to Thr68Gln mutant. No binding was observed with
 9 the mutant protein for any of the peptides under the same conditions explored for wild-type NUPR1
 10 (Fig. S3), and thus, we conclude that Thr68 is a key residue to achieve peptide association.

21 We also measured the K_D of the peptides by using fluorescence; in that experiment we were
 22 certain that upon binding to NUPR1 the environment around Trp4 in the peptides was modified upon
 23 binding to NUPR1. In all cases, the measured affinity constants were similar to those reported by
 24 ITC (Table 1, Fig. S4). These experiments confirm that the binding was specific and that it occurred
 25 around the sole tryptophan of the peptides.

26 As the peptides partially self-associate (26), if they were bound to NUPR1 in monomeric form,
 27 then the dimerization equilibrium of the peptide would be coupled to the binding equilibrium
 28 towards NUPR1. Because the binding to NUPR1 implies binding of the monomeric species or
 29 dissociation of the dimeric species of the peptide, then, the apparent dissociation constant for the
 30 peptide/NUPR1 interaction, K_D^{app} , will be larger than the intrinsic dissociation constant for the
 31 peptide/NUPR1 interaction, K_D (i.e., the dissociation constant if all peptide molecules are in
 32 monomeric form). In particular, the apparent dissociation constant for the peptide/NUPR1 interaction
 33 is given by:

$$K_D^{app} = K_D \left(1 + \frac{2[L]}{K_{sD, pep}} \right) = K_D \left(1 + \frac{2[L_2]}{[L]} \right),$$

1 where $K_{sd,pep}$ is the self-dissociation constant for the peptide monomer-dimer equilibrium
2 ($K_{sd,pep}=[L]^2/[L_2]$). From the above expression, we can conclude that if the peptide self-association
3 equilibrium is displaced towards the monomer ($K_{sd,pep}$ very high and $2[L_2]/[L]\ll 1$), then K_D^{app} is
4 roughly equal to K_D (the observed affinity is similar to the intrinsic affinity for the monomeric
5 peptide). However, if the peptide self-association equilibrium is displaced towards the dimer ($K_{sd,pep}$
6 very low and $2[L_2]/[L]\gg 1$), then K_D^{app} is much larger than K_D (i.e., the observed affinity is much
7 lower than that of the monomeric peptide).
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16 And at a third stage, we carried out far-UV CD experiments of the peptide/NUPR1 complexes.
17 For all the peptides, there were no changes in the addition spectrum (obtained from the sum of those
18 of the two isolated polypeptides) and that of the corresponding peptide/NUPR1 complex (Fig. S5 A),
19 indicating that the conformational ensemble of the peptide and/or wild-type NUPR1 remained the
20 same after binding. Furthermore, the absence of a sigmoidal behavior in the thermal denaturations
21 (Fig. S5 B) suggests that wild-type NUPR1 remained disordered upon peptide binding.
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31 To sum up, the three techniques (fluorescence, ITC and CD) showed that peptides and wild-
32 type NUPR1 interacted *in vitro* with a low affinity constant in the range of 4 μ M. This interaction
33 was hampered in the Thr68Gln mutant.
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38 3.3. The peptide binding mapped to a hydrophobic region located at the 30s of NUPR1 sequence

39 To characterize the molecular bases behind the formation of the peptide/NUPR1 complexes,
40 we sought to determine the NUPR1 regions involved in binding. Since we have previously reported
41 the NMR assignment of all residues of wild-type NUPR1 (22) (BMRB number 19364), we used 2D
42 $^1H, ^{15}N$ -HSQC spectra of wild-type NUPR1 to monitor changes upon peptide addition.
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51 At pH 4.5, upon addition of any of the peptides, the NUPR1 cross-peaks broadened (Fig. 3
52 shows as a representative example the data for P-2, whereas the rest of the HSQC spectra for the
53 other peptides are shown in three panels in Fig. S6); the changes were larger in P-2 (Table 2), for
54 which many of the NUPR1 resonances disappeared. It could be thought that the intrinsic self-
55 association of P-2 could be responsible for this general broadening of most of NUPR1 cross-peaks;
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1 however, the fact that the self-association constants for the four peptides is the same ($\sim 4 \mu\text{M}$, (26))
2 and that the assayed concentration of the peptides was nearly the same (Table 2), suggest that there
3 were other factors affecting peptide binding to NUPR1. In P-1, P-3 and P-4 peptides, only the
4 relative intensity of several cross-peaks were affected, suggesting that the binding was specific (as
5 further pinpointed by ITC and fluorescence). In general, the most affected regions of NUPR1 upon
6 addition of any of the four peptides were: (i) around Ala33, where the two aromatic residues of
7 NUPR1 are located (Tyr30 and Tyr36); and, (ii) the C-terminal region (close to Asn72) (Table 2). In
8 some of the peptides, the relative intensities of residues close to the N terminus of the protein were
9 also affected (Table 2). It is interesting to note that the relative intensity of the cross-peak of Thr68 in
10 the presence of any of the peptides did not show a variation larger than 15 %, whereas it has been
11 shown to change during the binding of NUPR1 to small molecules (24). However, the relative
12 intensities of cross-peaks of nearby residues (Glu63, Asn72 and Ser73) were largely affected upon
13 peptide binding (Table 2).
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31 As there were changes only in the intensities of the cross-peaks (and not in their chemical
32 shifts), the equilibrium-exchange between the peptide- free and bound wild-type NUPR1 must be
33 intermediate-to-slow within the NMR time scale. It could be thought that those variations in the
34 spectral intensity were due to a larger solution viscosity (which may be altered by the presence of the
35 peptides) (59); however, at the maximal total biomolecular concentration used in this work (~ 500
36 μM), we have not observed broadening in the spectra of isolated NUPR1 (22). In fact, NUPR1 does
37 not aggregate up to concentrations of 2 mM, where its NMR assignment has been carried out (22). In
38 addition, it could be suspected that the pH increased during the experiment (or upon addition of the
39 peptide), resulting in signal-broadening due to solvent-exchange. However, the pH measurements
40 before and after spectra acquisition indicate that the pH remained constant during the experiments
41 (see section 2.6). The fact that signal broadening was only observed upon peptide addition suggests
42 that the binding between NUPR1 and the peptides was responsible for that broadening.
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2 Furthermore, the titration experiments carried out by following the HSQC spectra at different
3 peptide concentrations suggest that the decrease in intensity of the cross-peaks of NUPR1 spectra
4 was peptide-dependent (Fig. S7). This decrease was larger for P-2, in agreement with the larger
5 affinity constant determined by ITC (Table 1), the gold-standard in determining thermodynamic
6 parameters of binding reactions (58). Moreover, the fact that the decrease in intensity does not occur
7 at the same pace for all the residues indicate that the binding was not a two-state process, as it could
8 be inferred from the results in Table 2 at a single peptide concentration: different cross-peaks show
9 several changes in their relative intensities. It could be thought that a K_D could be obtained from the
10 fitting of the decrease in absolute intensities of each of the different cross-peaks for each of the
11 peptides; however, a reliable value for a dissociation constant cannot be obtained from the changes
12 in intensities for intermediate-to-slow exchange rates within the NMR time scale, as it has been
13 widely discussed (60).
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28 In conclusion, the NMR experiments further confirm that there was binding between the
29 peptides and wild-type NUPR1, involving, at least at a first stage, mainly residues around the 30s
30 region of the NUPR1 sequence.
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36 *3.4. In silico calculations predicted the regions involved in the peptide binding*

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38 A systematic search was performed to study *in silico* the binding of NUPR1 with the peptides.
39 To overcome the computational limitation in treating large numbers of torsional degrees of freedom
40 of IDPs, molecular docking was carried out using seven-residue-long fragments spanning the whole
41 sequence of wild-type NUPR1, and an ensemble of the structures of the peptides as hosts. Although
42 the scoring obtained provides only an empirical approximation of the association affinity, molecular
43 docking has already proved to be effective with NUPR1 to identify binding to other small
44 compounds (24). Furthermore, an accurate determination of the conformational ensemble of the IDP
45 is not a prerequisite for the findings presented below, because docking is applied considering only
46 isolated fragments of the primary structure of NUPR1.
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Figure S8 shows the binding scores obtained for the segments of NUPR1, calculated as a function of the protein sequence. The energy differences among the four peptides were comparable with those observed in the docking of the fragments with different structures of the same peptide (~0.3 kcal/mol, on average); therefore, all the data for each peptide were combined to increase their statistical significance. The predicted segments of the NUPR1 sequence that contribute the most to the binding correspond to minima in the curve (Fig. S8), and include: the 30s region; the polypeptide patch around Thr54; and the two protein termini (with the lowest values found for Thr8 and Asn72/Ser73, which are close to the N and C terminus, respectively). These regions agree nicely with those observed to experience the larger broadening changes in all peptides by NMR (Table 2, see section 3.3). On the other hand, those values of the theoretical binding energies (Table 1) were slightly less favorable than the experimental ones (-5.5 kcal/mol *versus* -7.2 kcal/mol for P-3). Importantly enough, such predicted regions did not include the polypeptide patch around residue Thr68, which was important for the anchoring of small compounds (24), whereas they still include the other key region of relatively high hydrophobicity around Ala33, which was important for the interaction of NUPR1 with the other binding partners (20,22-24).

Once we had identified the NUPR1 region centered around Ala33 as that contributing most to the binding affinity of the peptides, we sought to use this information to pinpoint the preferred binding region in the latter. To this aim, we mapped the binding poses of the seven-residue fragment centered on Ala33 onto the structure of each of the peptides (Fig. 4). The results clearly indicate that only about half of the helical surface of the peptide was involved in the binding, and this interacting region was common to all of them. Interestingly, the binding patch was on the same side where residue 11 is placed, and regardless whether it is an Arg (as in P-1 and P-3) or a Val residue (P-2 and P-4). More importantly, the binding patch did not overlap with the most hydrophobic region of the helical peptides, suggesting that hydrophathy is not the sole determinant of the peptide/NUPR1 association. This binding patch (formed among others by Val1, Trp4, Met5 and Leu9) is the same involved to form the self-associated helical region in the intact capsid protein of HIV-1; thus, the

1 peptides use the same region to interact with other biomolecules (NUPR1) or with themselves (26).
2 These results support the experimental fluorescence experiments (see section 3.2.), in which the
3 environment around Trp4 in the peptides was affected upon binding to NUPR1.
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7 The findings above are apparently difficult to reconcile with the experimental observation that
8 the Thr68Gln mutant (which disrupts the hydrophobicity pattern in that specific region of the
9 NUPR1 sequence) did not bind to the peptides (Figs. S2 and S3). To investigate this aspect, MD
10 simulations were performed of both wild-type NUPR1 and the Thr68Gln mutant, with peptide P-2,
11 for which the largest changes were observed (Table 2, Fig. 3), placed on the protein surface in
12 correspondence to Ala33. For wild-type NUPR1, we observed a capture mechanism of the peptide
13 (Fig. 5), where P-2 appeared to maintain a stable α -helical conformation at the end of the binding
14 process. The protein had a more compact N-terminal region and an extended C-terminal patch,
15 which possesses a higher amount of positively charged residues. The presence of Thr68 provides a
16 slightly higher hydrophobicity and favors the collapse of the protein C terminus on the helical
17 peptide, allowing the flanking residues Glu63 and Asn72 to approach the ligand surface. Thus, the
18 gradient of hydrophobicity along the NUPR1 sequence plays a crucial role in the binding
19 mechanism, although it is not a main factor in maintaining the anchoring of the peptides.
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38 Although our simulations were performed starting with the hydrodynamic radius of the
39 NUPR1 ensemble close to the experimental value measured by NMR (24, 45) possible artifacts due
40 to overcompaction of an IDP are usually a concern during MD simulations. To exclude a potential
41 influence on the binding mechanism described above, additional control simulations were performed
42 using the TIP4P-D water model (47), which avoids disordered protein ensembles less expanded
43 compared to the experiment. As a result, we observed (Fig. S9) for wild type NUPR1 an interaction
44 with the peptide analogous to the one described in Fig. 5. In particular, Ala33 maintained the binding
45 with the helical peptide while Thr68 favored the approach of the protein C terminus, driving the
46 nearby residues Glu63 and Asn72 towards the ligand surface. Besides being an independent
47 validation under simulation conditions that do not favor the complex formation (because solute-
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1 solvent interactions are increased with the use of the TIP4P-D water model (47)), we point out that
2 this effect does not involve an overcompaction of NUPR1 structure, which would involve a stable
3 interaction of the Thr68 residue with the peptide. In contrast, the mechanism described explains why
4 mutation of Thr68 affects the binding without this residue being listed in a binding patch.
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9 In summary, although the docking procedure could not eventually discriminate among the
10 quantitative differences in the binding of the four peptides (Table 1), it provided an accurate
11 identification of the critical regions for the binding to NUPR1. Furthermore, the MD simulations
12 clarified the molecular details in the peptide association.
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19 3.5. Evidence for the interaction of peptides with NUPR1 within an intracellular environment 20

21 To test whether the interaction between the peptides and wild-type NUPR1 occurred within
22 cells and the peptides were capable of inhibiting the binding of NUPR1 with one of its natural
23 partners, we carried out studies *in cellulo*. Our previous results showed that C-RING1B efficiently
24 and specifically interacted with wild-type NUPR1 (23). Subsequently, we sought to detect whether
25 the peptides interacted with NUPR1 and blocked its interaction with C-RING1B by using the
26 Duolink *in situ* assay, which resolves the binding of proteins that occurs at distances lower than 16
27 Å. We performed controls experiments transfecting a FITC-antibody, as a control peptide included in
28 the kit (Fig. S10), in order to monitor the internalization of the peptides, and the possible effect in the
29 interaction between the two proteins. The control peptide did not change the interaction between the
30 two proteins *in cellulo*.
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45 When interaction between the two proteins takes place, red fluorescent spots appear,
46 corresponding to the positive PLA signal. Our results show that only peptides P-1 and P-3 efficiently
47 interacted with wild-type NUPR1 (Fig. 6) and blocked the interaction between the two proteins,
48 diminishing the red fluorescent spots, corresponding to the PLA signals. Different attempts to
49 replicate this behaviour with P-2 and P-4, by using several transfection times, were always
50 unsuccessful. We believe that the presence of the Arg (and its positive charge) adjacent to the C-
51 terminal residue of the peptides (Table 1) allows cell insertion, as the two peptides with such residue
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2 at this position led to successful PLAs. For the rational design of further peptides inhibiting PPIs of
3 NUPR1, the presence of this residue should be ensured.

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5 Thus, taken together, these results demonstrate that a specific peptide/NUPR1 complex forms
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7 within cultured cells. These findings confirm the biological relevance of our simulations and the *in*
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9 *vitro* biophysical and biochemical experiments.
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4. DISCUSSION

4.1. Peptides as inhibitors of NUPR1 functions

The importance of the network of interactions between proteins (known as interactome) for understanding the functioning of living organisms and the development of diseases is widely acknowledged (7). Despite this, drugs designed to target PPIs are relatively rare (when compared to other designed small molecules aiming to specific polypeptide binding sites). Moreover, drug design is particularly challenging when some of the macromolecules involved in the PPIs are IDPs. NUPR1 exerts several functions by specific interactions with other biomolecules (DNA or proteins). We have recently found a small drug, TFP, capable of binding to NUPR1 and inhibiting its functions, thus mimicking the cellular effects observed in the absence of this protein (24). However, the hot-spot region of NUPR1 is a discontinuous epitope (8) involving two distant polypeptide patches: (i) that at the 30s in the NUPR1 sequence; and, (ii) a region close to Thr68 and the C terminus. TFP shows a proximity to both Ala33 and Thr68 in the NMR experiments and in MD simulations of the complex (24). Mutations at those key positions (Ala33 and Thr68) have resulted in impaired NUPR1 mutants where binding to its natural partners is severely decreased (23).

We and others have designed small molecules against IDPs capable of interfering with their PPIs or nucleic-acid interactions (24, 61-68), and there are a few examples in the literature where peptides are used as inhibitors of PPIs involving IDPs (69-71), or several approaches for folded proteins (see 2 and references therein). In these examples, the peptides are designed based on either similarity with the sequence of the partner protein or, alternatively, fitting with the disordered region(s) of the IDP. In all of them, the protein (or protein region involved in the binding) remained basically disordered, with changes in the conformational ensemble of the protein (64). Furthermore, several α -helix peptidomimetics have also been designed based on the α -helix scaffold of the original protein sequence that the IDP binds to (72). In NUPR1, we have not designed peptides based on the sequence of one of its partners neither on any homologous sequence to NUPR1, but rather we have carried out hypothesis-driven experiments by exploiting physico-chemical principles based on

1 the amphipathy and helicity of a series of peptides with a natural tendency to adopt helical structures.
2 There were poor linear relationships between each of these isolated properties and the values of the
3 measured K_D (Table 1) with linear regression coefficients no larger than 60 %. Thus, it is the
4 interplay of both physico-chemical properties, helicity and amphipathy, which seems to allow the
5 specific binding to NUPR1, and not a single property. However, since we have a rough estimation of
6 the percentage of α -helix- or turn-like structures from the CD measurements, we cannot rule out
7 other factors determining the K_D values, such as the entropic contribution due to the solvent or the
8 shape of the peptides, with their corresponding structured conformation.
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10 4.2. Molecular basis of peptide/NUPR1 complex formation

11 We experimentally observed that the assayed peptides bound *in vitro* to NUPR1 (Table 2, Figs.
12 3 and S6) at the same regions involved in binding to other biomolecules: the 30s region and the
13 neighbourhood of Thr68. In fact, the region around Thr68 was critical for the binding of the peptides,
14 since experiments with the Thr68Gln mutant did not yield any binding (Figs. S2, S3). This residue
15 did not contribute energetically very much to the binding (as suggested by the simulation, Figs. 4 and
16 S8), but it was critical in directing the binding mechanism (Fig. 5). Therefore, we can conclude that
17 NUPR1 uses the same discontinuous epitopes (or “short lineal motifs”, SLiMs), which facilitate
18 multivalent interactions with other molecules (73) to bind to any partner. Some SLiMs in other IDPs
19 have also been shown to function as high promiscuous binders: for instance, the transactivation
20 domain of p53 interacts with multiple proteins, by using the same motif with structurally distinct
21 binding modes upon engagement with its several partners (74). In the case of the peptides, other
22 SLiMs of NUPR1 (together with the regions around Thr68 and Ala33) appeared to be involved in the
23 binding, probably due to the larger size of the peptides compared to that of small drugs (24).
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25 Upon binding to the peptides, NUPR1 remained disordered, as judged by the absence of
26 changes in chemical shifts in most of the cross-peaks in the NMR spectra (Fig. 3 and Fig. S6), and
27 the lack of sigmoidal behaviour in the thermal denaturations measured by CD (Fig. S5) and
28 fluorescence. The cross-peaks of residues, that became broadened, might indicate that the
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1 corresponding amino acids are involved in other conformations where they could form more local
2 structured regions, but these local structures must not be very rigid due to the lack of sigmoidal
3 transitions in thermal denaturations. In the particular case of P-2, since most of the signals became
4 broadened and disappeared (Fig. 3), we could not rule out the presence of more structured
5 conformations, although they cannot be very stable due to: (i) the features of the thermal
6 denaturation curves of the complexes; and, (ii) the absence of large changes in the CD spectrum of
7 the complex when compared to that of isolated NUPR1 (as it happens with P-3, Fig. S5). The fact
8 that the cross-peaks of different residues became broadened before those of other amino acids (Fig.
9 S7, Table 2) suggests also that the binding is not a two-state process, and it must involve several
10 populations of peptides with helix-like conformations, or, alternatively NUPR1 conformations with
11 several local residual structure. We suggest that peptide binding must involve a change of NUPR1
12 conformational ensemble from a random set in the isolated protein to a particular subset of
13 conformers (and that is why only a subset of cross-peaks were broadened in each peptide). This shift
14 must require a reduction in the entropy of the whole system (peptide and protein) caused by steric,
15 hydrophobic or electrostatic effects. The persistently disordered conformation of NUPR1 seems to be
16 a general feature (20, 22-24): whatever molecule (synthetic small molecule, DNA, protein and
17 peptide) binds to the IDP, it is not capable of altering its disordered nature. Furthermore, in all these
18 molecules, the binding equilibrium seems to have an intermediate-to-slow exchange rate within the
19 NMR time-scale (22-24, 60).

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At this stage, we do not know the structure acquired by the peptides upon binding to NUPR1,
but from MD simulations, the structure seems to be α -helical (Figs. 4, 5). The helical propensity of
the peptide was decreased in the control runs performed with the TIP4P-D water model (Fig. S9), but
it is not clear whether this is a cost of increasing the solute-solvent dispersion interactions that
improve the disorder propensity in the IDP (47). Furthermore, based on two pieces of evidence, we
believe that the helical structure acquired by the peptides upon binding, as suggested by the MD
simulations, is weak. First, we did not observe sigmoidal, co-operative thermal denaturations

1 monitored by CD or fluorescence for the complexes (Fig. 1C, Fig. S5 B). And second, the binding of
2 NUPR1 to the peptides is not a two-state process as concluded from the different decrease regimes in
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4 NMR cross-peaks intensities for several residues (Fig. S7); it is likely that the peptides, upon binding
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6 to NUPR1, shift the equilibrium towards more populated helical conformations, but all of those
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8 helical conformations are capable, to a larger or lesser extent, of binding to NUPR1. Finally, it is
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10 important to note that the peptides interacted with NUPR1 through residues located in the proximity
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12 of Trp4 (Fig. S4), as it has been observed when these peptides are bound to other proteins or in their
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14 own self-association (26); thus, peptides also employ the same (hydrophobic) residues to interact
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16 with different molecules.
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21 *4.3. Energetic and thermodynamic determinants of the peptide/NUPR1 interaction*

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24 The measured K_D of NUPR1 for interacting with the peptides (Table 1) was similar to that
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26 observed for MSL1 (~3 μM) (22), for prothymosin α (~6 μM) (20), C-RING1B (~12 μM) (23) and
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28 for different drugs capable of inhibiting the binding between NUPR1 and MSL1 (~5 μM) (24). Such
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30 affinity is relatively small, but it allows a proper regulation of the several pathways where NUPR1 is
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32 involved (18), achieving a high specificity despite a moderate binding energy. Moderate affinities
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34 (i.e., in the range 1-10 μM) are not rare for other IDPs, and they have been measured during the
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36 formation of “fuzzy” complexes, where the partners remain disordered after complex formation (75,
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38 76); these fuzzy complexes facilitate fast binding since no disorder-to-order transition is happening.
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40 Thus, binding between both molecules goes beyond the classical archetypes of conformational
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42 selection or induced fit since both are sampling a range of conformations continuously.
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49 It is important to note that our peptides self-associate in solution (with a self-association
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51 constant of ~4 μM) (26); therefore, peptide binding to NUPR1 would be coupled to peptide
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53 dissociation, and the apparent dissociation constant (K_D , Table 1) for the peptide/NUPR1 complex
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55 will contain an energetic penalty associated with the peptide dissociation event. Thus, the intrinsic
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57 dissociation constant for the peptide/NUPR1 interaction would be lower (i.e. more favorable binding
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1 affinity, see section 3.2.). The effect of the peptide self-association has not been considered
2 quantitatively and explicitly in the interaction experiments (fluorescence and ITC). For that, we
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4 would need to know the peptide self-dissociation constant (in the range of $\sim 4 \mu\text{M}$, as measured by
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6 CD titrations (26)) and the self-dissociation enthalpy. However, from the measured apparent
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8 dissociation constant for the peptide/NUPR1 interaction (Table 1), we can state that the intrinsic
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10 dissociation constant for the peptide/NUPR1 interaction would be somewhat smaller. Other
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12 biomolecules interacting with NUPR1 are either monomeric (prothymosin α (20)) or oligomeric (C-
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14 RING1B (23) and MSL1 (22)), although in the MSL1 case, we do not know its exact self-association
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16 state.
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21 *4.4. Perspectives in the design of helical peptides for inhibition of IDPs*

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24 Although there are several FDA approved peptides for therapeutic use (77) and the future holds
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26 promising for others to be approved (78), in general, the current therapeutic application of peptides is
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28 limited due to low cell penetration and marked propensity to proteolysis and metabolic degradation.
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30 We observed some of these features in our design: only peptides P-1 and P-3 were able to hamper the
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32 interaction of C-RING1B and NUPR1 *in cellulo*. Although we cannot rule out the cytotoxicity of
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34 peptides P-2 and P-4, we suggest that cell-permeation was facilitated in P-1 and P-3 by the presence
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36 of an Arg residue at the second-to-last position in peptide sequence (Table 1). An alternative
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38 explanation could be that this residue provides critical electrostatic interactions in a surface patch
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40 that is key for binding to the 30s region of NUPR1, as pointed out by our docking results. In either
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42 case, future designs of the peptides will probably require a positive charge at this position of the
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44 sequence. At this stage, it is tempting to speculate that P-1 and P-3 might be even particularly
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46 effective in conjunction with TFP to prevent PPIs of NUPR1.
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53 Furthermore, our studies suggest that helicity was not the sole factor governing peptide affinity
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55 of the peptides, since there was not a clear relationship between the helicity experimentally measured
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57 (or the value theoretically calculated) and the measured affinities (Table 1). Although we cannot rule
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1 out that peptide self-association could mask a clear relationship between those physico-chemical
2 parameters, it seems that there are other additional factors governing the peptide binding to NUPR1.
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4 In fact, our *in silico* results pinpoint a specific peptide region (that marked in Fig. 4) that may be
5 improved through a rational design, to modulate the binding to NUPR1. However, a delicate balance
6 in the sequence should be maintained to preserve the amphipathic properties of the peptides, since
7 the presence of a hydrophobic patch on their surface appears to be equally important to drive the
8 binding of the C-terminal region of NUPR1.
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16 **5. CONCLUSIONS**

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19 Our results represent a proof-of-concept for the use of peptides with tuned physico-chemical
20 properties to bind an IDP, and without any homology of sequence with respect to the target IDP.
21 Furthermore, they show that peptides can be effectively used *in cellulo* as inhibitors of PPIs essential
22 for IDPs biological functions. These findings open the possibility to use designed peptides (or
23 peptidomimetics) as regulators of these challenging "druggable-targets".
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31 **AUTHOR CONTRIBUTIONS AND COMPETING INTERESTS**

32
33 PSC, JLI, OA, AVC, BR and JLN designed the experiments and the research methodology.
34 PSC, AVC, OA, BR and JLN carried out the experiments and analyzed data. PSC, JLI, BR, AVC,
35 OA and JLN wrote the paper. The authors declare no competing interests.
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6 **SUPPLEMENTARY INFORMATION**

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9 There are ten figures describing: the interaction of P-1, P-2 and P-4 with wild-type NUPR1
10 followed by ANS fluorescence (Fig. S1); the interaction of P-3 with Thr68Gln mutant by
11 fluorescence (Fig. S2) and ITC (Fig. S3); the fluorescence titrations of P-1 and P-4 (Fig. S4); the
12 interaction of wild-type NUPR1 with P-3 monitored by CD (Fig. S5); the 2D ¹H, ¹⁵N-HSQC NMR
13 spectra of NUPR1 when P-1, P-3 and P-4 are added (Fig. S6); the decrease in absolute intensity for
14 selected residues measured in the HSQC cross-peaks for the four peptides (Fig. S7), as their
15 concentration was increased; the energetic results obtained by molecular docking (Fig. S8); the
16 capture mechanism of a helical peptide by NUPR1 in simulation under conditions favouring a
17 relatively less compacted protein state (Fig. S9); and the control experiment with the test peptide kit
18 for the *in cellulo* experiments (Fig. S10).
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Table 1: Sequences, percentage of helical structure and thermodynamic parameters in the NUPR1 binding of the peptides ^a.

Peptide	% of helical structure (AGADIR) ^b	% of helical structure (from CD measurements) ^b	K_a (M ⁻¹) ^c	K_D (μM) ^c	ΔH (kcal mol ⁻¹) ^c	K_D (μM) ^d
Ac-VKNWMTETLLRQ-NH ₂ (P-1)	13.90	8.4	9.4×10^5	1.1	1.3	2.3 ± 0.8
Ac-VKNWMTEYLLVQ-NH ₂ (P-2)	24.80	5.7	1.8×10^6	0.57	1.7	1.6 ± 0.7
Ac-VKNWMTEYLLRQ-NH ₂ (P-3)	52.88	8.2	3.1×10^5	3.2	1.2	2 ± 1
Ac-VKNWMTETLLVQ-NH ₂ (P-4)	4.59	6.9	4.4×10^5	2.2	1.5	0.4 ± 0.2

^aThe P-4 is the wild-type sequence. Errors in the calorimetric measurements are estimated to be 15-20 % for K_a and K_D and 5% for ΔH .

^bData from (26).

^cDetermined from ITC measurements. Measurements were carried out at 15 °C.

^dDetermined from fluorescence titrations. Errors are fitting errors to Eq. 1. Measurements were carried out at 25 °C.

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Table 2: Resonances of NUPR1 residues, whose relative intensities (with respect to Arg82) were affected by the presence of the peptide ^a

Peptide	Resonances ^b
Ac-VKNWMTETLLRQ-NH ₂ (385 μM) (P-1)	Thr8 (+); Glu18; Ala33 (+); Gly39; Gly44; Ala50 (+); Glu63; Asn72; Ser73
Ac-VKNWMTEYLLVQ-NH ₂ (385 μM) ^c (P-2)	All cross-peaks broadened
Ac-VKNWMTEYLLRQ-NH ₂ (405 μM) (P-3)	Thr3; Thr8 (+); Glu18; Gly39; Gly44; Ala50; Thr54; Glu63; Asn72
Ac-VKNWMTETLLVQ-NH ₂ (230 μM) (P-4)	Thr8 (+); Glu18; Ala33 (+); Gly39; Gly44; Glu63; Asn72

^aOnly residues whose signals could be unambiguously integrated (that is, non-overlapping cross-peaks) are indicated. The peptide concentration used is indicated within parenthesis beside each peptide sequence. NUPR1 concentration was 190μM. Experiments were carried out at pH 4.5 and 25 °C.

^bResidues with “+” sign indicate that the intensity variation was ~45 % of the relative intensity of the same cross-peak in the spectrum of the isolated NUPR1. Residues that showed variations in the relative intensity of cross-peaks (when compared to that in isolated NUPR1) larger than 15 % are indicated.

^cP-2 was also tested at 430 μM concentration.

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FIGURE LEGENDS

FIGURE 1: Intrinsic and ANS-fluorescence characterization of the binding between P-3 and wild-type NUPR1: (A) Fluorescence spectrum after excitation at 370 nm, in the presence of ANS, of the complex P-3/NUPR1 (15: 20 μ M) and that obtained by the addition of the spectra of each isolated molecule. (B) Fluorescence spectrum, after excitation at 280 nm, of the complex P-3/NUPR1 and that obtained by the addition of the spectra of each isolated molecule. (C) Fluorescence thermal denaturations followed by the emission at 480 nm, after excitation at 370 nm (in the presence of ANS) of the isolated NUPR1 (green, blank circles), P-3 (blue line) and of the complex (black line).

FIGURE 2: ITC of wild-type NUPR1 and the peptides: Calorimetric titrations for (from left to right): P-1, P-2, P-3 and P-4. Thermograms (top) and binding isotherms (bottom) are shown. Non-linear fittings according to a model considering a single ligand binding site (continuous lines) are shown for each peptide.

FIGURE 3: Interaction of wild-type NUPR1 and P-2 mapped by 2D 1 H, 15 N-HSQC spectra: Overlay of wild-type NUPR1 spectra at 0 (black); and 385 (red) μ M (in protomer units) of P-2. The same low contour level was used for both spectra. Experiments were acquired at pH 4.5 and 25 $^{\circ}$ C.

FIGURE 4: Preferred binding poses of the seven-residue-long NUPR1 sequence fragment centered on Ala33 onto the peptides. Binding modes calculated by using the AutoDock Vina (50) docking engine for the protein segment 30-36 of NUPR1 are mapped onto the ideal helical structure of the peptides (superimposed by rototranslation on the C^{α} atoms, and represented as a solid tube with the position of amino acids 1-7 shown as transparent volumes). The wheel scheme corresponding to an ideal α -helix is

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4 also displayed on the left, with the region showing the highest hydrophobicity indicated with a black line
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11 **FIGURE 5: Capture mechanism of an α -helical peptide by NUPR1.** The protein is oriented with the
12 more compact N-terminal region at the bottom and the extended C-terminal region on top, with selected
13 residues labeled. The P-2 peptide (in orange) is shown using a surface representation of the backbone.
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21 **FIGURE 6: *In situ* PLA of peptides and wild-type NUPR1.** Mouse anti-HA and rabbit anti-human
22 wild-type NUPR1 were used to reveal the interaction between the C-RING1B and NUPR1 in HeLa
23 cells. PLA was carried out as described (22,23) after cells were treated with the peptides (170 μ M) for 4
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Figure 1

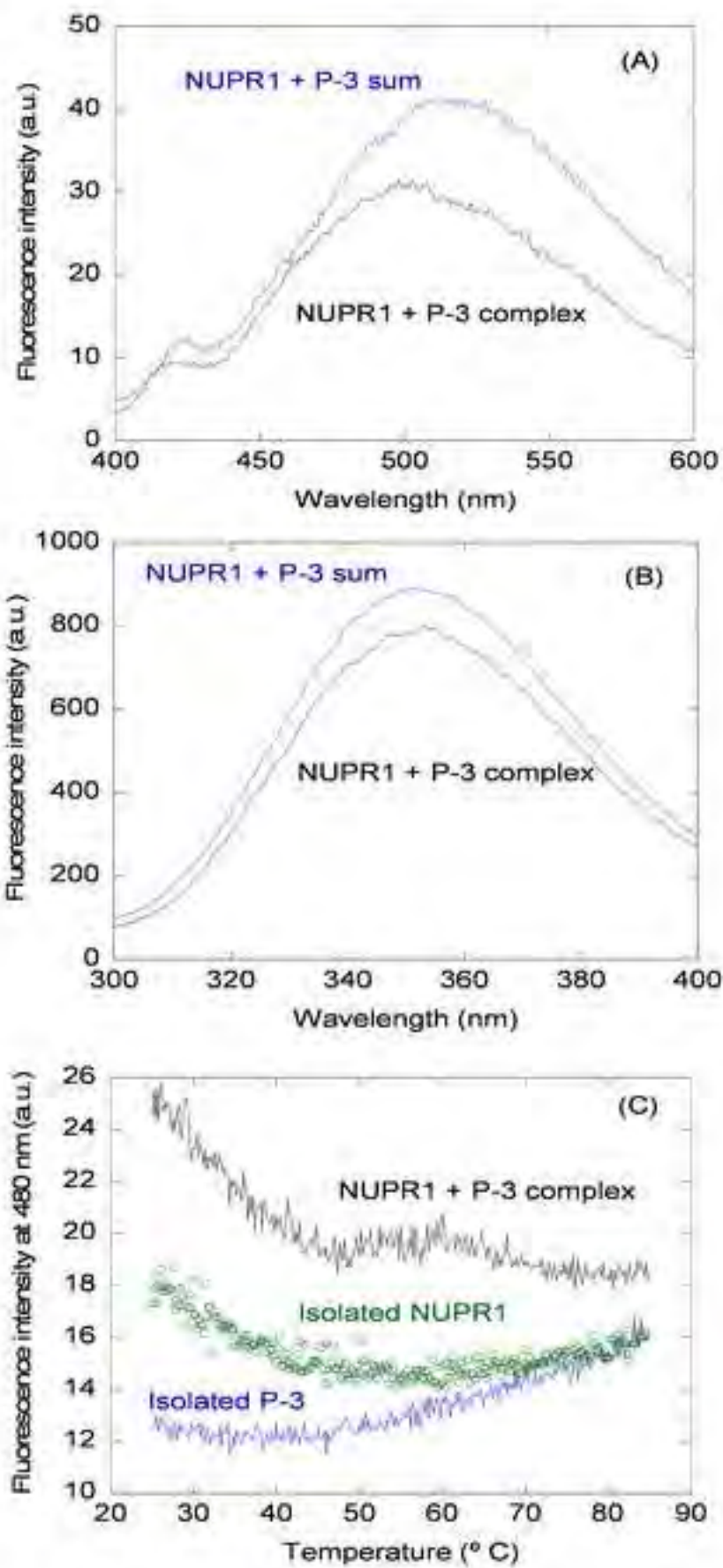


Figure 2

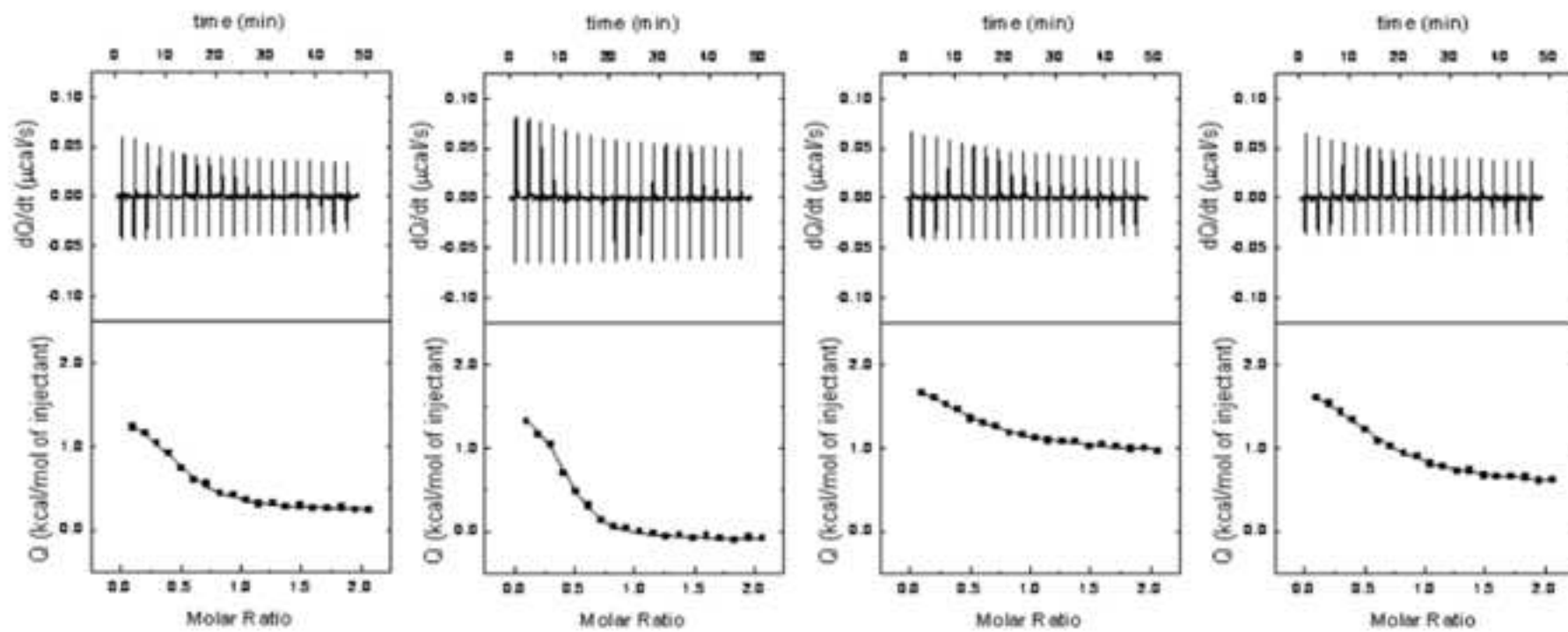


Figure 3

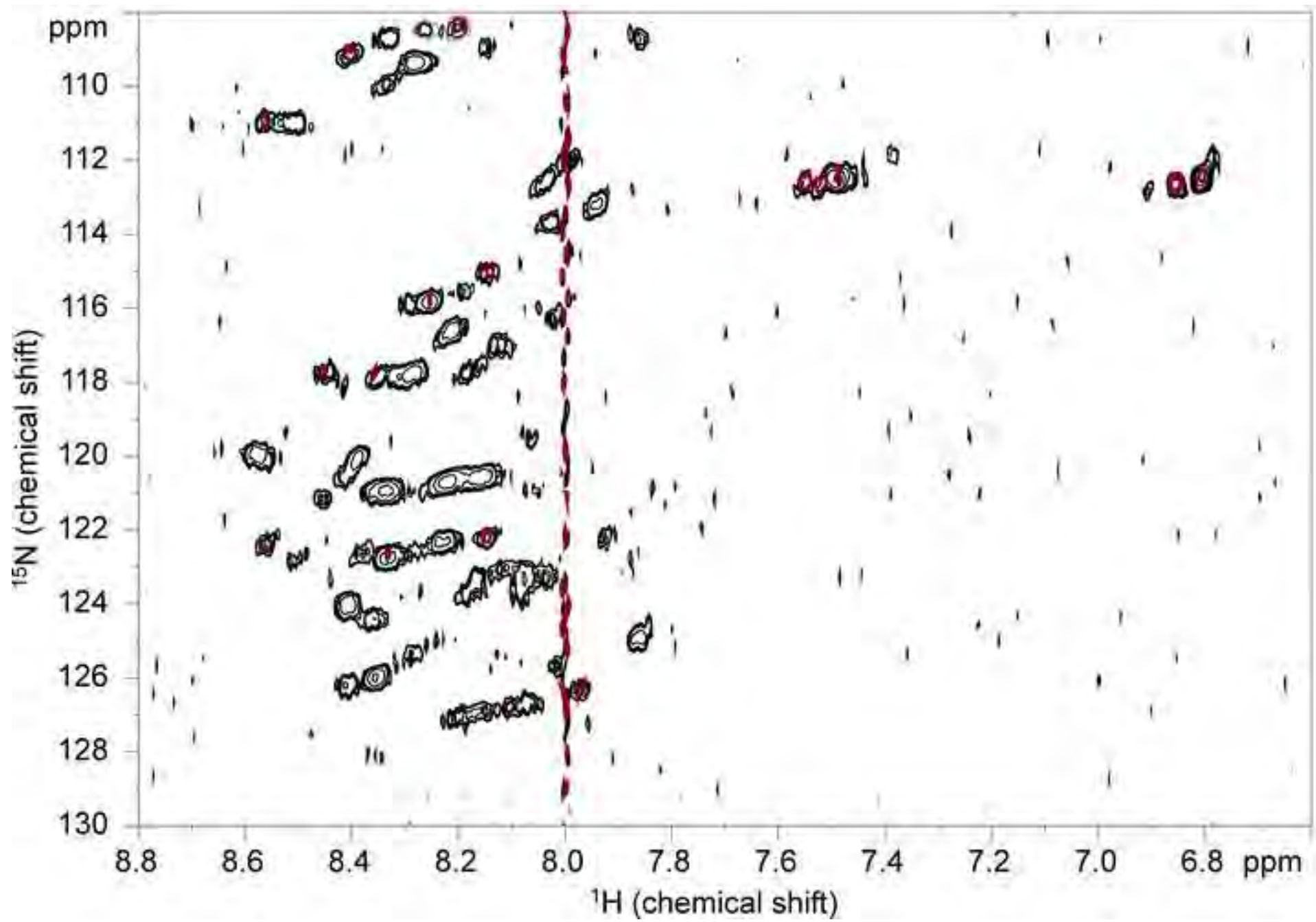


Figure 4

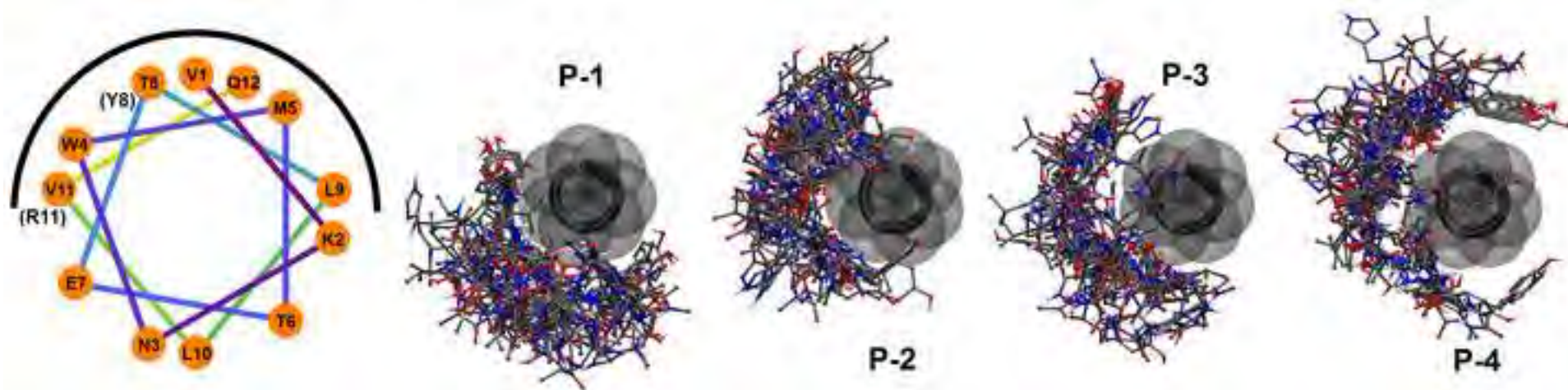


Figure 5

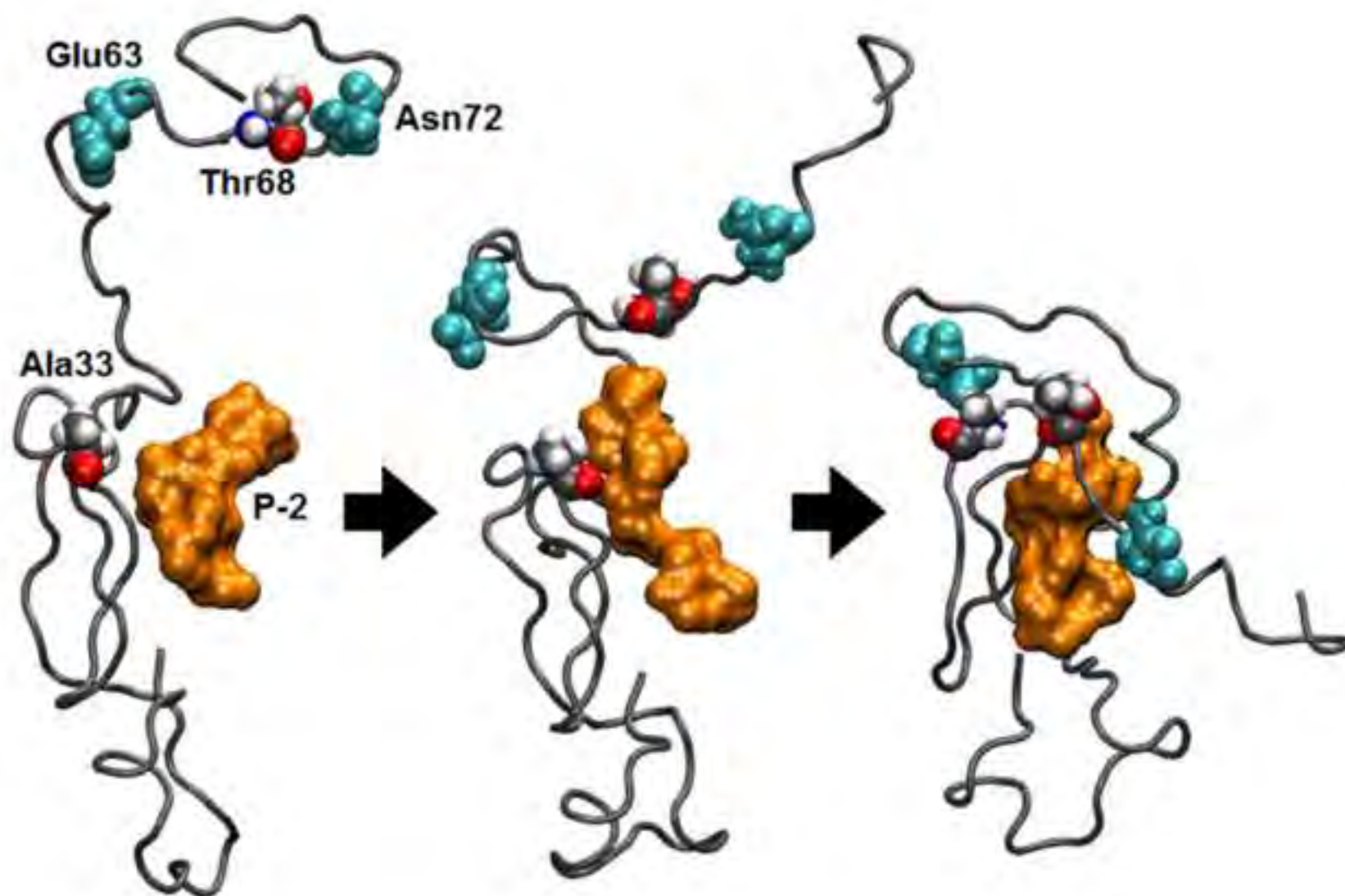
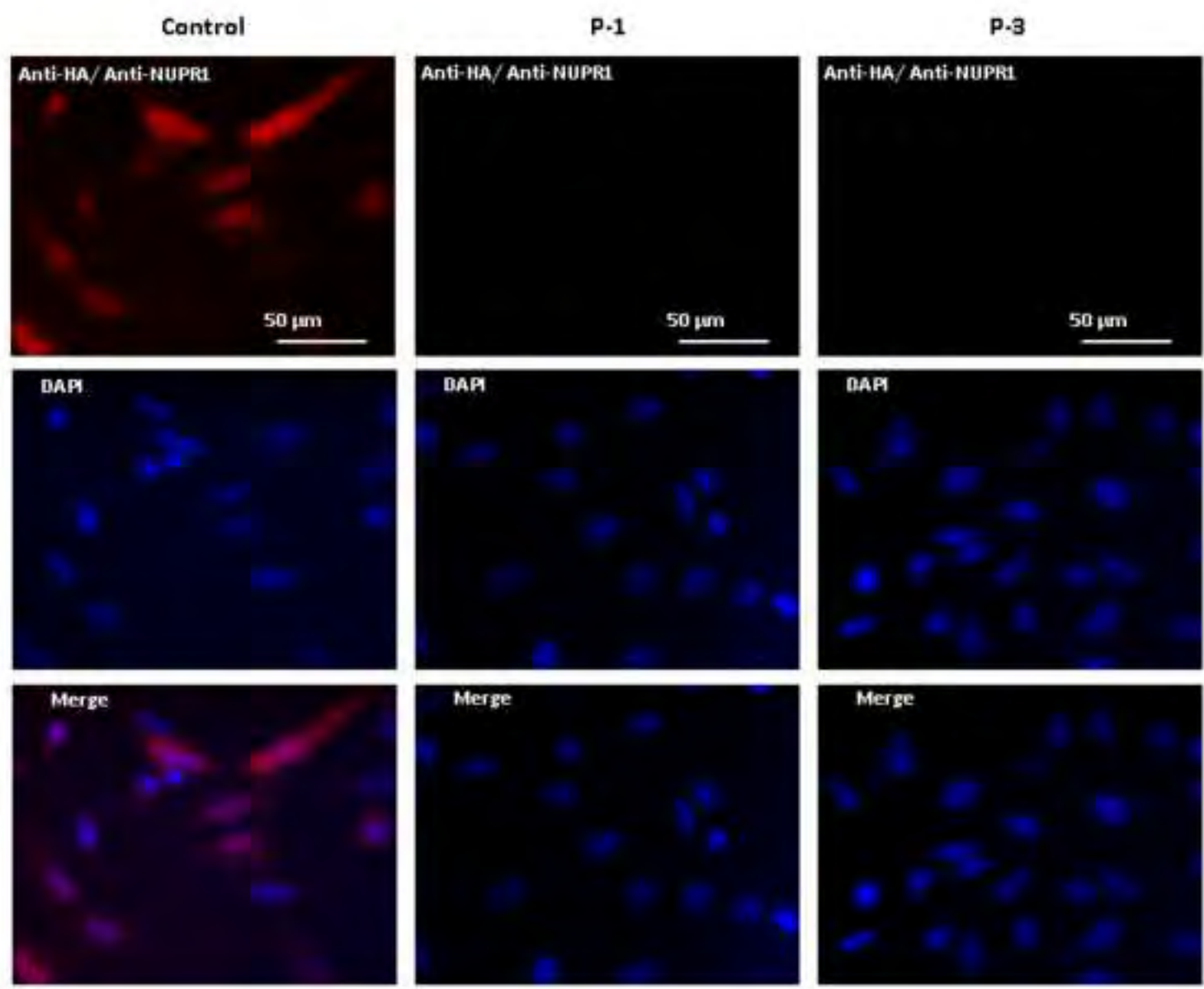
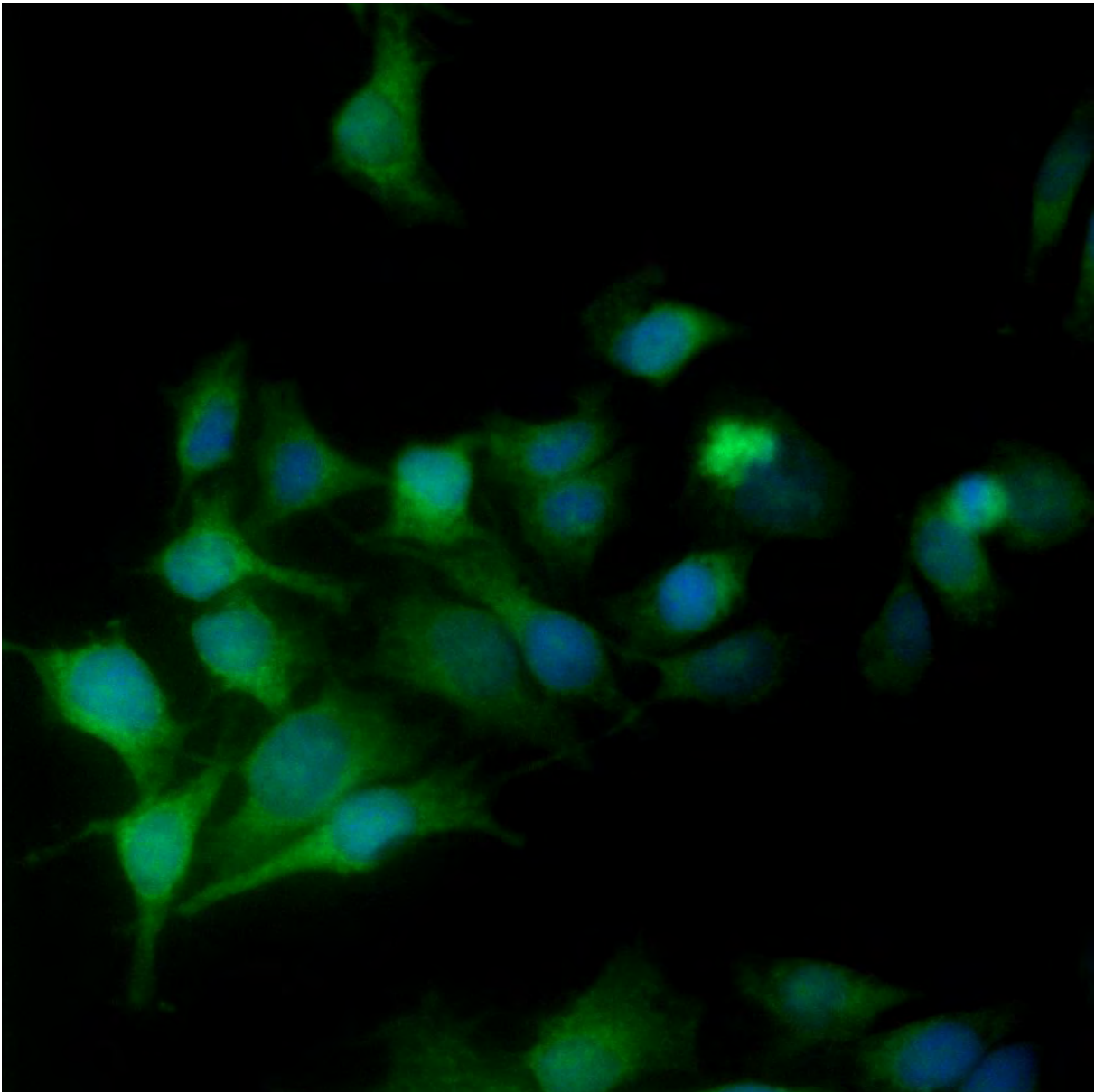


Figure 6





*Conflict of Interest form

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