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Autophagy and evasion of the immune system by SARS-CoV-2. Structural features of the non-structural protein 6 from wild type and Omicron viral strains interacting with a model lipid bilayer†

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The viral cycle of SARS-CoV-2 is based on a complex interplay with the cellular machinery, which is mediated by specific proteins eluding or hijacking the cellular defense mechanisms. Among the complex pathways induced by the viral infection, autophagy is particularly crucial and is strongly influenced by the action of the non-structural protein 6 (Nsp6) interacting with the endoplasmic reticulum membrane. Importantly, differently from other non-structural proteins, Nsp6 is mutated in the recently emerged Omicron variant, suggesting a possible different role of autophagy. In this contribution we explore, for the first time, the structural properties of Nsp6 thanks to long-timescale molecular dynamics simulations and machine learning analysis, identifying the interaction patterns with the lipid membrane. We also show how the mutation brought by the Omicron variant may indeed modify some of the specific interactions, and more particularly help anchor the viral protein to the lipid bilayer interface.

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Introduction

The emergence in late 2019 of a novel positive sense, single-stranded RNA β -coronavirus, shortly afterwards named SARS-CoV-2, has led to the outbreak of a pandemic, COVID-19, which has since then severely affected every country worldwide. Indeed, and despite its relatively low mortality ratio, the high transmissibility of SARS-CoV-2,^{1–3} coupled with the potential development of serious outcomes requiring intensive care treatment,⁴ has resulted in a considerable strain posed on health systems and has led to serious social distancing and containment measures, including full lockdowns. Though the development and the large-scale deployment, at least in Western countries, of efficient vaccines,^{5–7} including the revolutionary mRNA vaccines,^{8–11} has allowed a much better containment of the pandemic and a significant decrease in both

deaths and intensive care admissions, SARS-CoV-2 remains a considerable and not fully mastered threat. Indeed, the year 2021 has been characterized by the emergence of different SARS-CoV-2 variants, classified as variants of concern (VOCs) by the World Health Organization (WHO), which due to their higher transmissibility have rapidly become dominant, replacing the original viral strains, even if the mutation rate of SARS-CoV-2 is much slower compared to other RNA viruses, thanks to the presence of an exonuclease acting as a proofreading tool during viral genome replication. After the emergence first of Alpha¹² and Beta¹³ (beginning of 2021) and Delta variants¹⁴ (summer 2021), a novel strain, styled Omicron and accumulating a high density of point mutations, was reported in Southern African countries on the 25th of November 2021.¹⁵ The Omicron variant,^{16–19} also called B.1.1.529, is characterized by both a higher transmissibility and the partial capacity to infect subjects with prior immunity obtained either *via* vaccination or precedent infection.²⁰ Despite some preliminary data appearing to point to a lower severity rate of Omicron compared to the original variant,²¹ its high transmissibility and the partial evasion of precedent immunization constitute a drastic problem.²²

From a molecular biology point of view,^{23,24} the SARS-CoV-2 genome is constituted by a large, ~ 30 k base, positive-sense single-stranded RNA fragment, which is enveloped in a membrane virion. After cellular infection the viral genome is translated into two large polyproteins, PP1 and PP2, and some structural proteins, such as spike (S, responsible for the

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interaction with the cellular receptor and the membrane fusion, and concentrating most of the Omicron mutations),²⁵ nuclear (N) and envelope (E) proteins. In turn the original PPs are self-cleaved by two proteases giving rise to the so-called non-structural proteins (Nsps), which are responsible for crucial viral processes related to its replication and resistance to the host immune system.^{26,27} Indeed, among the non-structural proteins one should cite, in addition to the proteases, the RNA-dependent RNA polymerase^{28,29} which is responsible for the genome replication, *via* a temporary negative-stranded RNA template, the exonuclease complex,³⁰ and the SARS-Unique Domain (SUD) which may sequester RNA to impede triggering apoptotic signals.^{31,32} Furthermore, membrane Nsps are also present and tend to accumulate in the endoplasmic reticulum (ER), *i.e.* the replication compartment of SARS-CoV-2. Nsp6 has also been recognized as being capable of interfering with the type I interferon pathway by blocking the activation of Tank binding kinase allowing viral evasion of innate immune response.³³ Nsp6 has also been reported to be an important regulator of autophagy in infected cells.^{34,35} As a matter of fact, coronavirus-infected cells present a higher number of autophagosomes, the latter being much smaller than in non-infected cells, a phenomenon in which the Nsp6 protein plays a crucial role.³⁴ These different roles of NSP6 reflect the complex equilibrium between immune response and viral replication.³⁶ Interfering with different steps of autophagy favors viral survival and propagation.³⁷ Indeed, while autophagosomes incorporating exogenous and endogenous protein material may actively participate in the elimination/destruction of viral components and hence enhance adaptive immunity by delivering viral antigens, they may also form protective compartments in which viral replication can take place using autophagic products such as metabolites and substrates. Reducing the size of the autophagosome may, thus, hamper the fusion of autophagosomes with lysosomes, preventing the elimination of the viral material while maintaining a favorable environment for viral replication and maturation.³⁵

The structure of many key SARS-CoV-2 structural and non-structural proteins has been resolved,^{28,38,39} from the first day of the pandemic, and the relation between their structure and activity has since also been complemented by multiscale molecular modeling and simulation,^{24,31,40–42} also tackling enzymatic reactivity.^{26,27} Undoubtedly, the S protein, also in complex with the human ACE2 receptor, and the viral proteases have been the main target for structural biology and molecular modeling simulations.^{43–47} Of note, the proposition of some possible viral inhibitors has also been undertaken with some success. In contrast, the structure, and hence the mechanisms, of other Nsps including Nsp6 (ref. 33) has been much less studied despite its fundamental biological role. Notably, while no experimentally resolved structure of membrane-embedded Nsp6 is available, the combination of sequence homology and machine learning approaches⁴⁸ has allowed a putative starting structure to be proposed. In this contribution we aim at filling this gap validating the proposed Nsp6 structure through extended all-atom molecular dynamics (MD) simulation, identifying the key structural motifs that allow an efficient

interaction with the lipid bilayer. Our results are also consistent with those of Kumar *et al.* for the WT strain.⁴⁹

The SARS-CoV-2 variants exhibit a high density of mutations, mainly concentrated on the spike coding sequence, which influence greatly the binding to the human ACE2 receptor and its viral invasion capacity.⁵⁰ Interestingly, the Omicron variant also presents the deletion of three amino acids, namely L105, S106 and G107, from the Nsp6 sequence.²⁵ The three deleted amino acids are located at the polar head/water interface where they connect, *via* a distorted loop, two transmembrane α -helices. Their absence can clearly influence the protein/membrane interaction and hence have a non-negligible role in autophagy.⁵¹ Hence, in the following we provide MD simulations on the mutated protein for comparison with those originating from the native strain.

In addition to rationalizing the dynamical properties of Nsp6 and its interaction with lipid bilayers, our results also tackle the effects of the mutations of the Omicron variant in a crucial protein responsible for virus maturation and immune system elusion.

Results

As shown in Fig. 1 the structure of the Wild Type (WT) Nsp6 presents all the key features of typical transmembrane proteins: transmembrane helical bundles and extramembranous connectors. Furthermore, the structure of the protein retrieved by machine learning approaches is found to be remarkably stable. Indeed, throughout the 2 μ s-long MD simulation of Nsp6 embedded in a lipid bilayer only very limited and local deviation from the initial structure is observed. This is evidenced by the time series of the root mean square deviation (RMSD) of the

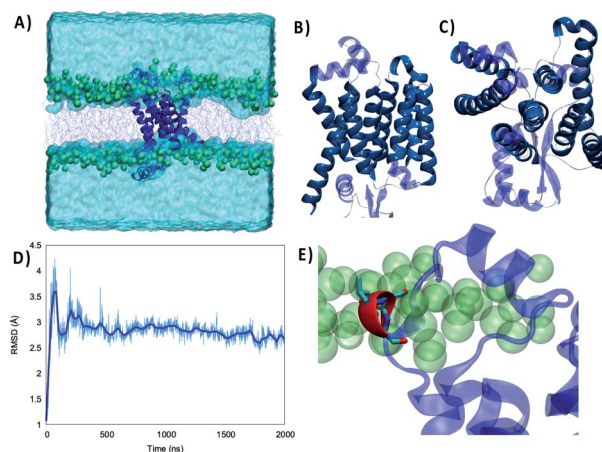


Fig. 1 (A) Representation of the simulation box showing the WT Nsp6 protein embedded in a model lipid bilayer surrounded by water buffer. Side (B) and top (C) views of the Nsp6 protein highlighting the secondary structure motifs. The transmembrane regions are represented in darker blue and opaque, while extramembranous areas are rendered in lighter blue and transparent. (D) Time series of the RMSD for Nsp6 from the MD simulation. (E) Zoomed-in view of the three amino acids which are deleted in the Omicron variant represented in red and licorice.



protein which after an equilibration period of about 200–330 ns exhibits an extended plateau to reach an average value of 2.88 ± 0.01 Å. Remarkably, even during the first equilibration period the maximum value of the RMSD barely exceeds 4 Å confirming the global stability of the protein structure and its favorable interaction with the lipid bilayer. Besides, the membrane structural parameters also do not exhibit remarkable fluctuations during the production run, confirming the stability of the overall system – see Fig. S1†. Hence, our results can also be regarded as a first independent confirmation of the proposed structure of Nsp6. From the analysis of the protein's secondary structure during the MD simulation we identified some crucial features, which deserve discussion. As shown in Fig. 1B and C, we identified 8 rather long transmembrane α -helices (TM1 to TM8), which are mainly composed of hydrophobic residues and which assume a clear transmembrane bundle structure, which is a widespread pattern in membrane proteins (see for instance rhodopsin and other receptors). Despite their slightly different length and some minor changes in their orientation with respect to the membrane axis, all eight α -helices are remarkably stable in terms of both their secondary and tertiary structures and exhibit only negligible displacements during the simulation. Once again, the rigidity of the transmembrane core may be regarded as a typical feature of stable membrane proteins. On top of the transmembrane core one can also identify six shorter and more flexible α -helices and 2 β -sheets extending at the interface between the lipid polar head and water. Interestingly, these interfacial structures tend to be positioned in a parallel conformation with respect to the membrane plane, and the β -helices are disposed almost symmetrically on the two membrane/water interfaces.

Differently from the transmembrane core structure, these shorter motifs present a higher density of polar residues, consistent with their positioning at the polar head region. Furthermore, some transient electrostatic interactions, as well as hydrogen bonds, are formed between the phosphate or the choline moieties of the lipids and the interfacial amino acids. However, the interaction network is highly dynamic and evolves continuously throughout the simulation without showing a dominant pattern. This fact may also partially contribute to justifying the higher flexibility of the extramembranous regions as compared to the core. These considerations are supported by an analysis of the secondary structure along the trajectory (Fig. S2†), showing the stability of the eight transmembrane helices and of the two isolated β -sheets, while all extramembranous α -helices are more flexible (especially between TM3 and TM4 and after TM8 toward the –C terminus), due to bending, turning and, more generally, the presence of non-structured short linkers.

Of particular interest (see Fig. 1E) is a very short α -helix composed of a triad of residues, namely L105, S106 and G107. Indeed, while the secondary structure is stable throughout the MD simulation, this triad can be found inside an unstructured loop connecting the transmembrane core to the extramembranous α -helix formed by residues 89 to 99. This short helix is one of the most flexible and mobile moieties of Nsp6 and experiences significant oscillation on the membrane plane.

Furthermore, L105, S106 and G107 are also the amino acids that are deleted from Nsp6 in the Omicron variant. Hence, this structural motif and the nearby areas may experience greater variability among the different strains.

For this reason, we performed an independent MD simulation of the Omicron Nsp6 variant, which has been obtained by manually mutating the WT strain. The results of the MD simulations are shown in Fig. 2. From the analysis of the MD simulation it is apparent that the Omicron variant is still able to favorably interact with the lipid bilayer producing a stable aggregate (Fig. 2A). Noteworthy, the structural parameters of the membrane (*i.e.*, thickness, position of the membrane center on the Z axis and area per lipid) remain stable during the simulation for both variants (Fig. S1†). Furthermore, as evidenced by the time evolution of the RMSD, we may see that the global deformation of the protein is still rather small, and more importantly after a first equilibration region where it reaches 3.5/4.0 Å it stabilizes to an extended plateau at 3.0 Å. Consistent with what was observed for the WT Nsp6 while the transmembrane core is extremely rigid and experiences barely any fluctuation over the course of the MD simulation, the peripheral helices present a greater mobility and flexibility. Hence, the peripheral motifs will also show a higher difference between the WT and Omicron variant, as pictorially shown in Fig. 2C and D where representative snapshots for the two variants are superimposed. As expected, the peripheral α -helix formed between the 89th and 99th residues, *i.e.* the region between TM3 and TM4, is indeed the one showing the larger deviation between the two structures (see Fig. S2†). As a matter of fact, the mutation in the Omicron strain involves the short loop connecting this helix to the protein core, hence justifying the higher variability of its structure. These global tendencies are also confirmed by the analysis of the flexibility profile at the residue level (Fig. 2E) and by the root mean square fluctuation (RMSF, reported in Fig. S3†). Indeed, most of the protein, and in particular the transmembrane core, presents a very low

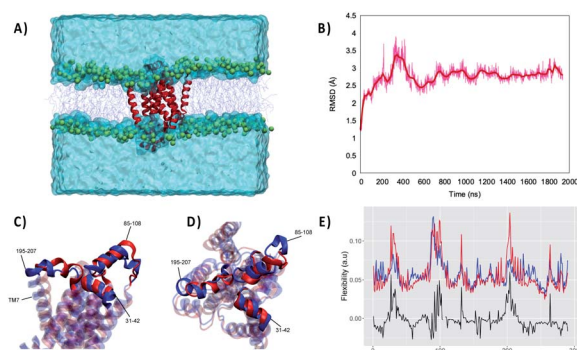


Fig. 2 (A) A snapshot of the full simulation box showing the stability of the interaction between Omicron Nsp6 and the lipid membrane. (B) The structural stability of the latter also revealed by the time series of the RMSD for the protein. Side (C) and top (D) views of the superposition of WT (blue) and Omicron (red) Nsp6; the extramembranous helices and the mobile transmembrane helix 7 (TM7) are evidenced. (E) Per amino acid flexibility profiles of the WT (blue) and Omicron (red) variants; the point-to-point difference is also reported in black.



flexibility, with the partial exception of TM7, which experiences considerable turning events in its $-C$ terminal region (see Fig. S2†). As a matter of fact, two peaks can be observed corresponding to residues 86–108 and 195–207 for the WT. These two regions represent two transmembrane helices protruding in the polar head region. Furthermore, the first flexible peak also encompasses the amino acid triad which is deleted in the Omicron variant. As for the comparison with the Omicron variant we observe that while the rigidity profile is similar to that of the WT a rather larger flexibility is observed corresponding to the two peaks. This may be related to the necessity of rearranging the local structure of the protein to account for the effects of the mutation.

To better analyze the global effects of the different flexibility and organization around the membrane we report in Fig. 3 the evolution of the relative position of the center of mass of the 89–99 helix with respect to the center of mass of the lipid bilayer for the two variants, evidencing a slightly different behavior of the two proteins. Indeed, concerning the projection of the distance along the (x,y) plane, *i.e.* the one parallel to the membrane, we see that the WT peaks around the $(-8.0, 0.0)$ Å position while the Omicron variant concentrates at $(-9.0, 0.0)$ Å. More importantly, the 2D-distribution shows a different topology between the two variants; the Omicron strain evidences a bimodal distribution with the presence in addition to the main peak of a metastable state at $(-13.0, -6.0)$ Å spanning for a couple of hundreds of ns. In contrast, the WT shows a more pronounced variability with the distribution of distances covering an extended region prior to its stabilization to the final stable state.

The analysis of the projection of the distance on the z axis, *i.e.* the one perpendicular to the membrane, is perhaps even

more significant and interesting. As shown by its time series reported in Fig. 3B, we evidence two very distinct behaviors between the two proteins. Indeed, in the case of the Omicron strain the peripheral helix is buried deeper inside the polar head region and assumes an average distance from the center of the bilayer of 24.8 ± 2.0 Å, compared with 27.7 ± 2.0 Å for the WT. Furthermore, even if less evident from the standard deviation, in the case of the WT we evidence the oscillation between a buried (*i.e.* shorter distance) and an exposed (*i.e.* longer distance) conformation, which translates to a rather broad distribution (see Fig. S4†). The oscillation period between the two WT conformations appears to be of the order of 400 ns, even though longer trajectories and a more extended sampling would be necessary to provide a quantitative estimation.

An analysis of the interactions between the extra-membranous α -helix, containing the triad of residues deleted in the Omicron variant, and the surrounding membrane was performed by measuring the distance between the centers of mass of each residue and the center of mass of the polar heads found in a radius of 60 Å from the Leu105–Gly107 triad (Fig. S5†). As can be seen, such deletion induced a consistent strengthening of the protein–membrane interaction, with an average distance that decreases from *ca.* 30 to *ca.* 15 Å, with charged Lys109 interacting at even shorter distances (peak at around 10 Å). Thus, we can state on firm ground that the Omicron variant clearly favors a more deeply buried conformation of this α -helix.

A similar behavior can also be observed for the 195–207 helix (Fig. S6†), which has also been identified as a flexibility hot-spot, and which is globally more deeply buried in the case of the Omicron variant, presenting an average distance from the center of the membrane of 15.6 ± 1 Å compared to the 17.3 ± 2 Å observed for the WT. Furthermore, the projection of the distance on the (x,y) plane is also significantly different between the two variants with the WT peaking at $(10.0, -1.0)$ Å and the Omicron variant at $(6.0, -1.5)$ Å as shown in Fig. S6†.

Conclusions and discussion

Nsp6 is a crucial protein in the SARS-CoV-2 viral cycle. In particular, its capacity to modulate the autophagy response and the autophagosome topology is fundamental in regulating the delicate equilibrium between the establishment of favorable replication conditions and an efficient immune system response. However, its structure and its interaction with lipid bilayers has not been reported yet. In this contribution we fill this gap by validating the proposed Nsp6 structure using large-scale MD simulations and confirming the stability of the protein while identifying the crucial interaction pattern established with a model lipid bilayer.

Furthermore, the mutation of Nsp6 in the Omicron variant, consisting in the deletion of a triad of amino acids at the polar head interface, has been pinpointed as a potential concerning mutation due to its possibility of altering the modulation of autophagy by favoring the interaction with the membrane. Indeed, the molecular basis underlying autophagy will be due to the recruitment of lipid, specifically PIP2, by Nsp6 and the

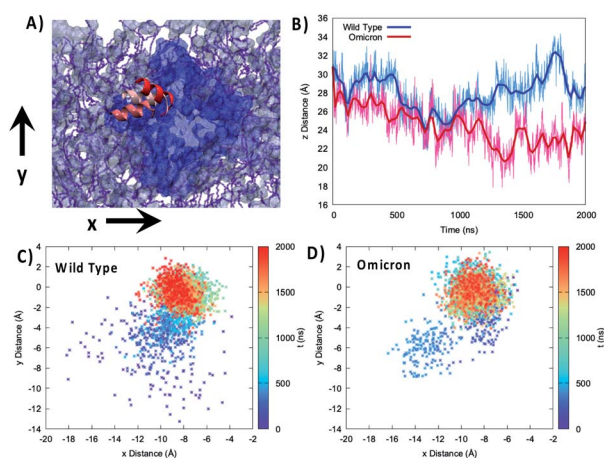


Fig. 3 (A) Representation of the mobility of the 89–99 α -helix by three different snapshots, shown with different reddish shades, reported on top of the membrane and protein on the xy plane. (B) Time series of the distance between the center of mass of the 85–108 helix and the lipid membrane projected on the z -axis parallel to the bilayer thickness for the WT (blue) and Omicron variant (red). Projection on the xy plane of the distance between the center of mass of the 85–108 helix and the lipid membrane for the WT (C) and Omicron variant (D). The time series is given as a color map. The histograms of the distance distribution at various time intervals are also reported in Fig. S7†.



subsequent formation of the autophagosome vesicle. Hence, any mutation which would favor the interaction with the membrane is prone to alter this mechanism. While we have shown that the global structure of Nsp6 is not altered by the triad deletion, in particular concerning the transmembrane core, we have evidenced an important modification of the structure and dynamic of the peripheral helices, as illustrated by flexibility signature perturbation. More specifically, we have shown that the peripheral helices in the Omicron variant are more deeply buried in the polar head region compared to those of the WT, hence suggesting stronger interaction and an increased capacity of recruiting specific lipids. Importantly, the structure of the POPC membrane does not show strong deviations between the two variants. Yet, further study of the complex effect of the membrane's composition on its mechanical properties would provide a clearer picture of the interplay between the membrane structure and the biological aspects of the viral cycle. This would also include the study of the effects of lipids such as phosphatidylinositol-4,5-bisphosphate (PIP2) which are known to play an important role in the autophagosome formation.⁵² However, while our results should be confirmed experimentally, they point to a different modulation of the autophagy between the two viral variants which may in part explain both the immune system resistance of the Omicron variant and its different pathological evolution. In the near future we plan to pursue this study on the one side explicitly including PIP2 in our simulations and on the other side experimentally comparing the induction of autophagy in cell lines in which WT or Omicron Nsp6 would have been expressed.

Methodology

The initial proposed structure of SARS-CoV-2 Nsp6 has been retrieved from Jumper *et al.*'s Deepmind web server⁴⁸ collecting putative structures of COVID-19 related proteins obtained *via* the AlphaFold machine learning methodology⁵³ fed by the sequences taken from the UniProt pre-release download.⁵⁴ The structure of the Omicron variant has been manually constructed from the WT by deleting the L105, S106, and G107 triad. The protocol used to construct the Omicron variant has been justified, by the AlphaFold-based⁵⁵ structural prediction which confirms that the deletion of the amino acid triad does not alter the global structure of the transmembrane α -helix bundle. Both WT and Omicron Nsp6 have been embedded in a lipid membrane bilayer composed of two leaflets of 100 1-palmitoyl-2-oleoylphosphatidylcholine (POPC) lipids, and a water buffer has been added complemented with a physiological concentration of K^+ and Cl^- ions. Although cellular membranes are clearly more complex and present variable concentrations of lipid species, including steroids, which can alter their rigidity, single lipid membranes may be considered as valid models of a biologically relevant lipid bilayer, and have already been used in different computational studies.^{56,57} The solvated membrane/protein structure has been prepared using the Charmm-gui web-server interface.⁵⁸ The lipid and the protein have been represented by the Amber ff14 (ref. 59 and 60) force field while water has been modeled with TIP3P.⁶¹ MD

simulations have been performed using the NAMD code,^{62,63} using periodic boundary conditions (PBCs) and the constant number of particle, pressure, and temperature (NPT) ensemble, using a Langevin thermostat and barostat.^{64,65} A time step of 4 fs has been used to integrate the Newton equation of motion thanks to the use of Hydrogen Mass Repartitioning (HMR)⁶⁶ in combination with RATTLE and SHAKE.⁶⁷ After minimization, thermalization and equilibration have been performed, progressively releasing harmonic constraints on the protein backbone and the lipid over 40 ns, after which a 2 μ s production run has been obtained for both the WT and the Omicron systems. All the trajectories have been visualized and analyzed with the VMD code.⁶⁸ The per residue Nsp6 flexibility profile was obtained through an in-house machine learning protocol, based on the one proposed by Fleetwood and coworkers,⁶⁹ considering the principal component analysis (PCA) of the protein based on the decomposition of the RMSD covariance matrix. In particular, we use the internal coordinates (inverse distance between the geometric centers of two residues) along the trajectories as the input of PCA instead of the cartesian coordinates to provide better performance. The per residue importance (*i.e.* flexibility) was calculated by taking the sum of the weights of the principal components up to 80%, where the weight is defined as the eigenvalue of the corresponding principal component over the sum of all the eigenvalues of the RMSD covariance matrix. The secondary structure analysis of both the WT and Omicron variant was performed by applying the Define Secondary Structure of Proteins (DSSP) algorithm,⁷⁰ as implemented in the AmberTools21 suite of programs.⁷¹

Data availability

Raw data are available upon request.

Author contributions

Conceptualization, planning and data collection: A. M. Data analysis and discussion: A. M., E. B., M. M., and S. G. All the authors contributed to the draft realization and writing.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 X. Tang, C. Wu, X. Li, Y. Song, X. Yao, X. Wu, Y. Duan, H. Zhang, Y. Wang, Z. Qian, J. Cui and J. Lu, *Natl. Sci. Rev.*, 2020, **7**, 1012–1023.
- 2 Y. C. Liu, R. L. Kuo and S. R. Shih, *Biomed. J.*, 2020, **43**, 328–333.
- 3 I. Astuti and Ysrafil, *Diabetes Metab. Syndr.: Clin. Res. Rev.*, 2020, **14**, 407–412.
- 4 J. L. Domingo, M. Marquès and J. Rovira, *Environ. Res.*, 2020, **188**, 109861.
- 5 M. Sadarangani, A. Marchant and T. R. Kollmann, *Nat. Rev. Immunol.*, 2021, **21**, 475–484.
- 6 F. X. Heinz and K. Stiasny, *npj Vaccines*, 2021, **6**, 1–13.
- 7 T. Fiolet, Y. Kherabi, C.-J. MacDonald, J. Ghosn and N. Peiffer-Smadja, *Clin. Microbiol. Infect.*, 2022, **28**, 202–221.
- 8 F. P. Polack, S. J. Thomas, N. Kitchin, J. Absalon, A. Gurtman, S. Lockhart, J. L. Perez, G. Pérez Marc, E. D. Moreira, C. Zerbini, R. Bailey, K. A. Swanson, S. Roychoudhury, K. Koury, P. Li, W. V. Kalina, D. Cooper, R. W. Frenck, L. L. Hammitt, Ö. Türeci, H. Nell, A. Schaefer, S. Ünal, D. B. Tresnan, S. Mather, P. R. Dormitzer, U. Şahin, K. U. Jansen and W. C. Gruber, *N. Engl. J. Med.*, 2020, **383**, 2603–2615.
- 9 E. J. Rubin and D. L. Longo, *N. Engl. J. Med.*, 2022, **386**, 183–185.
- 10 Q. Huang, J. Zeng and J. Yan, *J. Genet. Genomics*, 2021, **48**, 107–114.
- 11 T. Charmet, L. Schaeffer, R. Grant, S. Galmiche, O. Chény, C. Von Platen, A. Maurizot, A. Rogoff, F. Omar, C. David, A. Septfons, S. Cauchemez, A. Gaymard, B. Lina, L. H. Lefrançois, V. Enouf, S. van der Werf, A. Mailles, D. Levy-Bruhl, F. Carrat and A. Fontanet, *The Lancet Regional Health–Europe*, 2021, **8**, 100171.
- 12 P. Domingo and N. de Benito, *EBioMedicine*, 2021, **74**, 103703.
- 13 G. Cosentino, M. Bernard, J. Ambroise, J. M. Giannoli, J. Guedj, F. Débarre and F. Blanquart, *J. Infect.*, 2021, **84**, 94–118.
- 14 R. Thiruvengadam, A. Binayke and A. Awasthi, *Lancet Infect. Dis.*, 2021, **22**, 301–302.
- 15 S. S. A. Karim and Q. A. Karim, *Lancet*, 2021, **398**, 2126–2128.
- 16 V. M. Ferré, N. Peiffer-Smadja, B. Visseaux, D. Descamps, J. Ghosn and C. Charpentier, *Anaesth. Crit. Care Pain Med.*, 2021, 100998.
- 17 V. Thakur and R. Kanta Ratho, *J. Med. Virol.*, 2022, **94**, 1821–1824.
- 18 J. R. C. Pulliam, C. van Schalkwyk, N. Govender, A. von Gottberg, C. Cohen, M. J. Groome, J. Dushoff, K. Mlisana and H. Moultrie, *Science*, 2022, DOI: [10.1101/2021.11.11.21266068](https://doi.org/10.1101/2021.11.11.21266068).
- 19 X. He, W. Hong, X. Pan, G. Lu and X. Wei, *MedComm*, 2021, **2**, 838–845.
- 20 L. Zhang, Q. Li, Z. Liang, T. Li, S. Liu, Q. Cui, J. Nie, Q. Wu, X. Qu, W. Huang and Y. Wang, *Emerging Microbes Infect.*, 2022, **11**, 1–5.
- 21 N. Wolter, W. Jassat, S. Walaza, R. Welch, H. Moultrie, M. Groome, D. G. Amoako, J. Everatt, J. N. Bhiman, C. Scheepers, N. Tebeila, N. Chiwandire, M. du Plessis, N. Govender, A. Ismail, A. Glass, K. Mlisana, W. Stevens, F. K. Treurnicht, Z. Makatini, N. Hsiao, R. Parboosing, J. Wadula, H. Hussey, M.-A. Davies, A. Boulle, A. von Gottberg and C. Cohen, *The Lancet*, 2022, **399**, 437–446.
- 22 S. K. Saxena, S. Kumar, S. Ansari, J. T. Paweska, V. K. Maurya, A. K. Tripathi and A. S. Abdel-Moneim, *J. Med. Virol.*, 2022, **94**, 1738–1744.
- 23 A. A. T. Naqvi, K. Fatima, T. Mohammad, U. Fatima, I. K. Singh, A. Singh, S. M. Atif, G. Hariprasad, G. M. Hasan and M. I. Hassan, *Biochim. Biophys. Acta, Mol. Basis Dis.*, 2020, **1866**, 165878.
- 24 A. Francés-Monerris, C. Hognon, T. Miclot, C. García-Iriepa, I. Iriepa, A. Terenzi, S. Grandemange, G. Barone, M. Marazzi and A. Monari, *J. Proteome Res.*, 2020, **19**, 4291–4315.
- 25 L. Wang and G. Cheng, *J. Med. Virol.*, 2022, **94**, 1728–1733.
- 26 C. A. Ramos-Guzmán, J. J. Ruiz-Pernía and I. Tuñón, *ACS Catal.*, 2020, **10**, 12544–12554.
- 27 K. Świderek and V. Moliner, *Chem. Sci.*, 2020, **11**, 10626–10630.
- 28 A. A. Elfiky, *J. Biomol. Struct. Dyn.*, 2021, **39**, 3204–3212.
- 29 H. S. Hillen, G. Kokic, L. Farnung, C. Dienemann, D. Tegunov and P. Cramer, *Nature*, 2020, **584**, 154–156.
- 30 F. Robson, K. S. Khan, T. K. Le, C. Paris, S. Demirbag, P. Barfuss, P. Rocchi and W.-L. Ng, *Mol. Cell*, 2020, **80**, 1136–1138.
- 31 C. C. Hognon, T. Miclot, C. G. Iriepa, A. Francés-Monerris, S. Grandemange, A. Terenzi, M. Marazzi, G. Barone, A. Monari, C. Garcia-Iriepa, A. Francés-Monerris, S. Grandemange, A. Terenzi, M. Marazzi, G. Barone and A. Monari, *J. Phys. Chem. Lett.*, 2020, **11**, 5661–5667.
- 32 J. Tan, Y. Kusov, D. Mutschall, S. Tech, K. Nagarajan, R. Hilgenfeld and C. L. Schmidt, *Biochem. Biophys. Res. Commun.*, 2007, **364**, 877–882.
- 33 H. Xia, Z. Cao, X. Xie, X. Zhang, J. Y. C. Chen, H. Wang, V. D. Menachery, R. Rajsbaum and P. Y. Shi, *Cell Rep.*, 2020, **33**, 108234.
- 34 E. M. Cottam, M. C. Whelband and T. Wileman, *Autophagy*, 2014, **10**, 1426–1441.
- 35 W. W. Y. Yim and N. Mizushima, *Dev. Cell*, 2021, **56**, 400–402.
- 36 J. Mao, E. Lin, L. He, J. Yu, P. Tan and Y. Zhou, in *Advances in Experimental Medicine and Biology*, Springer, Singapore, 2019, vol. 1209, pp. 55–78.
- 37 Y. Choi, J. W. Bowman and J. U. Jung, *Nat. Rev. Microbiol.*, 2018, **16**, 341–354.



- 38 J. Lan, J. Ge, J. Yu, S. Shan, H. Zhou, S. Fan, Q. Zhang, X. Shi, Q. Wang, L. Zhang and X. Wang, *Nature*, 2020, **581**, 215–220.
- 39 A. Dömling and L. Gao, *Chem*, 2020, **6**, 1283–1295.
- 40 L. Casalino, Z. Gaieb, J. A. Goldsmith, C. K. Hjorth, A. C. Dommer, A. M. Harbison, C. A. Fogarty, E. P. Barros, B. C. Taylor, J. S. McLellan, E. Fadda and R. E. Amaro, *ACS Cent. Sci.*, 2020, **6**, 1722–1734.
- 41 C. García-Iriepa, C. Hognon, A. Francés-Monerris, I. Iriepa, T. Miclot, G. Barone, A. Monari and M. Marazzi, *J. Phys. Chem. Lett.*, 2020, **11**, 9272–9281.
- 42 A. Spinello, A. Saltalamacchia and A. Magistrato, *J. Phys. Chem. Lett.*, 2020, **11**, 4785–4790.
- 43 Y. Zhang, N. Zheng, P. Hao, Y. Cao and Y. Zhong, *Comput. Biol. Chem.*, 2005, **29**, 254–257.
- 44 R. Yan, Y. Zhang, Y. Li, L. Xia, Y. Guo and Q. Zhou, *Science*, 2020, **367**, 1444–1448.
- 45 M. Amin, M. Amin, M. Amin, M. K. Sorour and A. Kasry, *J. Phys. Chem. Lett.*, 2020, **11**, 4897–4900.
- 46 A. C. Henry, *C&EN Global Enterprise*, 2020, **98**, 24–25.
- 47 S. Chakraborti, P. Prabakaran, X. Xiao and D. S. Dimitrov, *Virol. J.*, 2005, **2**, 73.
- 48 K. T. John Jumper, A. T. suvunakool, P. Kohli and D. Hassabis, *Deep. Res.*, 2020, <https://deepmind.com/research/open-source/computat>.
- 49 A. Kumar, P. Kumar, K. U. Saumya and R. Giri, *Microb. Pathog.*, 2021, **161**, 105236.
- 50 M. Koehler, A. Ray, R. A. Moreira, B. Juniku, A. B. Poma and D. Alsteens, *Nat. Commun.*, 2021, **12**, 6977.
- 51 D. Benvenuto, S. Angeletti, M. Giovanetti, M. Bianchi, S. Pascarella, R. Cauda, M. Ciccozzi and A. Cassone, *J. Infect.*, 2020, **81**, e24–e27.
- 52 X. Tan, N. Thapa, Y. Liao, S. Choi and R. A. Anderson, *Proc. Natl. Acad. Sci.*, 2016, **113**, 10896–10901.
- 53 A. W. Senior, R. Evans, J. Jumper, J. Kirkpatrick, L. Sifre, T. Green, C. Qin, A. Židek, A. W. R. Nelson, A. Bridgland, H. Penedones, S. Petersen, K. Simonyan, S. Crossan, P. Kohli, D. T. Jones, D. Silver, K. Kavukcuoglu and D. Hassabis, *Nature*, 2020, **577**, 706–710.
- 54 A. Bateman, M. J. Martin, S. Orchard, M. Magrane, R. Agivetova, S. Ahmad, E. Alpi, E. H. Bowler-Barnett, R. Britto, B. Bursteinas, H. Bye-A-Jee, R. Coetzee, A. Cukura, A. Da Silva, P. Denny, T. Dogan, T. G. Ebenezer, J. Fan, L. G. Castro, P. Garmiri, G. Georgiou, L. Gonzales, E. Hatton-Ellis, A. Hussein, A. Ignatchenko, G. Insana, R. Ishtiaq, P. Jokinen, V. Joshi, D. Jyothi, A. Lock, R. Lopez, A. Luciani, J. Luo, Y. Lussi, A. MacDougall, F. Madeira, M. Mahmoudy, M. Menchi, A. Mishra, K. Moulang, A. Nightingale, C. S. Oliveira, S. Pundir, G. Qi, S. Raj, D. Rice, M. R. Lopez, R. Saidi, J. Sampson, T. Sawford, E. Speretta, E. Turner, N. Tyagi, P. Vasudev, V. Volynkin, K. Warner, X. Watkins, R. Zaru, H. Zellner, A. Bridge, S. Poux, N. Redaschi, L. Aimò, G. Argoud-Puy, A. Auchincloss, K. Axelsen, P. Bansal, D. Baratin, M. C. Blatter, J. Bolleman, E. Boutet, L. Breuza, C. Casals-Casas, E. de Castro, K. C. Echioukh, E. Coudert, B. Cuche, M. Doche, D. Dornevil, A. Estreicher, M. L. Famiglietti, M. Feuermann, E. Gasteiger, S. Gehant, V. Gerritsen, A. Gos, N. Gruaz-Gumowski, U. Hinz, C. Hulo, N. Hykounoussipikel, F. Jungo, G. Keller, A. Kerhornou, V. Lara, P. Le Mercier, D. Lieberherr, T. Lombardot, X. Martin, P. Masson, A. Morgat, T. B. Neto, S. Paesano, I. Pedruzzi, S. Pilbout, L. Pourcel, M. Pozzato, M. Pruess, C. Rivoire, C. Sigrist, K. Sonesson, A. Stutz, S. Sundaram, M. Tognolli, L. Verbregue, C. H. Wu, C. N. Arighi, L. Arminski, C. Chen, Y. Chen, J. S. Garavelli, H. Huang, K. Laiho, P. McGarvey, D. A. Natale, K. Ross, C. R. Vinayaka, Q. Wang, Y. Wang, L. S. Yeh and J. Zhang, *Nucleic Acids Res.*, 2021, **49**, D480–D489.
- 55 J. Jumper, R. Evans, A. Pritzel, T. Green, M. Figurnov, O. Ronneberger, K. Tunyasuvunakool, R. Bates, A. Židek, A. Potapenko, A. Bridgland, C. Meyer, S. A. A. Kohl, A. J. Ballard, A. Cowie, B. Romera-Paredes, S. Nikolov, R. Jain, J. Adler, T. Back, S. Petersen, D. Reiman, E. Clancy, M. Zielinski, M. Steinegger, M. Pacholska, T. Berghammer, S. Bodenstein, D. Silver, O. Vinyals, A. W. Senior, K. Kavukcuoglu, P. Kohli and D. Hassabis, *Nature*, 2021, **596**, 583–589.
- 56 M. Marazzi, H. Gattuso, A. Giussani, H. Zhang, M. Navarrete-Miguel, C. Chipot, W. Cai, D. Roca-Sanjuán, F. Dehez and A. Monari, *J. Phys. Chem. Lett.*, 2019, **10**, 7133–7140.
- 57 M. Girardon, S. Parant, A. Monari, F. Dehez, C. Chipot, E. Rogalska, N. Canilho and A. Pasc, *ChemPhotoChem*, 2019, **3**, 1034–1041.
- 58 S. Jo, T. Kim, V. G. Iyer and W. Im, *J. Comput. Chem.*, 2008, **29**, 1859–1865.
- 59 A. T. P. Carvalho and M. Swart, *J. Chem. Inf. Model.*, 2014, **54**, 613–620.
- 60 C. J. Dickson, B. D. Madej, Å. A. Skjervik, R. M. Betz, K. Teigen, I. R. Gould and R. C. Walker, *J. Chem. Theory Comput.*, 2014, **10**, 865–879.
- 61 P. Mark and L. Nilsson, *J. Phys. Chem. A*, 2001, **105**, 9954–9960.
- 62 J. C. Phillips, R. Braun, W. Wang, J. Gumbart, E. Tajkhorshid, E. Villa, C. Chipot, R. D. Skeel, L. Kalé and K. Schulten, *J. Comput. Chem.*, 2005, **26**, 1781–1802.
- 63 J. C. Phillips, D. J. Hardy, J. D. C. Maia, J. E. Stone, J. V. Ribeiro, R. C. Bernardi, R. Buch, G. Fiorin, J. Hénin, W. Jiang, R. McGreevy, M. C. R. Melo, B. K. Radak, R. D. Skeel, A. Singharoy, Y. Wang, B. Roux, A. Aksimentiev, Z. Luthey-Schulten, L. V. Kalé, K. Schulten, C. Chipot and E. Tajkhorshid, *J. Chem. Phys.*, 2020, **153**, 044130.
- 64 S. E. Feller, Y. Zhang, R. W. Pastor and B. R. Brooks, *J. Chem. Phys.*, 1995, **103**, 4613–4621.
- 65 R. L. Davidchack, R. Handel and M. V. Tretyakov, *J. Chem. Phys.*, 2009, **130**, 234101.
- 66 C. W. Hopkins, S. Le Grand, R. C. Walker and A. E. Roitberg, *J. Chem. Theory Comput.*, 2015, **11**, 1864–1874.
- 67 S. Miyamoto and P. A. Kollman, *J. Comput. Chem.*, 1992, **13**, 952–962.
- 68 W. Humphrey, A. Dalke and K. Schulten, *J. Mol. Graphics*, 1996, **14**, 33–38.
- 69 O. Fleetwood, M. A. Kasimova, A. M. Westerlund and L. Delemotte, *Biophys. J.*, 2020, **118**, 765–780.



- 70 W. Kabsch and C. Sander, *Biopolymers*, 1983, **22**, 637–2577.
- 71 D. A. Case, H. M. Aktulga, K. Belfon, I. Y. Ben-Shalom, S. R. Brozell, D. S. Cerutti, T. E. Cheatham III, G. A. Cisneros, V. W. D. Cruzeiro, T. A. Darden, R. E. Duke, G. Giambasu, M. K. Gilson, H. Gohlke, A. W. Goetz, R. Harris, S. Izadi, S. A. Izmailov, C. Jin, K. Kasavajhala, M. C. Kaymak, E. King, A. Kovalenko, T. Kurtzman, T. S. Lee, S. LeGrand, P. Li, C. Lin, J. Liu, T. Luchko, R. Luo, M. Machado, V. Man, M. Manathunga, K. M. Merz, Y. Miao, O. Mikhailovskii, G. Monard, H. Nguyen, K. A. O'Hearn, A. Onufriev, F. Pan, S. Pantano, R. Qi, A. Rahnamoun, D. R. Roe, A. Roitberg, C. Sagui, S. Schott-Verdugo, J. Shen, C. L. Simmerling, N. R. Skrynnikov, J. Smith, J. Swails, R. C. Walker, J. Wang, H. Wei, R. M. Wolf, X. Wu, Y. Xue, D. M. York, S. Zhao and P. A. Kollman, *Amber 2021*, University of California, San Francisco, p. 2021.

