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# Restricted binding of a model protein on C<sub>3</sub>N<sub>4</sub> nanosheets suggests an adequate biocompatibility of the nanomaterial†

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Recently, C<sub>3</sub>N<sub>4</sub>, a carbon nitride nanomaterial, has attracted great attention in many scientific fields due to its outstanding properties. Specifically, this nanomaterial has displayed non- or low-toxicity in biological systems suggesting its excellent biocompatibility and biosafety. Nevertheless, few studies address the structural consequences from the direct interaction between C<sub>3</sub>N<sub>4</sub> and biomolecules that could imply the physical origin of its bio-effect, particularly from the molecular level. Herein, we explored the interaction of a C<sub>3</sub>N<sub>4</sub> nanosheet and a model protein, the λ-repressor protein. We found that the C<sub>3</sub>N<sub>4</sub> nanosheet has a limited influence on the structure of the λ-repressor protein, which substantiates the outstanding biocompatibility of the nanomaterial. Detailed analyses showed that upon absorption on the C<sub>3</sub>N<sub>4</sub> nanosheet, the λ-repressor protein remains located in a relatively fixed position without compromising the structural integrity of the protein. Furthermore, the protein-nanomaterial interaction is mediated by positively charged residues located on the surface of the protein and by the regional negatively charged center on the C<sub>3</sub>N<sub>4</sub> nanosheet (*i.e.*, N-rich defects). These findings provide further molecular-level insights into the good biocompatibility of the C<sub>3</sub>N<sub>4</sub> nanomaterial and also suggest its potential usage as a protein drug delivery platform.

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## Introduction

The rapid development of carbon-based nanomaterials (CBNs) has inspired tremendous interests in biomedical applications,<sup>1,2</sup> including nanotherapeutics,<sup>3–5</sup> optical imaging,<sup>6</sup> and gene delivery,<sup>7</sup> because of their outstanding optical, mechanical, and electrical properties.<sup>8–10</sup> Recently, ordered carbon nitrides (*e.g.* C<sub>2</sub>N,<sup>11</sup> C<sub>3</sub>N,<sup>12,13</sup> and C<sub>3</sub>N<sub>4</sub> (ref. 14)), a new family class of CBNs, have also greatly stimulated research enthusiasm, due to their inherent electronic and optoelectronic properties. Based on their surprising electronic conductivity and optical properties, carbon nitrides were successfully utilized to produce various devices, such as photoelectrical devices, sensors, field-effect transistor devices, and so on.<sup>12,15,16</sup> Carbon nitrides also exhibited interesting catalytic activity for the H<sub>2</sub> evolution and oxygen reduction reaction.<sup>17–19</sup> Moreover, some carbon nitrides exhibited promising potential in biomedical applications. For instance, ultrathin graphitic-

phase C<sub>3</sub>N<sub>4</sub> (g-C<sub>3</sub>N<sub>4</sub>) nanosheets, prepared by a “green” liquid exfoliation pathway from bulk g-C<sub>3</sub>N<sub>4</sub> in water, displayed outstanding properties including (i) good stability in both acidic and alkaline solvents, (ii) intensive photoabsorption and photoresponse, and (iii) excellent biocompatibility.<sup>20</sup> The g-C<sub>3</sub>N<sub>4</sub> has also been utilized as a platform carrying up-conversion nanoparticles for cancer photodynamic therapy.<sup>21</sup> Meanwhile, g-C<sub>3</sub>N<sub>4</sub> was considered an excellent biosensor, due to its metal-free, high fluorescence quantum yield, easy preparation, high stability as well as adequate biocompatibility.<sup>22,23</sup> For example, a g-C<sub>3</sub>N<sub>4</sub> single-layer quantum dot was introduced for fluorescence imaging of the cellular nucleus.<sup>24</sup> Through hybridization with gold nanoparticles, the g-C<sub>3</sub>N<sub>4</sub> material was exploited as an electrochemiluminescence immunosensor.<sup>25</sup> The ultrathin g-C<sub>3</sub>N<sub>4</sub> nanosheet was developed for the detection of alkaline phosphatase exhibiting label-free high sensitivity in biological systems.<sup>26</sup> The protonated g-C<sub>3</sub>N<sub>4</sub> nanosheets without metal/label functionalization were established as a sensing platform for highly sensitive and selective detection of heparin.<sup>27</sup> Although hybridized with some components, the g-C<sub>3</sub>N<sub>4</sub> material was also exploited for biosensing of other molecules, *e.g.*, hybridized with MOF for amyloid-β protein sensing,<sup>28</sup> hybridized with MnSe for H<sub>2</sub>O<sub>2</sub> and glucose sensing,<sup>29</sup> hybridized with molecularly imprinted polymers for H<sub>2</sub>O<sub>2</sub> sensing,<sup>22</sup> *etc.*<sup>30–32</sup>

Despite the widespread biomedical applications, thus far, there is very limited knowledge regarding the interaction of the C<sub>3</sub>N<sub>4</sub> nanomaterial with biomolecules. Detailed knowledge of

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the biomolecule- $C_3N_4$  (bio-nano) interaction is central to support the desired biocompatibility of  $C_3N_4$ . Recently, Zhao *et al.* investigated the interaction between amino acid residues and a  $C_3N_4$  nanosheet by employing density functional theory (DFT) and molecular dynamics simulations.<sup>33</sup> They found that  $C_3N_4$  preferentially attracted the backbone amino ( $NH_2$ ) group, yet this group appeared infrequently in any protein due to the dehydration synthesis from amino acids to protein. However, the positively charged side-chain groups from lysine and arginine may follow similar behavior regarding their preference to contact the  $C_3N_4$ . In this study, we probed the binding of a model protein, the  $\lambda$ -repressor protein, to a  $C_3N_4$  nanosheet using all-atomic molecular dynamics (MD) simulations. We observed that the transverse migration of the  $\lambda$ -repressor protein on the  $C_3N_4$  plane was significantly restricted, while the same protein on the graphene plane moved relatively free and randomly. Also, the protein structure exhibited little change throughout the binding process, suggesting excellent biocompatibility of the  $C_3N_4$  nanomaterial. In addition, the positively charged residues were robustly attracted by the N-rich defects on the  $C_3N_4$  surface, which was critical to the confined combination. Our works revealed the interaction basis at the molecular level of the  $C_3N_4$  nanosheet and a model protein, which might be directly correlated with the nanomaterial's adequate biocompatibility that is central for its utility in biomedical devices.

## Results

In order to investigate the binding of the  $\lambda$ -repressor protein to the  $C_3N_4$  surface, we built an all-atom system consisting of a single biomolecule (the  $\lambda$ -repressor protein) and the  $C_3N_4$  nanosheet as shown in Fig. 1. The system was investigated by unbiased MD simulations and five independent 200 ns-long trajectories were generated (see Methods for details). To assess the stability of the secondary and tertiary structures of the protein upon surface adsorption, we performed analyses of several parameters along the different simulations, including the root mean square deviations (RMSDs) of  $C\alpha$  atoms, the fraction of native contacts ( $Q$  values) and the  $\alpha$ -helix ratio (Fig. 2). Herein, a direct contact of the protein with the  $C_3N_4$  surface was considered when any heavy atom of the  $\lambda$ -repressor

protein was within 0.6 nm of any atom of  $C_3N_4$  nanomaterial. Using the same direct contact criterion for the cut-off distance, the fraction of native contacts of protein over time  $t$ ,  $Q(t)$ , was defined as the ratio of the total number of native contacts at time  $t$  relative to that at time zero (*i.e.*, the total number of native contacts as observed in the X-ray crystal structure); for this analysis, only residue pairs apart by at least 3 consecutive residues from each other were considered. The analysis indicates that the protein conserved its structural integrity in the five independent MD trajectories. Specifically, the RMSD values for the protein  $C\alpha$  atoms remained lower than 0.5 nm and even maintained values of  $\sim 0.2$  nm in three out of five trajectories, suggesting that the  $C_3N_4$  nanosheet had little influence on the overall protein structure (*i.e.*, the nanomaterial displays adequate biocompatibility). Moreover, both the  $Q$  values and the  $\alpha$ -helix ratios also indicated a minor impact on the conformations sampled by the protein during the entire adsorption process, which substantiates the aforementioned conclusion regarding the protein structure. The mild effect of the  $C_3N_4$  nanosheet on the protein tertiary structure substantially emerges from the particular molecular features of the surface (see detailed discussion below), which varies from that of pristine graphene.<sup>34–36</sup> Fig. 3a illustrated the distributions of the center of mass (CoM) of the  $\lambda$ -repressor protein mapping onto the  $X$ - $Y$  plane in the five trajectories. Clearly, during the entire simulation time, the CoMs of the protein was restricted on the  $C_3N_4$  nanosheet in all five trajectories, showing a localized binding pattern. For comparison, we also performed another simulation in which the  $C_3N_4$  nanosheet was replaced by a graphene surface (Fig. 3b). We similarly tracked the CoM of the  $\lambda$ -repressor protein on the graphene surface and found a different situation: the same protein moved quickly and randomly on the graphene plane. This difference was fundamentally caused by the ordered charge distribution and abundant defects on the  $C_3N_4$  nanosheet which is in stark contrast to graphene's zero charge and uniform surface.

In order to explore the adsorption molecular determinants and interaction mechanism of the  $\lambda$ -repressor protein binding to the  $C_3N_4$  nanosheet, we calculated the heavy atom contact number, the van der Waals (vdW), and coulombic energy contributions between the protein and the nanosheet, as well as the residue-dependent heavy atom contact number map



Fig. 1 (a) The  $C_3N_4$  nanosheet. The cyan and blue dots indicated the carbon and nitrogen atoms, respectively. The black circle showed the defect surrounded by nitrogen atoms. (b) Side view of the simulated system. (c) Top view of simulated system where the  $\lambda$ -repressor protein was shown in lime cartoon representations.

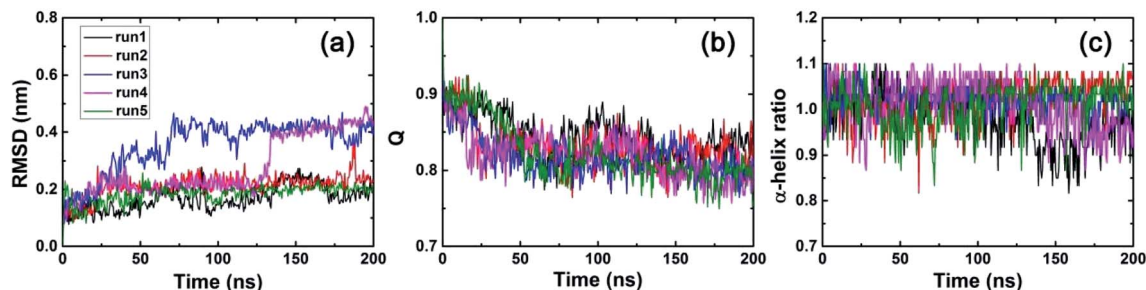


Fig. 2 (a) Root mean square deviations (RMSDs) of the  $C_{\alpha}$  atoms of the  $\lambda$ -repressor protein in five independent trajectories. (b)  $Q$  values for the  $\lambda$ -repressor protein in five trajectories. (c) The  $\alpha$ -helix ratio of the  $\lambda$ -repressor protein during simulation trajectories with respect to the values in the crystal structure.

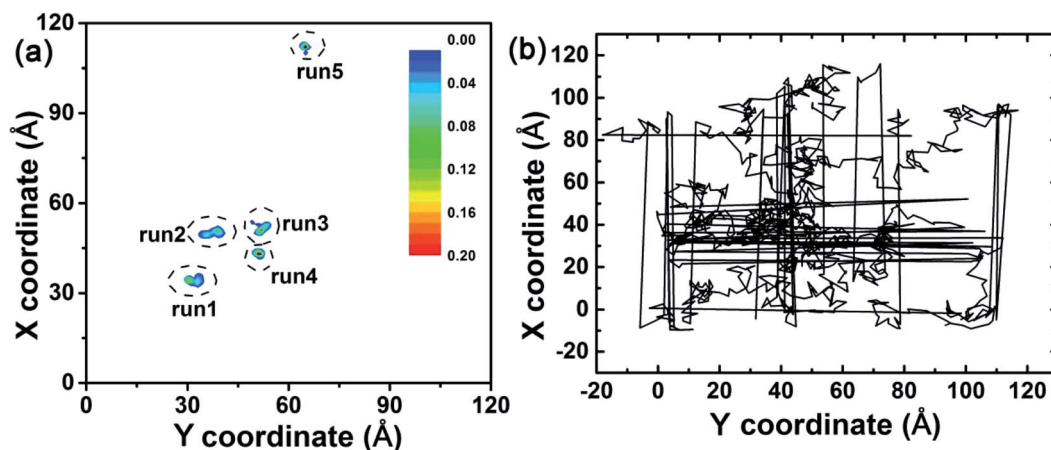


Fig. 3 (a) Center of mass (CoM) distributions of the  $\lambda$ -repressor protein mapping onto the  $X$ - $Y$  plane in five trajectories. (b) CoM track of the  $\lambda$ -repressor protein when bound to a graphene surface.

(Fig. 4a–c). Additionally, according to their tendencies, we described in detail the adsorption process by plotting some snapshots at selected key time points along the entire trajectory (Fig. 4d). Specifically, at early stages,  $t = 10$  ns, the  $\lambda$ -repressor protein approached the  $C_3N_4$  nanosheet, with some residues directly contacting the surface, *i.e.*, residues E22, K24, K25, S72, E75, and Y85. As expected, the values for the heavy atom contact number sharply increased to a value of around 43. Meanwhile, the vdW energy and the Coulombic energy decreased their values to  $-72.46$  kcal mol $^{-1}$  and  $-17.13$  kcal mol $^{-1}$ , respectively. The larger decrement of its magnitude from the former energy term suggests that the vdW interactions may drive the initial protein-surface interactions. From 10 ns to 37 ns, the binding profile fluctuated significantly. At around  $t = 37$  ns, the binding seems to reach a metastable state with minor increments in the protein-surface contact values mediated mainly by the interaction of the E89 and S92 residues (heavy atom contact number:  $\sim 47$ , vdW energy:  $\sim -88.66$  kcal mol $^{-1}$ , coulombic energy:  $\sim -37.49$  kcal mol $^{-1}$ ). At around  $t = 102$  ns, and after the packing of residues E89, Y88, and Y85 (Fig. 4c), the adsorption event was further strengthened, where the heavy atom contact number reached values of close to 61 while the vdW and coulombic energies reached close to  $-122.29$  and  $-31.10$  kcal mol $^{-1}$ , respectively. Around  $t = 188$  ns, the protein

contact intensified slightly after the adsorption of the R82 residue. Even though the  $C_3N_4$  nanomaterial comprised various negatively charged defects in its structure, during the absorption event it is the vdW interactions, and not the coulombic interactions, that seem to dominate the binding process. Yet interestingly, the positively charged residues account for around 30% of all the contact residues.

Due to the particular chemical topology of the  $C_3N_4$  surface, the defects on the  $C_3N_4$  nanosheet are surrounded by highly-concentrated negative charges (*i.e.*, nitrogen atoms, as indicated in Fig. 1a) which enabled a specific attraction of these sites to the positively charged residues *via* electrostatic attraction. Therefore, we next explored the binding character of positively charged residues (choosing residue K24 as an example) as shown in Fig. 5. We first calculated the interaction energies between the K24 sidechain and the  $C_3N_4$  nanosheet. Interestingly, both the vdW and the coulombic energies showed a sharp decline in the initial value of the contact number ( $\sim 10$  ns). To interpret the absorption process of the K24 residue in the proximity of the  $C_3N_4$  surface, we described in detail some snapshots from the MD trajectory as shown in Fig. 5c. Clearly, from 22.2 ns to 22.3 ns, the positively charged amino group of K24 quickly approaches the N-encircling defect in the surface. This event seems to be driven by the conjunct contributions,

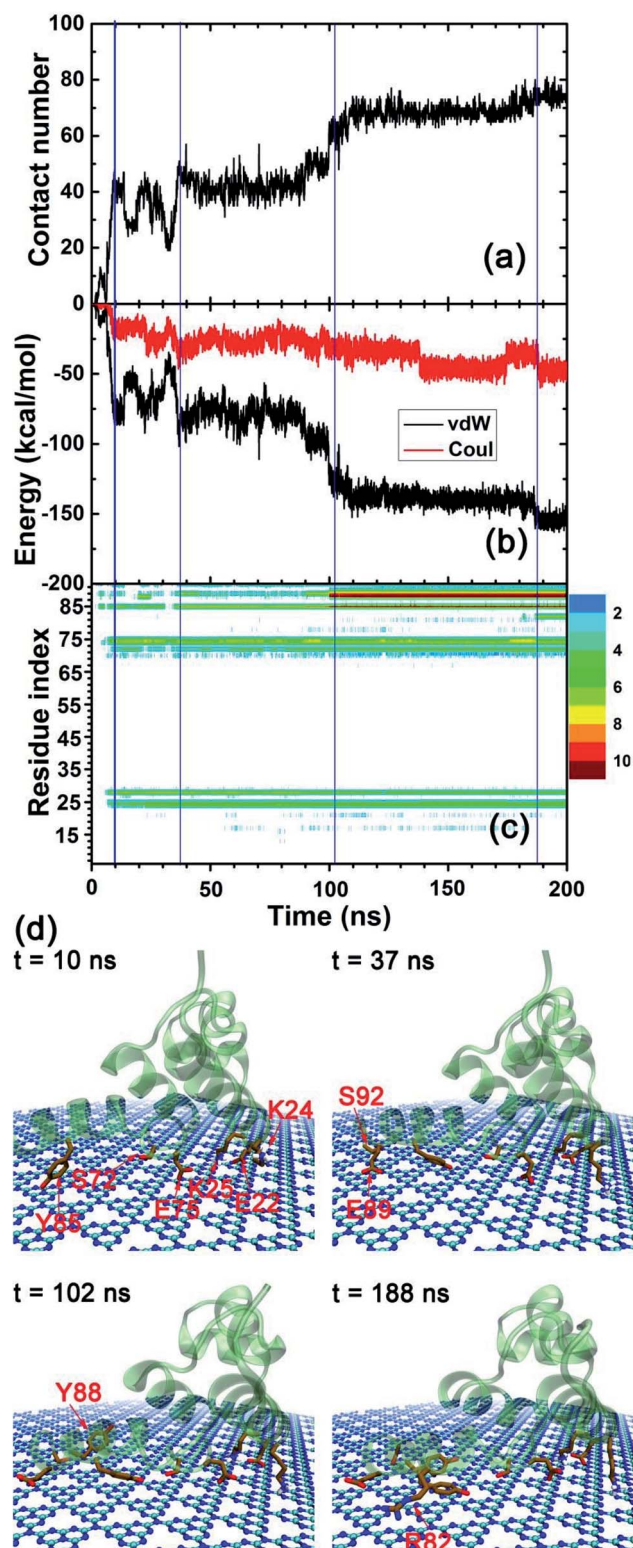


Fig. 4 (a) Heavy atom contact number of the  $\lambda$ -repressor protein adsorbed onto the  $C_3N_4$  nanosheet. (b) van der Waals and coulombic energies between the  $\lambda$ -repressor protein and the  $C_3N_4$  nanosheet. (c) Residue-specific heavy atom contact number map for the  $\lambda$ -repressor protein. (d) Representative snapshots for the binding event at some key time points according to the tendencies of (a–c) figures.

vdW and coulombic; the coulombic energy values range from  $-0.05 \text{ kcal mol}^{-1}$  to  $-13.43 \text{ kcal mol}^{-1}$  while the vdW energy values range from  $-4.21 \text{ kcal mol}^{-1}$  to  $-12.21 \text{ kcal mol}^{-1}$ . Yet, soon after, at  $t = 22.4 \text{ ns}$  and due to minor conformational perturbations of the K24 sidechain, the binding strength weakened significantly with a major contribution by the coulombic energy, which suggests that small conformational changes of the lysine residue can cause a large change in the value of this energy term. At  $t = 23.0 \text{ ns}$ , the conformation recovered completely. We identified that the conformations where the K24 amino sidechain is directly positioned on the  $C_3N_4$  surface defect were the structurally most stable and energetically more favorable binding modes; where the main contributions arise from the coulombic energy term. Moreover, our previous study also found a similar interacting pattern by the HP35 protein binding onto the  $C_2N$  surface, suggesting more general binding features by this type of nanomaterials.<sup>37</sup> Next, we mapped the location of the nitrogen atom from the amino group of residue K24 during the last 100 ns trajectory, on the  $X$ - $Y$  plane (same plane as the  $C_3N_4$  surface) as well as on the  $Z$  direction (perpendicular to the  $C_3N_4$  plane), respectively (Fig. 5d and e). Remarkably, the largest distribution is located at the center of the N-encircling defect. Moreover, along the perpendicular direction, the position of this atom is mostly maintained at a distance of 0.20–0.23 nm over the  $C_3N_4$  plane. This finding demonstrates the significance of the specific binding of basic residues to the  $C_3N_4$  defects.

To probe if the aforementioned electrostatic interaction played a role in the observed fixed binding, we conducted a simulation by changing all atomic charges of  $C_3N_4$  to zero. Notably, by tracking the CoM of the protein (as shown in Fig. S2†), we still observed a restricted binding of the protein on the  $C_3N_4$  nanostructure, which thereby suggests that the electrostatic interactions are not the main factor for the observed fixed binding pattern of the protein but that the vdW interactions alone can induce the determinants of the observed binding event. We have demonstrated that the protein can freely move on graphene (Fig. 3b), and since the only difference between the two 2D nanomaterials is the surface topological structure, we concluded that the vdW interactions are enough to restrain the movement of the adsorbed protein and the key reason for the observed fixed binding pattern is the presence of significant defects in the nanomaterial surface topology. These defects could generate high free energy barriers and would affect the free protein movement, resembling the case for the  $C_2N$  nanosheet.<sup>37</sup> Further analysis of the surface water distribution on the  $C_3N_4$  structure (Fig. S3†) showed that the surface defects aggregate many water molecules (with one water located nearby a structural defect), which forms a specific water layer in the proximity of the  $C_3N_4$  material. This water layer may also contribute to the fixed binding of protein by forming a steric hindrance to prevent the lateral displacement of the protein, which resembles the results from our previous study regarding the dsDNA binding to the 2D  $C_2N$  nanosheet.<sup>38</sup> To sum up, the restricted protein binding on the  $C_3N_4$  nanosheet is mainly attributed to the inherent porous structure of the  $C_3N_4$  sheet as well as the water aggregation in the proximity of the surface.

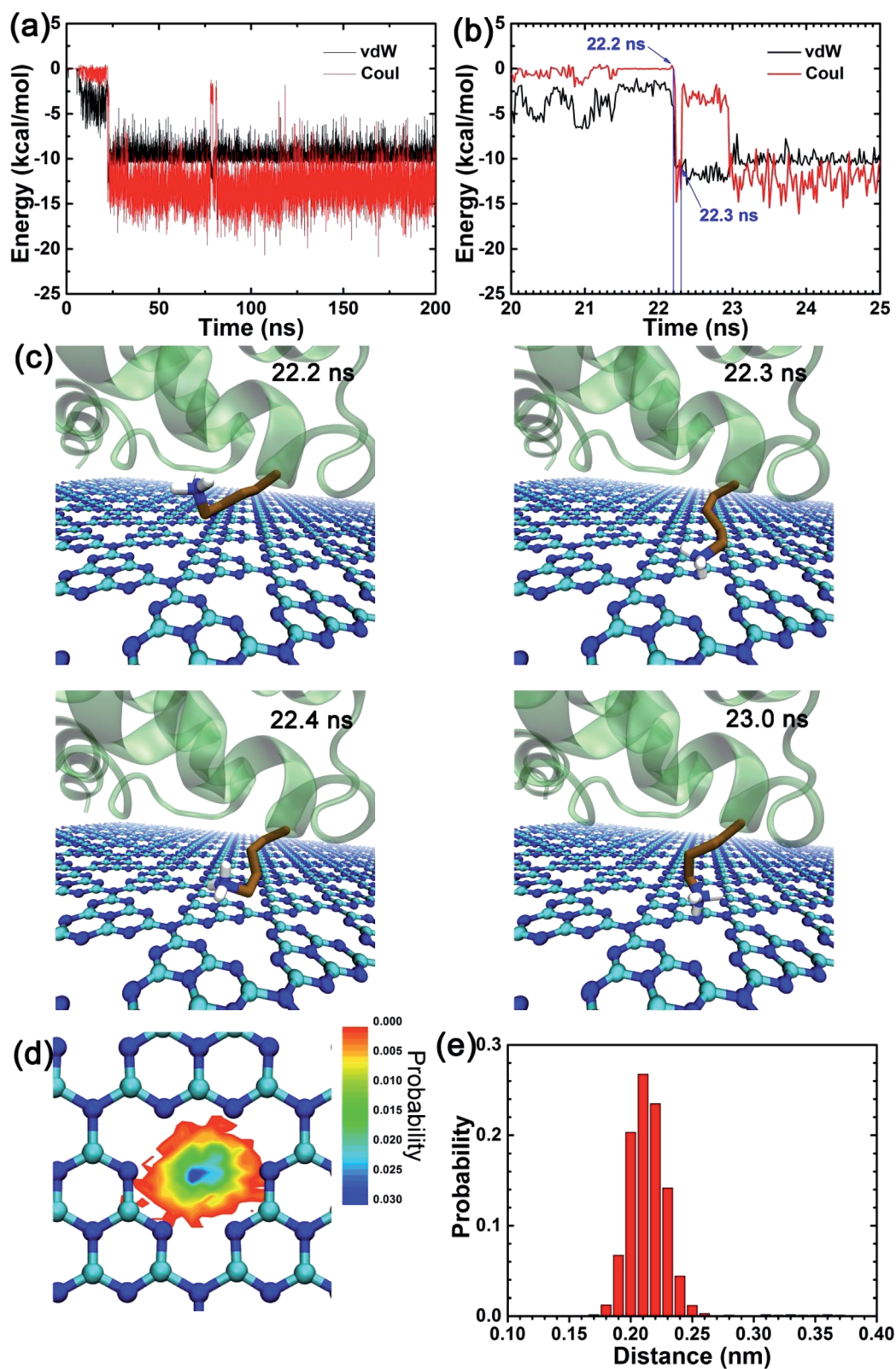


Fig. 5 (a and b) The van der Waals (vdW) and coulombic (Coul) energies between the side chain of residue K24 and the  $C_3N_4$  nanosheet. (c) Snapshots at key time points showing the K24 different conformations for the amino side-chain group. (d) The distribution of nitrogen atom from the amino side chain group of K24 mapped onto the X-Y plane after its tight binding (*i.e.*, after 23.0 ns). (e) The distribution probability of same nitrogen atom over the  $C_3N_4$  nanosheet along the vertical direction.

To further confirm if the observed restricted binding pattern occurred in other biomolecules upon binding to the  $C_3N_4$  nanosheet, we conducted MD simulations where we replaced the  $\lambda$ -receptor protein with (i) a double-stranded DNA segment (dsDNA) and (ii) the HP35 protein. As shown in Fig. S4,<sup>†</sup> it is notable that, despite their different physicochemical properties, both the HP35 protein and the dsDNA can bind to the  $C_3N_4$  surface. Also, upon binding, these two biomolecules explored restrained conformations on the  $C_3N_4$  surface comparable to those observed in the case of the  $\lambda$ -receptor protein bound to the  $C_3N_4$  nanosheet. These findings substantiate our conclusion that the observed fixed binding pattern of biomolecules on the  $C_3N_4$  nanosheet is mainly attributed to the inherent porous structure of the  $C_3N_4$  nanomaterial as well as the water distribution in the surface proximity.

Usually, under real experimental environmental conditions, most water-soluble proteins (*e.g.*, serum proteins) present a significant amount of hydrophilic residues on their surface for stability reasons in the water environment, particularly residue with positively charged side chains. In our previous study,<sup>39</sup> we found that solvent-exposed basic residues (Lys and Arg) played an important role in mediating the interactions at the bio-nano interface, *e.g.*, in the serum proteins–graphene interaction. Therefore, an abundance of basic residues may strengthen the protein absorption on the  $C_3N_4$  surface in a physiologically relevant environment, because of the potent attraction between basic residues and the  $C_3N_4$  pores. Similarly, by using the combination of MD simulations and experiments, we discovered that this kind of specific interactions between positively charged groups of bacterial lipid head groups and the  $C_3N_4$  pores, also weakened the  $C_3N_4$  insertion into the bacterial membrane due to the strong binding interactions between the lipid head groups and the  $C_3N_4$  nanomaterial, reflecting the weak antibacterial activity of pristine  $C_3N_4$  (*ref.* 40). However, after nitrogen plasma treatment, the pores of  $C_3N_4$  were sealed, endowing the  $C_3N_4$  nanosheet with a more homogeneous surface that results in an increase in membrane insertion and the concomitant increase in the  $C_3N_4$  antibacterial activity. These findings further demonstrate that the interaction between positively charged chemical moieties (groups/side chains) and the  $C_3N_4$  pores was extremely relevant for the behavior of the  $C_3N_4$  nanomaterial in real experiments. Furthermore, surface functionalizations and larger topological defects on the  $C_3N_4$  nanosheet should also be taken into account, due to the frequent presence during nanomaterial chemical synthesis. We have demonstrated that the main factors leading to the observed fixed protein binding is the porous  $C_3N_4$  surface structure and the specific water distribution in the proximity of the nanosheet (as shown in Fig. S3<sup>†</sup>), instead of the electrostatic interaction. The functionalizations on the  $C_3N_4$  surface will not have a significant impact on the determinants observed in the protein binding to the  $C_3N_4$  nanosheet. The larger defects generate higher free energy barriers for lateral movement (resembling  $C_2N$  nanosheet),<sup>37</sup> which in turn intensifies the fixed binding of biomolecules. Thereby, surface functionalizations may have a limited influence on the observed fixed protein binding pattern.

We also compared our findings with other works regarding the interactions between other 2D nanomaterials and biomolecules as seen in Table S1,<sup>†</sup> particularly focusing on the consequences on the biomolecular structural integrity. Notably, abundant surface defects, such as nitrogen-enriched pores on the  $C_3N_4$  and  $C_2N$  nanomaterials, maybe more biocompatible to biomolecules than more homogeneous materials because their inherent porous structure can significantly restrict the lateral movement of the biomolecules. The 2D nanomaterials without abundant surface pores, such as graphene,  $C_3N$ , boron nitride, and  $MoS_2$ , may have an influence on the tertiary structure of the systems, due to the potential fast lateral migration of biomolecules. Moreover, surface wrinkles (*e.g.*, in phosphorene) and hydrophilic groups (*e.g.*, in silicene) may contribute to protect the biomolecular integrity. However, if the pores on the 2D nanomaterial are distributed less close-packed (the separation distance among these pores was larger than the size of adsorbed biomolecule), the biomolecule may suffer partial or complete denaturation, *e.g.*, the HP35 protein on carboxyl functionalized graphene defects.<sup>41</sup> Furthermore, surface-specific atomic arrangements also impact the structural change of biomolecules, *e.g.*, Pt nanocrystals with (111) and (100) facets. In general, the factors mediating the structural influence of biomolecules after binding to 2D nanomaterials are complex, but it may be highly correlated to the surface characters of these 2D nanomaterials.

## Conclusion

In summary, we have performed unbiased all-atom MD simulations to investigate the binding of the  $\lambda$ -repressor protein to a  $C_3N_4$  nanosheet. Upon the protein's adsorption onto the  $C_3N_4$  surface, the  $\lambda$ -repressor protein retained its structural integrity, that is, the conformations sampled upon adsorption do not change its secondary and tertiary structure. The main driving force involved in the protein's adsorption event was dominated by the van der Waals (vdW) interactions while the coulombic energy term exhibits a more discreet contribution. Moreover, our analysis of the mobility of the center of mass (CoM) of the  $\lambda$ -repressor protein found that the protein remains relatively fixed on the  $C_3N_4$  nanosheet. Detailed analyses showed that the positively charged residues were mainly responsible for this specific binding pattern in which both interactions, vdW and coulombic, guided the determinants of the adsorption process. Our findings demonstrated a special interface of the protein and the  $C_3N_4$  nanosheet and revealed a potential molecular mechanism that substantiates the adequate biocompatibility of the  $C_3N_4$  nanomaterial observed in previous experiments.

## Model and methods

The  $C_3N_4$  nanosheet used here consists of 1344 nitrogen and 1792 carbon atoms with a dimension of  $9.98 \times 9.88 \text{ nm}^2$  as shown in Fig. 1a. Notably,  $C_3N_4$  was a porous 2D nanomaterial, in the sense that every pore is surrounded by six nitrogen atoms forming a negatively charged center (due to inherent charge transfer from the C atoms to the N atoms). The force field parameters for  $C_3N_4$  were obtained from our previous study,<sup>40</sup>

and the details were shown in Fig. S1.† The  $\lambda$ -repressor protein (PDB code 1LMB<sup>42</sup>) was utilized as a protein model to investigate its interaction with the  $C_3N_4$  nanomaterial, due to its previous use in investigating the interactions of biomolecules with graphene.<sup>43</sup> Next, the  $C_3N_4$  nanosheet and the  $\lambda$ -repressor protein were embedded in a box (size:  $9.98 \times 9.88 \times 6.89$  nm<sup>3</sup>) of 21 028 water molecules, with their minimum initial distance setting to 1.5 nm. Two sodium ions were added to neutralize the entire system. Additionally, some other simulation systems were also built to support our conclusion, such as the  $\lambda$ -repressor protein with non-charged  $C_3N_4$ , HP35/ $C_3N_4$ , dsDNA/ $C_3N_4$ .

The MD simulation was carried out with the GROMACS software package (version: 4.6.6).<sup>44</sup> The VMD software<sup>45</sup> was used to visualize the MD trajectories and rendered the molecular snapshots. The CHARMM36 force field<sup>46</sup> was utilized for the protein in this system. TIP3P<sup>47</sup> water model was applied to describe the water molecules. The system's temperature was maintained at 300 K using the velocity-rescale<sup>48</sup> method while its pressure was coupled at 1 atm using the Parrinello–Rahman<sup>49</sup> algorithm with semiisotropic type (coupling along  $Z$  direction). Periodic boundary conditions (PBC) were applied in all directions. The  $C_3N_4$  nanosheet was frozen throughout the simulation which was similar to the protocol used in our previous works.<sup>38,50</sup> The particle mesh Ewald (PME)<sup>51,52</sup> method was used to treat long-range electrostatic interaction (with a cutoff distance of 1.2 nm) and the van der Waals (vdW) interactions were computed with a cutoff distance of 1.2 nm. The LINCS<sup>53</sup> and SETTLE<sup>54</sup> algorithms were employed to constrain the solute bonds associated with hydrogen atoms to their equilibrium values and water geometry, respectively. During the simulations, the time step was set to 2.0 fs and the coordinates were saved every 20 ps. The system was investigated by unbiased MD simulations and performed five independent 200 ns-long trajectories, for a total aggregated simulation time of 1.0  $\mu$ s.

## Conflicts of interest

There are no conflicts to declare.

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## References

- 1 J. S. Lee, H.-A. Joung, M.-G. Kim and C. B. Park, Graphene-based chemiluminescence resonance energy transfer for homogeneous immunoassay, *ACS Nano*, 2012, **6**(4), 2978–2983.
- 2 C. Y. Cha, S. R. Shin, N. Annabi, M. R. Dokmeci and A. Khademhosseini, Carbon-based nanomaterials: multifunctional materials for biomedical engineering, *ACS Nano*, 2013, **7**(4), 2891–2897.
- 3 M. Li, X. Yang, J. Ren, K. Qu and X. Qu, Using graphene oxide high near-infrared absorbance for photothermal treatment of alzheimer's disease, *Adv. Mater.*, 2012, **24**(13), 1722–1728.
- 4 K. Yang, J. Wan, S. Zhang, B. Tian, Y. Zhang, *et al.*, The influence of surface chemistry and size of nanoscale graphene oxide on photothermal therapy of cancer using ultra-low laser power, *Biomaterials*, 2012, **33**(7), 2206–2214.
- 5 Z. Yang, S.-g. Kang and R. Zhou, Nanomedicine: *de novo* design of nanodrugs, *Nanoscale*, 2014, **6**(2), 663–677.
- 6 B. Li, Y. Cheng, J. Liu, C. Yi, A. S. Brown, *et al.*, Direct optical imaging of graphene *in vitro* by nonlinear femtosecond laser spectral reshaping, *Nano Lett.*, 2012, **12**(11), 5936–5940.
- 7 H. Bao, Y. Pan, Y. Ping, N. G. Sahoo, T. Wu, *et al.*, Chitosan-functionalized graphene oxide as a nanocarrier for drug and gene delivery, *Small*, 2011, **7**(11), 1569–1578.
- 8 V. C. Sanchez, A. Jachak, R. H. Hurt and A. B. Kane, Biological interactions of graphene-family nanomaterials: an interdisciplinary review, *Chem. Res. Toxicol.*, 2012, **25**(1), 15–34.
- 9 A. K. Geim, Graphene: status and prospects, *Science*, 2009, **324**(5934), 1530–1534.
- 10 L. Feng and Z. Liu, Graphene in biomedicine: opportunities and challenges, *Nanomedicine*, 2011, **6**(2), 317–324.
- 11 J. Mahmood, E. K. Lee, M. Jung, D. Shin, I.-Y. Jeon, *et al.*, Nitrogenated holey two-dimensional structures, *Nat. Commun.*, 2015, **6**, 6486.
- 12 S. W. Yang, W. Li, C. C. Ye, G. Wang, H. Tian, *et al.*,  $C_3N_4$ -a 2d crystalline, hole-free, tunable-narrow-bandgap semiconductor with ferromagnetic properties, *Adv. Mater.*, 2017, **29**(16), 1065625.
- 13 J. Mahmood, E. K. Lee, M. Jung, D. Shin, H. J. Choi, *et al.*, Two-dimensional polyaniline ( $C_3N$ ) from carbonized organic single crystals in solid state, *Proc. Natl. Acad. Sci. U. S. A.*, 2016, **113**(27), 7414–7419.
- 14 Z. X. Zhou, Y. F. Shen, Y. Li, A. R. Liu, S. Q. Liu, *et al.*, Chemical cleavage of layered carbon nitride with enhanced photoluminescent performances and photoconduction, *ACS Nano*, 2015, **9**(12), 12480–12487.
- 15 Y. Bu, Z. Chen and W. Li, Using electrochemical methods to study the promotion mechanism of the photoelectric conversion performance of ag-modified mesoporous g- $C_3N_4$  heterojunction material, *Appl. Catal., B*, 2014, **144**, 622–630.
- 16 X.-L. Zhang, C. Zheng, S.-S. Guo, J. Li, H.-H. Yang, *et al.*, Turn-on fluorescence sensor for intracellular imaging of glutathione using g- $C_3N_4$  nanosheet-MnO<sub>2</sub> sandwich nanocomposite, *Anal. Chem.*, 2014, **86**(7), 3426–3434.
- 17 X. F. Chen, J. S. Zhang, X. Z. Fu, M. Antonietti and X. C. Wang, Fe-g- $C_3N_4$ -catalyzed oxidation of benzene to phenol using hydrogen peroxide and visible light, *J. Am. Chem. Soc.*, 2009, **131**(33), 11658–11659.
- 18 J. Mahmood, S. M. Jung, S. J. Kim, J. Park, J. W. Yoo, *et al.*, Cobalt oxide encapsulated in  $C_2N$ -h2d network polymer as

- a catalyst for hydrogen evolution, *Chem. Mater.*, 2015, **27**(13), 4860–4864.
- 19 Y. Zheng, Y. Jiao, J. Chen, J. Liu, J. Liang, *et al.*, Nanoporous graphitic-C<sub>3</sub>N<sub>4</sub>@carbon metal-free electrocatalysts for highly efficient oxygen reduction, *J. Am. Chem. Soc.*, 2011, **133**(50), 20116–20119.
- 20 X. D. Zhang, X. Xie, H. Wang, J. J. Zhang, B. C. Pan, *et al.*, Enhanced photoresponsive ultrathin graphitic-phase C<sub>3</sub>N<sub>4</sub> nanosheets for bioimaging, *J. Am. Chem. Soc.*, 2013, **135**(1), 18–21.
- 21 L. L. Feng, F. He, B. Liu, G. X. Yang, S. L. Gai, *et al.*, G-C<sub>3</sub>N<sub>4</sub> coated upconversion nanoparticles for 808 nm near-infrared light triggered phototherapy and multiple imaging, *Chem. Mater.*, 2016, **28**(21), 7935–7946.
- 22 Y. H. Wu, Q. Chen, S. Liu, H. Xiao, M. L. Zhang, *et al.*, Surface molecular imprinting on g-C<sub>3</sub>N<sub>4</sub> photooxidative nanozyme for improved colorimetric biosensing, *Chin. Chem. Lett.*, 2019, **30**(12), 2186–2190.
- 23 Z. X. Zhou, Y. Y. Zhang, Y. F. Shen, S. Q. Liu and Y. J. Zhang, Molecular engineering of polymeric carbon nitride: advancing applications from photocatalysis to biosensing and more, *Chem. Soc. Rev.*, 2018, **47**(7), 2298–2321.
- 24 X. D. Zhang, H. X. Wang, H. Wang, Q. Zhang, J. F. Xie, *et al.*, Single-layered graphitic-C<sub>3</sub>N<sub>4</sub> quantum dots for two-photon fluorescence imaging of cellular nucleus, *Adv. Mater.*, 2014, **26**(26), 4438–4443.
- 25 L. C. Chen, X. T. Zeng, P. Si, Y. M. Chen, Y. W. Chi, *et al.*, Gold nanoparticle-graphite-like C<sub>3</sub>N<sub>4</sub> nanosheet nanohybrids used for electrochemiluminescent immunosensor, *Anal. Chem.*, 2014, **86**(9), 4188–4195.
- 26 M. H. Xiang, J. W. Liu, N. Li, H. Tang, R. Q. Yu, *et al.*, A fluorescent graphitic carbon nitride nanosheet biosensor for highly sensitive, label-free detection of alkaline phosphatase, *Nanoscale*, 2016, **8**(8), 4727–4732.
- 27 T. Y. Ma, Y. Tang, S. Dai and S. Z. Qiao, Proton-functionalized two-dimensional graphitic carbon nitride nanosheet: an excellent metal-/label-free biosensing platform, *Small*, 2014, **10**(12), 2382–2389.
- 28 Y. Wang, Y. Zhang, H. Sha, X. Xiong and N. Jia, Design and biosensing of a ratiometric electrochemiluminescence resonance energy transfer aptasensor between a g-C<sub>3</sub>N<sub>4</sub> nanosheet and Ru@mof for amyloid-beta protein, *ACS Appl. Mater. Interfaces*, 2019, **11**(40), 36299–36306.
- 29 F. M. Qiao, Q. Q. Qi, Z. Z. Wang, K. Xu and S. Y. Ai, MnSe-loaded g-C<sub>3</sub>N<sub>4</sub> nanocomposite with synergistic peroxidase-like catalysis: synthesis and application toward colorimetric biosensing of H<sub>2</sub>O<sub>2</sub> and glucose, *Sens. Actuators, B*, 2016, **229**, 379–386.
- 30 Y. X. Dong, J. T. Cao, B. Wang, S. H. Ma and Y. M. Liu, Spatial-resolved photoelectrochemical biosensing array based on a cds@g-C<sub>3</sub>N<sub>4</sub> heterojunction: a universal immunosensing platform for accurate detection, *ACS Appl. Mater. Interfaces*, 2018, **10**(4), 3723–3731.
- 31 Y. F. Zhang, R. Xu, Q. Kang, Y. Zhang, Q. Wei, *et al.*, Ultrasensitive photoelectrochemical biosensing platform for detecting n-terminal pro-brain natriuretic peptide based on SnO<sub>2</sub>/SnS<sub>2</sub>/mpg-C<sub>3</sub>N<sub>4</sub> amplified by PbS/SiO<sub>2</sub>, *ACS Appl. Mater. Interfaces*, 2018, **10**(37), 31080–31087.
- 32 X. L. Fu, F. Hou, F. R. Liu, S. W. Ren, J. T. Cao, *et al.*, Electrochemiluminescence energy resonance transfer in 2d/2d heterostructured g-C<sub>3</sub>N<sub>4</sub>/MnO<sub>2</sub> for glutathione detection, *Biosens. Bioelectron.*, 2019, **129**, 72–78.
- 33 X. F. Zhao, P. K. Panda, D. Singh, X. Y. Yang, Y. K. Mishra, *et al.*, 2d g-C<sub>3</sub>N<sub>4</sub> monolayer for amino acids sequencing, *Appl. Surf. Sci.*, 2020, **528**, 146609.
- 34 G. Zuo, X. Zhou, Q. Huang, H. Fang and R. Zhou, Adsorption of villin headpiece onto graphene, carbon nanotube, and C60: effect of contacting surface curvatures on binding affinity, *J. Phys. Chem. C*, 2011, **115**, 23323–23328.
- 35 X. C. Zhao, Self-assembly of DNA segments on graphene and carbon nanotube arrays in aqueous solution: a molecular simulation study, *J. Phys. Chem. C*, 2011, **115**(14), 6181–6189.
- 36 Y. Tu, M. Lv, P. Xiu, T. Huynh, M. Zhang, *et al.*, Destructive extraction of phospholipids from *Escherichia coli* membranes by graphene nanosheets, *Nat. Nanotechnol.*, 2013, **8**(8), 594–601.
- 37 B. Li, W. Li, J. M. Perez-Aguilar and R. Zhou, Mild binding of protein to C<sub>2</sub>N monolayer reveals its suitable biocompatibility, *Small*, 2017, **13**(12), 1603685.
- 38 Z. Gu, L. Zhao, S. Liu, G. Duan, J. M. Perez-Aguilar, *et al.*, Orientational binding of DNA guided by the C<sub>2</sub>N template, *ACS Nano*, 2017, **11**(3), 3198–3206.
- 39 Z. L. Gu, Z. X. Yang, L. L. Wang, H. Zhou, C. A. Jimenez-Cruz, *et al.*, The role of basic residues in the adsorption of blood proteins onto the graphene surface, *Sci. Rep.*, 2015, **5**, 10873.
- 40 H. Y. Cui, Z. L. Gu, X. C. Chen, L. Lin, Z. G. Wang, *et al.*, Stimulating antibacterial activities of graphitic carbon nitride nanosheets with plasma treatment, *Nanoscale*, 2019, **11**(39), 18416–18425.
- 41 Z. Gu, W. Song, S. H. Chen, B. Y. Li, W. F. Li, *et al.*, Defect-assisted protein hp35 denaturation on graphene, *Nanoscale*, 2019, **11**(41), 19362–19369.
- 42 L. J. Beamer and C. O. Pabo, Refined 1.8 crystal structure of the λ repressor-operator complex, *J. Mol. Biol.*, 1992, **227**(1), 177–196.
- 43 J. Guo, X. Yao, L. Ning, Q. Wang and H. Liu, The adsorption mechanism and induced conformational changes of three typical proteins with different secondary structural features on graphene, *RSC Adv.*, 2014, **4**(20), 9953–9962.
- 44 B. Hess, C. Kutzner, D. van der Spoel and E. Lindahl, Gromacs 4: Algorithms for highly efficient, load-balanced, and scalable molecular simulation, *J. Chem. Theory Comput.*, 2008, **4**(3), 435–447.
- 45 W. Humphrey, A. Dalke and K. Schulten, Vmd: Visual molecular dynamics, *J. Mol. Graphics Modell.*, 1996, **14**(1), 33–38.
- 46 J. Huang and A. D. MacKerell Jr, Charmm36 all-atom additive protein force field: validation based on comparison to NMR data, *J. Comput. Chem.*, 2013, **34**(25), 2135–2145.
- 47 W. L. Jorgensen, J. Chandrasekhar, J. D. Madura, R. W. Impey and M. L. Klein, Comparison of simple



- potential functions for simulating liquid water, *J. Chem. Phys.*, 1983, **79**(2), 926–935.
- 48 G. Bussi, D. Donadio and M. Parrinello, Canonical sampling through velocity rescaling, *J. Chem. Phys.*, 2007, **126**(1), 014101.
- 49 M. Parrinello and A. Rahman, Polymorphic transitions in single-crystals – a new molecular-dynamics method, *J. Appl. Phys.*, 1981, **52**(12), 7182–7190.
- 50 Z. Gu, J. Perez-Aguilar, L. Meng and R. Zhou, Partial denaturation of villin headpiece upon binding to a carbon nitride polyaniline (C<sub>3</sub>N) nanosheet, *J. Phys. Chem. B*, 2020, **124**(35), 7557–7563.
- 51 T. Darden, D. York and L. Pedersen, Particle mesh ewald - an n.Log(n) method for ewald sums in large systems, *J. Chem. Phys.*, 1993, **98**(12), 10089–10092.
- 52 U. Essmann, L. Perera, M. L. Berkowitz, T. Darden, H. Lee, *et al.*, A smooth particle mesh ewald method, *J. Chem. Phys.*, 1995, **103**(19), 8577–8593.
- 53 B. Hess, H. Bekker, H. J. C. Berendsen and J. Fraaije, Lincs: a linear constraint solver for molecular simulations, *J. Comput. Chem.*, 1997, **18**(12), 1463–1472.
- 54 S. Miyamoto and P. A. Kollman, Settle – an analytical version of the shake and rattle algorithm for rigid water models, *J. Comput. Chem.*, 1992, **13**(8), 952–962.