

What Can AlphaFold Do for Antimicrobial Amyloids?

Running title: AlphaFold: Favoring Cross- α Predictions

Peleg Ragonis-Bachar¹, Gabriel Axel², Shahar Blau¹, Nir Ben-Tal², Rachel Kolodny³, and
Meytal Landau^{1,4,5,6*}

¹ Department of Biology, Technion-Israel Institute of Technology, Haifa 3200003, Israel

² Department of Biochemistry and Molecular Biology, George S. Wise Faculty of Life Sciences, Tel Aviv University, Tel Aviv 6997801, Israel

³ Department of Computer Science, University of Haifa, Mount Carmel, Haifa 3498838, Israel

⁴ CSSB Centre for Structural Systems Biology, Deutsches Elektronen-Synchrotron DESY, 22607 Hamburg, Germany

⁵ The Center for Experimental Medicine, Universitätsklinikum Hamburg-Eppendorf (UKE), 20251 Hamburg, Germany

⁶ European Molecular Biology Laboratory (EMBL), Hamburg, Germany

* To whom correspondence should be addressed: mlandau@technion.ac.il (M.L)

Abstract

Amyloids, protein and peptide assemblies in various organisms, are crucial in physiological and pathological processes. Their intricate structures, however, present significant challenges, limiting our understanding of their functions, regulatory mechanisms, and potential applications in biomedicine and technology. This study evaluated the AlphaFold2 ColabFold method's structure predictions for antimicrobial amyloids, using eight antimicrobial peptides (AMPs), including those with experimentally determined structures and AMPs known for their distinct amyloidogenic morphological features. Additionally, two well-known human amyloids, amyloid- β and IAPP, were included in the analysis due to their disease relevance, short sequences, and antimicrobial properties. Amyloids typically exhibit tightly mated β -strand sheets forming a cross- β configuration. However, certain amphipathic α -helical subunits can also form amyloid fibrils adopting a cross- α structure. Some AMPs in the study exhibited a combination of cross- α and cross- β amyloid fibrils, adding complexity to structure prediction. The results showed that the AlphaFold2 ColabFold models favored α -helical structures in the tested amyloids, successfully predicting the presence of α -helical mated sheets and a hydrophobic core resembling the cross- α configuration. This implies that the AI-based algorithms prefer assemblies of the monomeric state which was frequently predicted as helical, or capture an α -helical membrane-active form of toxic peptides, which is triggered upon interaction with lipid membranes.

Keywords

AlphaFold, Amyloids, Antimicrobial peptides, Cross-alpha, Cross-beta

Introduction

Antimicrobial peptides (AMPs) play a critical role in the immune system of organisms across all kingdoms of life and are the first line of defense against pathogens (1). AMPs often self-assemble and some form ordered supramolecular structures that resemble amyloids (2–21). Amyloids vary in sequence, length, composition, and function, and are secreted by organisms from all kingdoms of life. In some instances, they can accumulate in a harmful way or adopt alternative structures, resulting in conditions like neurodegeneration and amyloidosis. However, amyloids can still fulfill their intended functions in their native, oligomeric, or fibrillar states.

Amyloid fibrils are composed of subunits stacked perpendicular to the fibril axis, typically forming cross- β sheets (22). A cross- α form of amyloid also exists in which subunits form α -helical mated sheets (2,3,15). Despite the tightly mated sheets that anchor their fibril structure, amyloids are highly polymorphic, even within the same sequence, with the extreme possibility of a different secondary structure at the fibril form (2,3,15,23,24). This latter observation has been demonstrated by recent structures of amyloidogenic AMPs (14,15,18). The self-assembly properties and polymorphism of amyloids distinguish them from globular and membrane proteins, which constitute the majority of available protein structures (25). This makes amyloid structure determination and prediction difficult (25–29), while structural information remains necessary for understanding their functions and mechanisms, discovering modulators of their activities, and designing novel amyloid sequences for biomedical and technological applications.

With the recent development of new methods for predicting protein structure, such as AlphaFold, there is potential for predicting amyloid structure. In this challenging field, we investigated the current capabilities of these methods. AlphaFold2 (AF2) was introduced at the 14th Critical Assessment of Protein Structure Prediction (CASP) in 2020 and uses innovative neural network architectures and training methods (30). The training of AF2 is based on the evolutionary, physical, and geometric constraints of experimental three-dimensional protein structures deposited in the Protein Data Bank (PDB) to predict the structure of proteins or their complexes from sequence (30). To make the AF2 algorithm more accessible, a collaborative "Jupyter notebook" hosted by Google called ColabFold has been established (31). This platform allows researchers without computational expertise or

resources to use the AF2-ColabFold algorithm to model the structures of protein monomers, homo- and hetero-oligomeric complexes (31).

In this study, our aim was to evaluate the capability of AF2-ColabFold-multimer in accurately predicting the structures of a new class of amyloidogenic antimicrobial peptides (AMPs). These AMPs, namely PSM α 3, uperin 3.5, and aurein 3.3, have recently been characterized at a high resolution and found to form cross- α and cross- β amyloid fibrils (2,3,14,15). Additionally, we examined a specific fragment (residues 17-29) of LL37, a human antimicrobial and immunomodulator, which assembles into a distinct α -helical fibril structure that is non-amyloid in nature (17). Furthermore, we included models of AMPs without experimentally determined structures but with known morphological features, such as citropin 1.3, cyanophlyctin, bombinin H4, and dolabellandin-B2 (18). Lastly, our analysis also encompassed two well-known pathological amyloids, namely amyloid- β (A β) and islet amyloid polypeptide (IAPP). We chose these particular human amyloids because of their short lengths, with 42 and 37 amino acids respectively, and because both are known to display antimicrobial activity (32–36).

Results

Table 1 lists the sequences of the peptides analyzed, and the number of subunits tested. Table S1 lists the key features of the leading AF2-ColabFold predicted models.

Table 1. Tested peptide sequences and AF2-ColabFold multimeric models analysed.

Protein	Sequence	Number of subunits							
		1	10	15	20	25	30	35	40
Citropin-1.3	GLFDIIKKVASVIGGL	V	V		V	V	V		
Cyanophlyctin	FLNALKNFAKTAGRLKSLN	V	V		V		V	V	V
Bombinin H4	LIGPVLGLVGSALGGLLKKI	V	V		V		V		
Dolabellandin-B2	SHQDCYEALHKCMASHSKPFSCSMKF HMCLQQQ	V	V		V	V	V		
Aurein-3.3	GLFDIVKKIAGHIVSSI	V	V		V		V		

Uperin-3.5	GVGDLIRKAVSVIKNIV	V	V		V	V	V		
LL37 (17-29)	FKRIVQRIKDFLR	V	V		V		V		
PSM α 3	MEFVAKLKFVKDLLGKFLGNN	V	V	V	V	V	V		
IAPP	KCNTATCATQRLANFLVHSSNNFGAIL SSTNVGSNTY	V	V	V	V				
Amyloid beta (1-42)	DAEFRHDSGYEVHHQKLVFFAEDVGS NKGAIIGLMVGGVVIA	V	V		V				

Names, sequences, and number of subunits used as input in various AF2-ColabFold experiments. ‘V’ indicate experiment that has been conducted.

AF2-ColabFold models partially recapitulate the mated α -helical sheets of the cross- α amyloid fibril of bacterial PSM α 3

PSM α 3 is a cytotoxic and lytic 22-residue peptide of the PSM family secreted by *Staphylococcus aureus*. The crystal structure of PSM α 3 (PDB id 5I55) revealed, for the first time, a cross- α fibril, a unique amyloid morphology composed on amphipathic α -helices (2,3). The cross- α is an assembly of molecules stacked perpendicular to the fibril axis, further forming tightly mated sheets, as in the canonical cross- β amyloid, but with each molecule forming α -helices rather than β -strands. Recently determined cryo-EM structures of PSM α 3 (PDB id 7T0X, 7SZZ) confirmed the cross- α configuration and further showed a supramolecular assembly of the mated α -helical sheets into nanotubes (37). Notably, the cross- α configuration was also observed for the longer, 44-residue PSM β 2, which forms a helix-turn-helix motif, demonstrating another possibility for mated helical sheet-based assembly (37). To compare the AF2-ColabFold models with the crystal and cryoEM structures, we evaluated the modeled monomer, decamer (10-mer), pentadecamer (15-mer), eicosamer (20-mer), 25-mer and 30-mer PSM α 3.

AF2-ColabFold prediction for monomeric PSM α 3 yielded an α -helical structure with a high confidence measure, namely the predicted local-distance difference test (pLDDT) score, for all models (Figure S1a). For multimer predictions, the closest model with similarity to the experimental cross- α structure was the 1st ranked model of the decameric (10-mer) PSM α 3. This model showed two mated parallel α -helical sheets, as in the experimental structures, yet the α -helices and mated sheets were not as tightly packed (Figure

1a,c). Specifically, in the crystal structure, the distance between α -helical subunits along each sheet is 10.5Å, and the distance between mated sheets, orthogonal to the fibril axis, is 12.6Å (2,3). In the AF2-ColabFold 10-mer model, the corresponding distances are 12.8-16.6Å between α -helices and ~16Å between sheets, resulting in a more loosely packed fibril (Figure 1c), that lacks most of the interhelical polar interactions observed in the experimental structures. In both the experimental and predicted structures, the amphipathic α -helices are arranged so that the interface between sheets is hydrophobic, forming a large hydrophobic core along the fibril (Figure 1b,d). To quantify the tightness of the packing, we compared the solvent accessible surface area (SASA) buried within the fibril. Values for all SASA and interhelical and inter-sheet distances are presented in Table S1. In the PSM α 3 crystal structure, the buried surface area of a single α -helix is 1150Å², representing 50% of its total SASA (2,3), compared to 622Å² (24% of the total SASA) in the 10-mer model. Overall, for a multimer of ten, AF2-ColabFold succeeded in predicting the general configuration of mated α -helical sheets, but with looser packing of the helices compared to the experimentally determined structures, which present an apparently much less stable configuration. Notably, although the pLDDT of all five models is similar and all individual subunits were predicted as α -helices, the four models ranked 2nd to 5th did not resemble the cross- α configuration, and included a barrel and unlikely assemblies with clashes between subunits (Figure S2d,e).

Another AF2-ColabFold model that resembles the cross- α configuration is that of a pentadecamer (15-mer), and in particular the 2nd ranked model, which shows two mated parallel α -helical sheets with a hydrophobic core, but again, not as tightly packed as in the experimental structures (Figure 2a). Specifically, the distances between the α -helices range from 13.7Å to 18.8Å, and between the sheets from 14Å to 19.5Å. Corresponding to the large distance between subunits, the SASA of a central α -helix was 361Å² (14% of the total SASA), which is much lower (less is buried) compared to the experimental structure. Although all five models share the same pLDDT score, the 1st ranked model of the 15-mer also showed two mated α -helical sheets but with several subunits detached from the assembly. The 3rd and 5th ranked models showed an arrangement resembling a globular shape, and the 4th ranked model resembled a barrel (Figure S3).

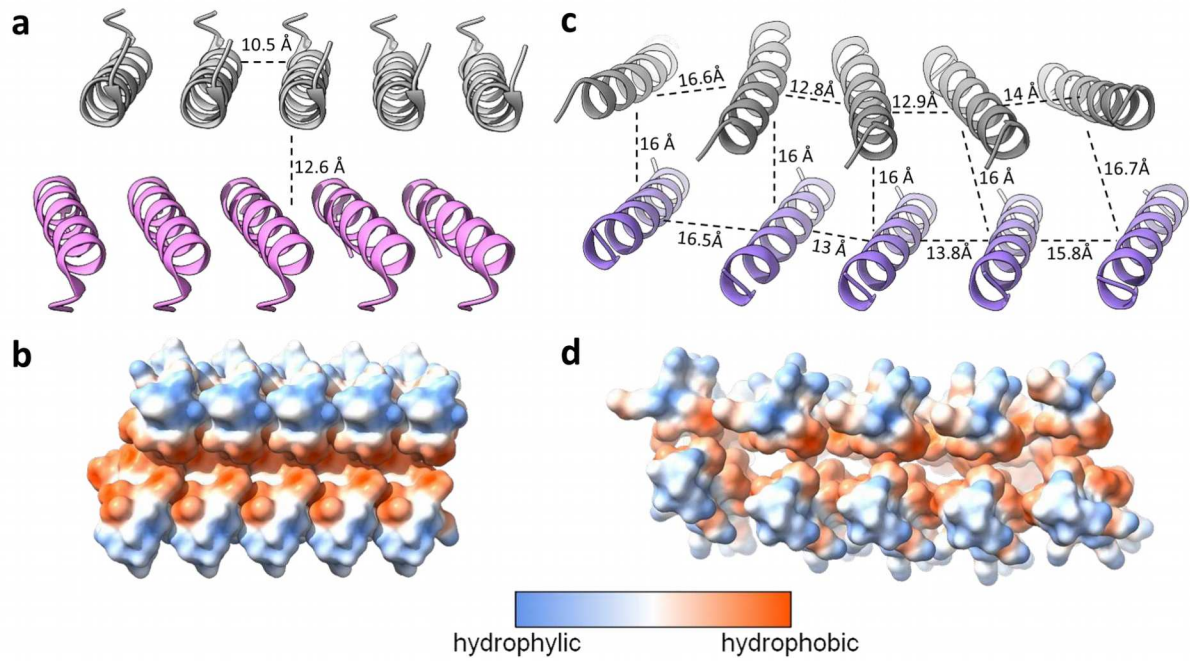


Figure 1. Comparison between experimental and predicted structures of PSM α 3. (a-b) The crystal structure of PSM α 3 (PDB: 5I55). (c-d) The AF2-ColabFold 1st ranked model of a decamer (10-mer). In panels a&c, the view is along the fibril axis, with PSM α 3 colored by sheet, with marked distances between sheets and between α -helical subunits along the sheet. In panels b&d, PSM α 3 is shown in a surface representation colored by hydrophobicity according to the scale bar.

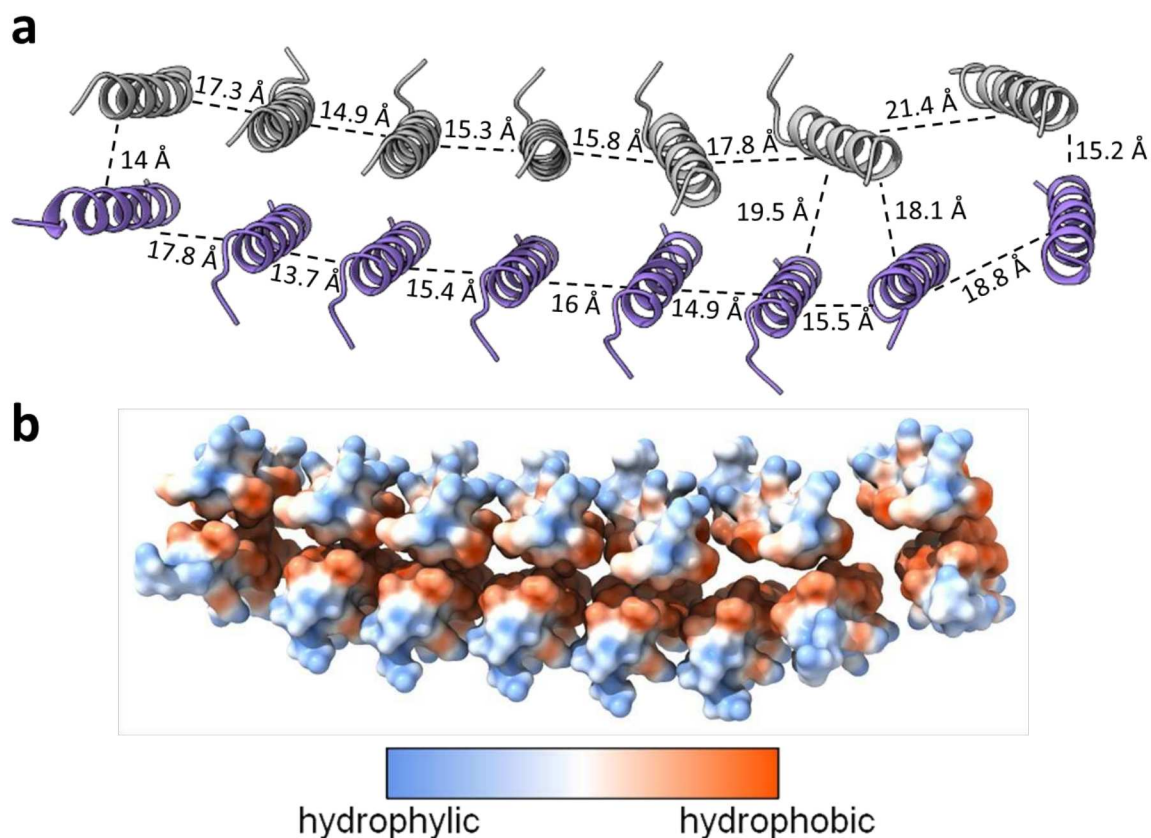


Figure 2. AF2-ColabFold 2nd ranked model of pentadecameric (15-mer) PSMα3. (a) PSMα3 model is viewed along the fibril axis and colored by sheets, showing distances between subunits and sheets. (b) Surface representation colored by hydrophobicity as indicated by the scale bar.

The 2nd ranked model of the eicosamer (20-mer) PSMα3 showed a 12-subunit core of mated sheets with six peptides from each sheet, with distances of 14Å - 19.6Å between α -helices along the sheet, and 15.4Å - 19.5Å between the two sheets. The buried SASA of one helix was 466Å² (18%), indicating a packing that is looser than the 10-mer but tighter than the 15-mer. The rest of the subunits were scattered around this core. This suggests that adding more subunits surprisingly predicts disassembly of the fibril-like arrangement compared to the 10- and 15-mers. The other ranked models showed diverse assemblies of α -helices, including helical sheets, spirals, and barrel-like helical clusters that are not very tightly packed (Figure S4). Adding more subunits, of 25- and 30-mers, leads to further deviation from the cross- α configuration, showing different shapes that attempt more globular-like clusters, but with loose packing of the helices compared to the cross- α tight fibril packing (Figures S5&S6).

Overall, the predictions of PSM α 3 multimers show a pLDDT score around 40 on a scale of 0 to 100, indicating low confidence, and structural diversity. Some of the models predicted mated α -helical sheets similar to the cross- α fibril arrangement, but with a looser packing compared to the experimental structure and often with surrounding detached subunits. The supramolecular nanotube structure shown by the cryoEM structure of PSM α 3 (37) was not recapitulated, even by modeling of a larger number of subunits. It is noteworthy that in some of the predicted models, the PSM α 3 α -helices spiral, echoing the structures shown by DeGrado and co-workers using designed peptides (38).

AF2-ColabFold models of uperin 3.5 recapitulate some cross- α features, and fail to predict the cross- β configuration

Uperin 3.5 is an antimicrobial peptide secreted on the skin of *Uperoleia mjobergii* (Australian toadlet). This peptide displays a secondary structure switch and can form both cross- β and cross- α amyloid fibrils, both of which have demonstrated by high-resolution structures and biophysical measurements (14,15). The cross- α crystal structure (PDB Id: 6GS3) (15) resembled the PSM α 3 fibril with mated sheets of stacked α -helices, but while the latter showed a parallel orientation of the α -helices along the sheets, uperin 3.5 showed an interesting feature of an antiparallel orientation of the α -helices (2,15) (Figure 3a,b). This revealed a surprising feature of parallel and antiparallel sheets, formed not only by β -strands, but also by α -helices. The cross- β form of uperin 3.5 was determined by cryo-EM (PDB Id: 7QV5), showing a three-blade symmetric propeller of nine peptides per fibril layer including tight β -sheet interfaces (14) (Figure 3e). This revealed a remarkable ability of a natural sequence to switch between different secondary structures depending on the environmental conditions. In particular, lipids mimicking a membrane environment induced the transition into the cross- α form (15,39–42). To compare the AF2-ColabFold models with the crystal and cryoEM structures, we evaluated the modeled monomer, decamer (10-mer), eicosamer (20-mer), 25-mer and 30-mer uperin 3.5.

The AF2-ColabFold prediction for monomeric uperin 3.5 is an α -helical structure in all five ranked models, with an average pLDDT score of $\sim$ 90 (Figure S7). All models of the decamer (10-mer), with an average pLDDT score of $\sim$ 40, consisted of clusters of helices reassembling a globular fold (Figure S8). One of the subunits in the 4th ranked model of the 10-mer showed a random coil with no determined secondary structure (Figure S8). The five models of the 20-mer uperin 3.5 also showed diverse arrangements despite having similar

pLDDT scores. The 2nd and 3rd ranked 20-mer models showed some remote resemblance to spiraling α -helical mated sheets (Figure S9).

Starting with 25-meric subunits, a higher similarity to cross- α was observed. The 3rd ranked 25-meric uperin 3.5 shows two mated sheets of α -helices, one with eight subunits and the other with seven (Figure 3), while the rest of the α -helices are scattered around (not shown). The sheets contain α -helices parallel to each other, as in PSM α 3 (2), and not antiparallel as in the crystal structure of uperin 3.5 (15). The antiparallel orientation of the uperin 3.5 α -helices allowed for a tight packing, with an interhelical distance of 9.8Å and inter-sheet distance of 8.6Å (Figure 3a). In the model, the distances are much larger, with interhelical distances of ~15Å in one sheet and 14.5-19Å in the other, and inter-sheet distances of 13-18.6Å (Figure 3c). Accordingly, while the buried SASA of one α -helix in the crystal structure is 752Å² (40% of the total SASA), in the model it is only 194Å² (10%) for a representative helix, highlighting the much looser packing of the model. With all 25-mer models having the same averaged pLDDT score of ~40, they showed diverse arrangements including clusters of helices in globular or spiral shapes (Figure S10).

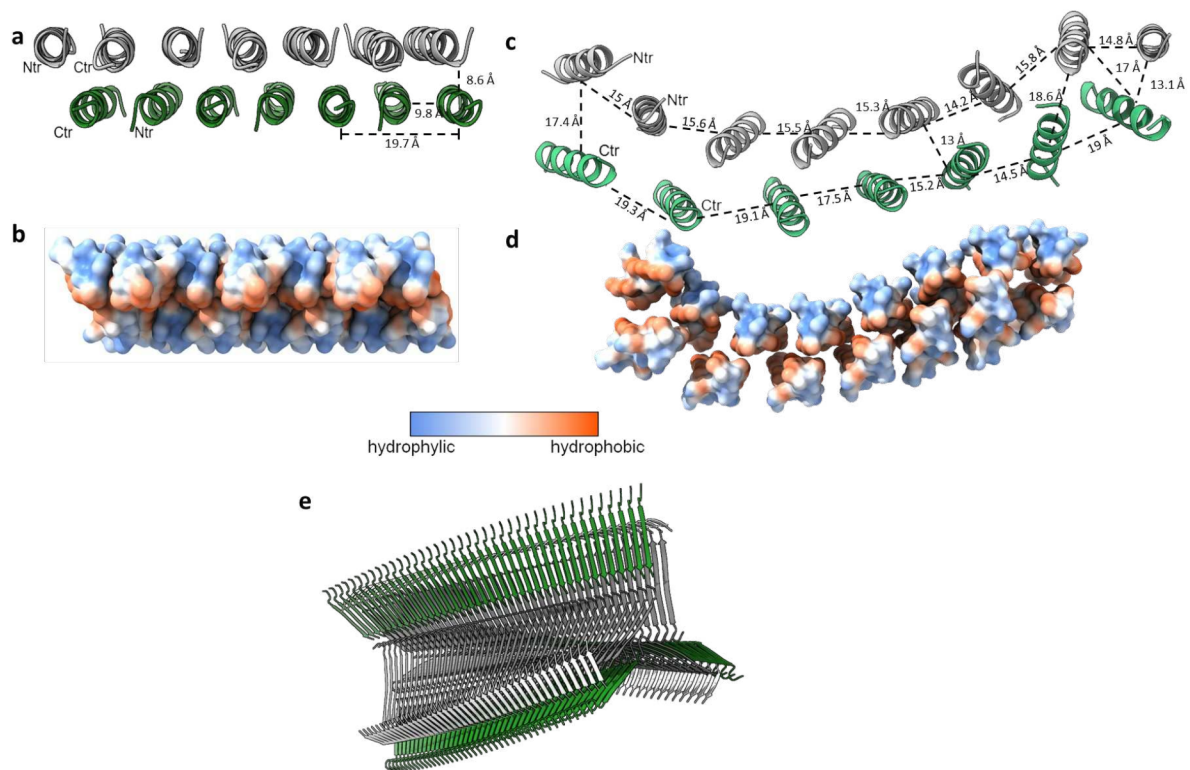


Figure 3. Comparison between experimental and predicted structures of uperin 3.5. (a-b) The crystal structure of uperin 3.5 (PDB: 6GS3). (c-d) The AF2-ColabFold 3rd 25-meric ranked model. (e) The cryo-EM cross- β structure (PDB: 7QV5). In panels a-d, the view is along the fibril axis. For clarity, only the core arrangement of 15 helices is shown, without

the scattered subunits present in the model. In panels a&c, uperin 3.5 is colored by sheets, and the distances between sheets and between two antiparallel and parallel neighboring subunits along the sheet are marked. In panels b&d, uperin 3.5 is shown in a surface representation colored by hydrophobicity, as indicated by the scale bar. In panel e, the view is along the fibril axis with a slight tilt, with uperin 3.5 colored by β -sheets.

An interesting observation is that the top-ranked model for the 30-meric uperin 3.5 showed α -helical mated sheets that were curved and spiraling in a parallel orientation (Figure 4). In this model, each sheet has an equal number of subunits and is more tightly packed than the 25-mer model, but not as tightly as the crystal structure. The model can be divided into two regions, with one section consisting of straight sheets and the other of spiraling sheets. The straighter part showed interhelical distances of $\sim 14.6\text{\AA}$ in one sheet and $\sim 12.5\text{\AA}$ in the other, and an inter-sheet distance of $\sim 12\text{\AA}$ (Figure 4a). The buried SASA of a representative helix is $454\text{\AA}^2$ (24% of the total SASA). The spiraling part had interhelical distances of $\sim 14.4\text{\AA}$ in one sheet and $\sim 14.6\text{\AA}$ in the other, and an inter-sheet distance of $\sim 12.3\text{\AA}$ (Figure 4b). The buried SASA of one representative helix is $363\text{\AA}^2$ (19% of the total SASA). Importantly, the sheets are oriented to maintain a hydrophobic core between the sheets. All 30-mer models have the same averaged pLDDT score of ~ 40 , and they showed diverse arrangements including clusters of helices in globular or spiraling shapes (Figure S11), similar to the 25-mer models. In the 4th and 5th ranked 30-mer models some of the subunits form random coils with no defined secondary structure. Notably, none of the AF2-ColabFold models we evaluated predicted an antiparallel orientation of the helical sheets (Figure 3a). Furthermore, the predictions did not anticipate the cross- β configuration which is formed by a three-blade symmetric propeller of nine peptides per fibril layer including tight β -sheet interfaces (Figure 3e).

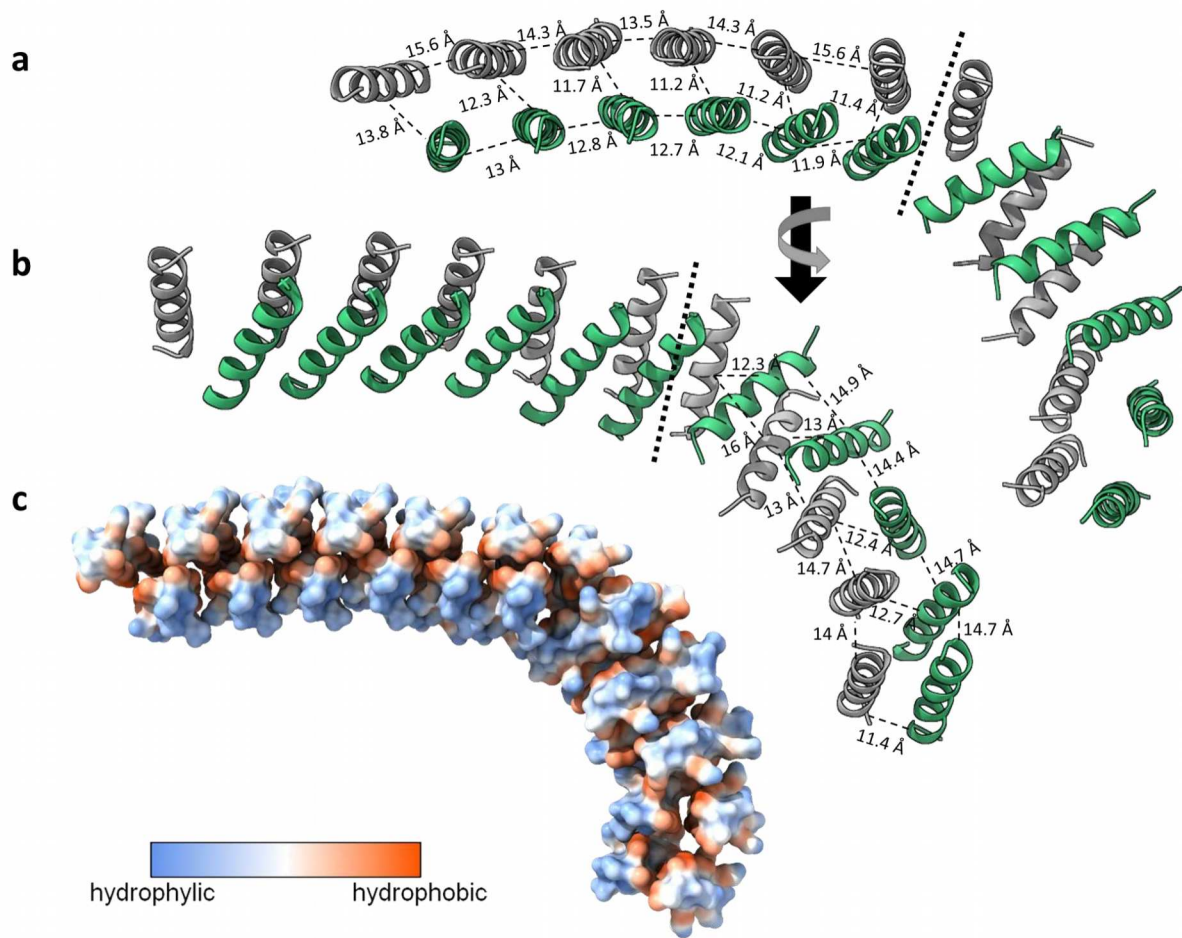


Figure 4. AF2-ColabFold prediction of the 1st ranked model of the 30-mer uperin 3.5. Only the core arrangement of 22 helices is shown, excluding eight scattered subunits that exist in the model but are not shown for clarity. (a&b) Two orientations of the model are shown, colored by sheets, indicating distances between subunits and sheets. The bold dotted line indicates the transition from aligned to spiraling subunits. (c) Surface representation colored by hydrophobicity, as indicated by the scale bar.

AF2-ColabFold failed to predict the kinked cross- β fibril of aurein 3.3 and instead showed arrangements of spiraling and curved α -helical mated sheets

Aurein 3.3 is an amphibian antimicrobial peptide isolated from the *Litoria/Ranoidea raniformis* (southern bell frog). Like uperin 3.5, aurein 3.3 showed a secondary structure switch in the fibrillar form to include both α -helical and β -rich conformations (18). The cryo-EM structure of aurein 3.3 revealed a cross- β fibril consisting of six peptides per fibril layer, all with kinked β -sheets allowing for a rounded compactness of the fibril (14) (Figure 5e). This unique assembly has been correlated with reversible and functional amyloid formation

(43–48). We modeled the monomer, decamer (10-mer), eicosamer (20-mer), 25-mer and 30-mer and 40-mer aurein 3.3.

The predicted monomeric aurein 3.3 was α -helical, with a high pLDDT score of ~ 90 for the first 13 of 16 residues (Figure S12). The multimer predictions were all α -helical subunits and failed to predict the β -rich configuration. The decamer (10-mer) models, which had a similar pLDDT score, showed a clustering of helices into a globular-like shape with clashes between subunits (Figure S13). From 20 subunits and up, the models were better packed into intriguing fibrils composed of α -helices. The 1st ranked model of the eicosamer (20-mer) showed 16 subunits arranged in two α -helical sheets, resembling cross- α , but with a twist between helices and curved sheets, so that the helices along each sheet are not strictly parallel or antiparallel to each other, resulting in clashes between atoms (Figure S14d,e). The 2nd ranked model showed a core of eight helices, similar to the spiraling cross- α amyloid-like structures (38) (Figure 5a). Within this region, the averaged distances between α -helical subunits along the two mated sheets were 12.7Å and 12.6Å, and the averaged distance between sheets was 14.7Å. The buried SASA of a representative helix was 811Å² (43.5% of the total SASA), indicating tight packing, similar to the cross- α structure of PSM α 3. This arrangement also recapitulated the hydrophobic core between the sheets (Figure 5b). The 3rd ranked model, similar to the 5th ranked model, showed pairs of mated spiraling α -helical sheets curved into a doughnut shape, retaining the core between the paired sheets (Figure 5c,d). The 20-mer models differed from each other and showed a similar pLDDT score of ~ 40 (Figure S14).

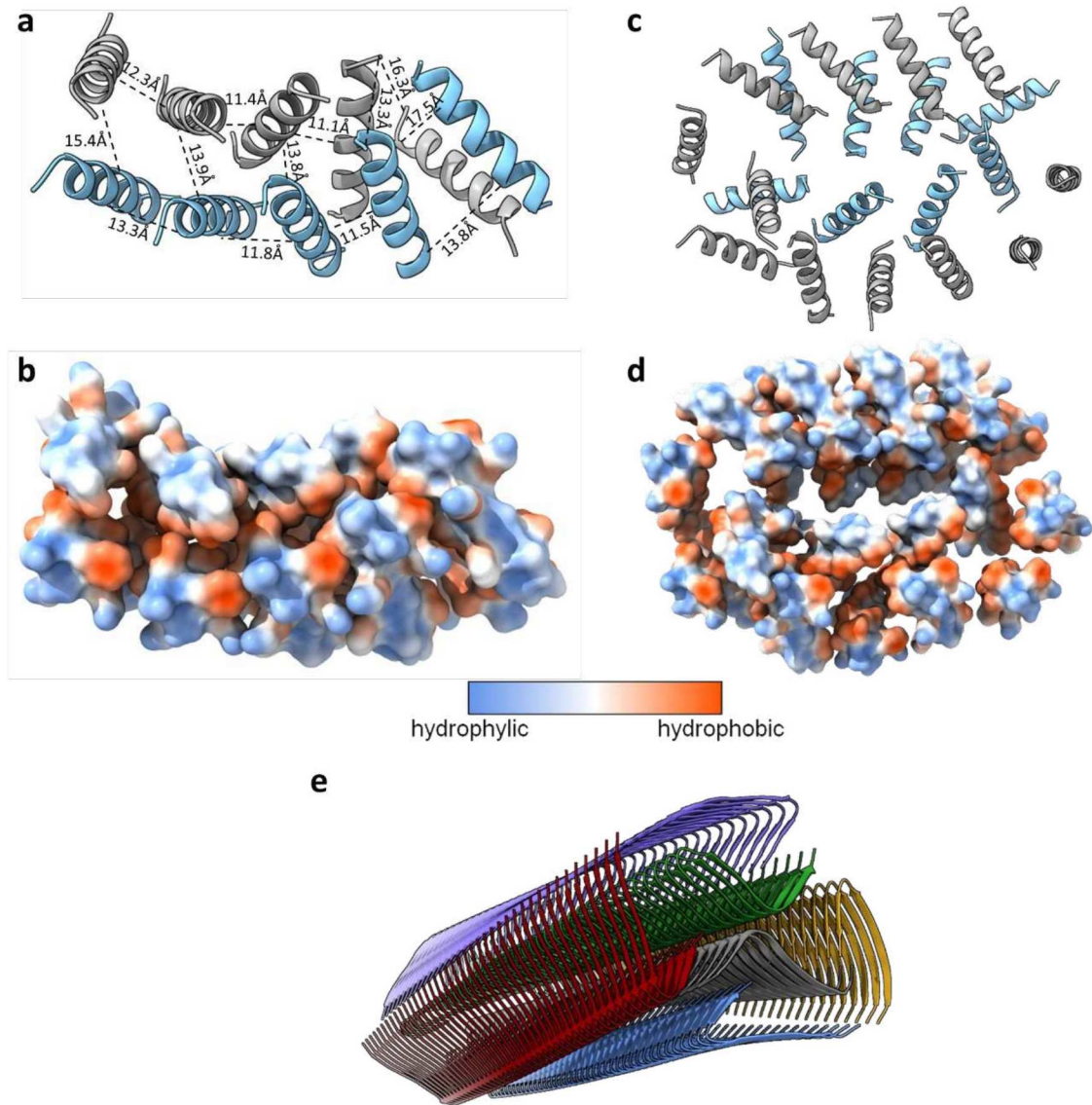


Figure 5. Comparison between experimental and predicted structures of aurein 3.3. (a-d) AF2-ColabFold models of 20-mer aurein 3.3, including the spiraling cross- α core of the 2nd ranked model (a-b) and the 3rd ranked model (c-d). The structures are either colored by sheets, showing the distances between subunits and sheets (a&c) or shown in a surface representation colored by a hydrophobicity scale as indicated (c&d). (e) The cryo-EM cross- β structure of aurein 3.3 (PDB: 7QV6) is shown along the fibril axis the β -sheets in different colors.

Models with a higher number of aurein 3.3 subunits (25-, 30- and 40-mers) supported the spiraling and curving cross- α into a spiraling fibril rather than the typical elongated fibril (the 25-mer models are shown in Figure 6). The 2nd, 4th and 5th ranked models of the 25-mer aurein 3.3 showed an average inter-sheet distance of 13.6Å, 21Å, and 15Å, respectively. The distance between subunits along the two sheets was 17.4Å and 16.5Å for the 2nd ranked

model, 16.4Å and 15.9Å for the 4th ranked model, and 18Å and 18.6Å for the 5th ranked model (Figure 6a,c and Figure S15f). The buried SASA for a single helix in the 2nd ranked model varied along the structure, with two examples of 722Å² and 150Å² (38.6% and 7.8% of the total SASA, respectively). In the 4th and 5th ranked models, the buried SASA for a single helix was 183Å² (9.8%) and 171Å² (9.1%), respectively. Overall, these spiraling curved fibrils are not as tightly packed as the cross- α crystal structures of PSM α 3 or uperin 3.5, but share the hydrophobic core between the helical sheets (Figure 6b,d and Figure S15g). The 3rd ranked model of the 25-mer showed a different, rounded shape (Figure S15). Surprisingly, in the 1st ranked 25-mer model, some of the subunits were random coils, while its average pLDDT score was higher than the other models (Figure S15). The models of the 30-mer aurein 3.3 recapitulated some of the features of the 10- and 25-mer models (Figure S16). Overall, the AF2-ColabFold predictions for aurein 3.3 were mostly similar to PSM α 3 and the spiraling cross- α of designed peptides (2,3,38), along with more extreme curving and spiraling of the fibril itself. There was no evidence of kinked β -sheets as determined by cryo-EM (Figure 5e).

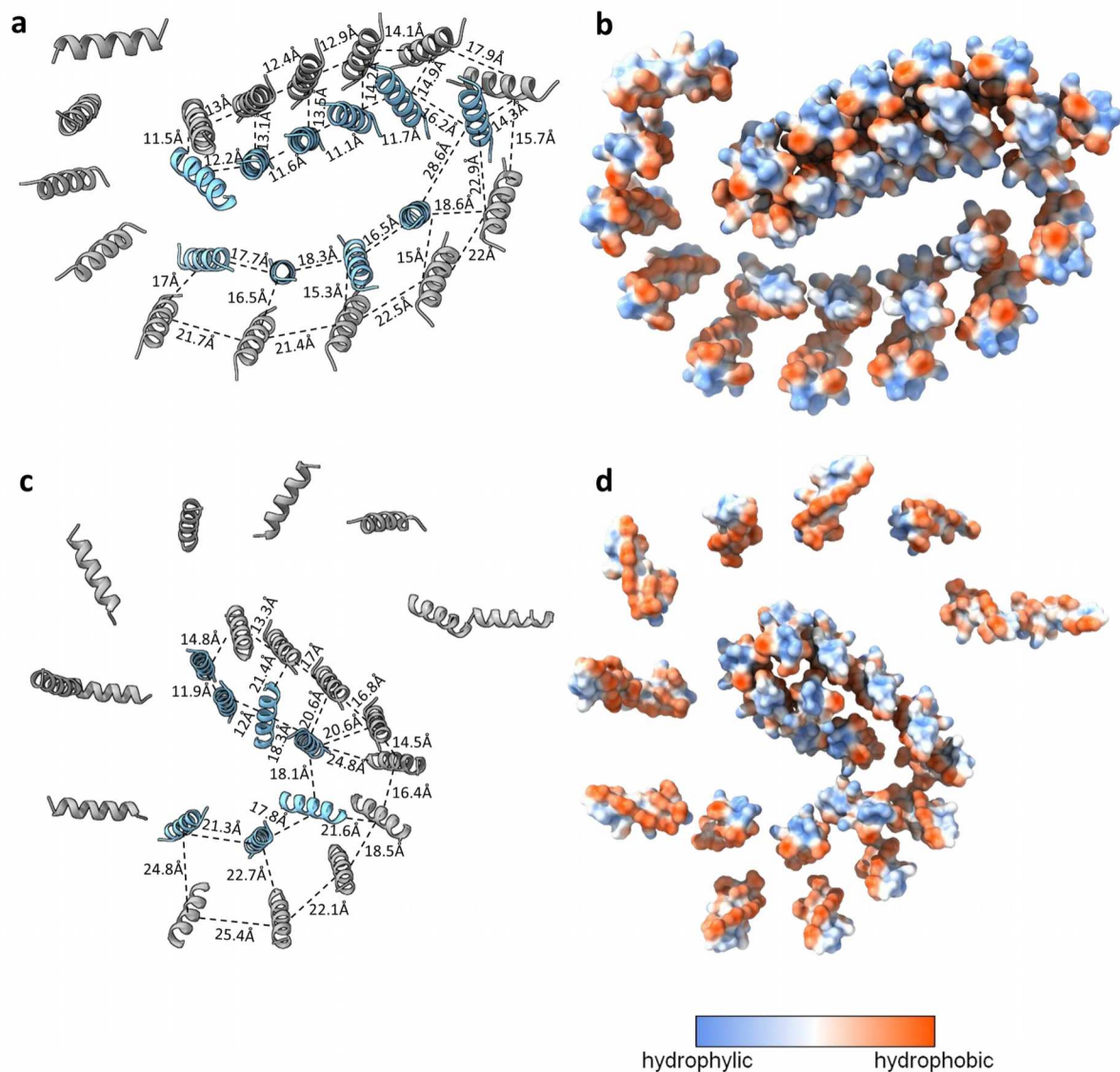


Figure 6. AF2-ColabFold predictions of the 25-mer aurein 3.3.

The 2nd (a&b) and 4th (c&d) ranked models are colored by sheet, showing inter-sheet and interhelical distances (a&c), or displayed in a surface representation colored by hydrophobicity, as indicated by the scale bar (b&d).

AF2-ColabFold failed to predict the non-amyloid α -helical fibril structure of human LL37(17-29)

LL37 is a host defense peptide cleaved from hCAP18, the only human cathelicidin (49,50).

LL37 forms an amphipathic α -helix that self-assembles into fibrils, which are critical for DNA binding and activity (17,49–51). The LL37 fragment containing residues 17-29 is an active antibacterial core and the shortest fragment that retains antiviral activity (12) (52). The

crystal structure of LL37(17-29) (PDB id 6S6M) (Figure S17a,b) revealed a unique self-assembly of amphipathic helices into a densely packed, elongated hexameric fibril with a central pore. The fibril is composed of four-helix bundles with a hydrophobic core that are further assembled through a network of polar interactions (17). To evaluate the predictive ability of AF2-ColabFold for this unique fibril structure, we modeled the monomer, decamer (10-mer), eicosamer (20-mer), and 30-mer LL37(17-29).

The monomeric model of LL37(17-29) is α -helical with flexible termini (Figure S17). The models for the decamer (10-mer), eicosamer (20-mer), and 30-mer LL37(17-29) showed different assemblies of helices with no specific order, are mostly loose, do not resemble the crystal structure or any known ordered fibril, and had a wide range of pLDDT scores (Figures S18-S20). Several individual subunits also lost the α -helical structure and exhibited random coils (as shown in Figure S19f). Overall, AF2-ColabFold was not successful in predicting the unique helical fibril of LL37(17-29) compared to the predictions of PSM α 3 and uperin 3.5, which recapitulated some of the cross- α features.

Amyloid-forming AMPs without experimental structures

We have previously identified and characterized AMPs that self-assemble into cross- α and cross- β amyloids, some with the ability to switch conformations over time or induced by lipids (18). These include the amphibian citropin 1.3, cyanophlyctin, and bombinin H4 (18,53–55), as well as dolabellandin-B2 secreted by sea hares (18,56). We selected these four peptides because they have different structural properties (18). While cyanophlyctin is unstructured in solution, also in the presence of lipids, it adopts a cross- α conformation in the fibril form, and its helicity is further induced by the presence of membrane lipids. Citropin-1.3 and bombinin H4 are unstructured in solution but adopt a helical conformation in the presence of membrane lipids. In the fibril state, citropin-1.3 forms a cross- β structure, but the presence of lipids induces the helical state. Bombinin H4 forms a cross- α fibril, and its helicity is also promoted by lipids. In contrast to most AMPs tested, which are unstructured or helical in solution, dolabellandin-B2 adopts a β -rich conformation in solution and in the fibril form, with a minor helical population in the fibril form, and is insensitive to the presence of lipids. Considering these biophysical properties (18), we evaluated the AF2-ColabFold models of the monomeric and multimeric forms of these AMPs (Figures 7, S21-S41).

All monomeric models of the four peptides were α -helical with relatively high pLDDT scores, with dolabellandin-B2 showing a helix-turn-helix prediction (Figures S21,S26,S32,S37). There was no evidence for a β -rich structure for any of these AMPs in the monomeric or any of the multimeric models described below (Figures S21-S41), even for dolabellandin-B2 which is predominantly β -rich, even in solution and with lipids (18). Among all AF2-ColabFold models of these peptides, three recapitulate the continuous α -helical mated sheets, one for citropin 1.3 and two models of bombinin H4. None of the models of cyanophlyctin, tested up to 40-mers (Figures S26-S31), or dolabellandin-B2, tested up to 30-mers (Figures S37-S41), showed a fibril-like form or mated sheets.

The 1st ranked eicosameric (20-mer) model of citropin 1.3 resembled the parallel cross- α structure of PSM α 3 and was even as tightly packed (Figure 7a). Specifically, the averaged distances between α -helical subunits along each of the two sheets were 10.7Å and 10.1Å, respectively, and the averaged distance between the sheets was 11.4Å. The buried SASA of one helix was 944.3Å² (55.3% of the total SASA), thereby showing the most densely packed model among all peptides evaluated here. The addition of citropin 1.3 subunits (in the 25-mer and 30-mer) resulted in the loss of the fibril-like assembly (Figure S24-S25), similar to the case of PSM α 3. Nevertheless, we observed a smaller assembly of three-helical bundles forming a triangular shape with a hydrophobic core, within the 2nd ranked model of the 20-mer and 5th ranked model of the 25-mer citropin 1.3 (Figure S23-S24). We wonder whether this assembly might represent a stable oligomeric structure.

The 2nd ranked eicosameric (20-mer) model of bombinin H4 contained clashes between atoms although the packing is rather loose (Figure 7c). Specifically, the averaged interhelical distances along the two sheets were 13.4Å and 13.9Å, and the averaged inter-sheet distance was 20.7Å. The buried SASA of one helix is 146.1Å² (21.4% of the total SASA). The 3rd ranked 25-mer bombinin H4 model also showed a loose packing, with averaged interhelical distances along the two sheets of 14.6Å and 14.9Å, and an averaged inter-sheet distance of 19.4Å (Figure 7e). The buried SASA of one helix is 100.2Å² (4.7% of the total SASA). In all three models shown, the hydrophobic pattern indicates a hydrophobic core and a hydrophilic surface similar to the experimentally determined cross- α structures (Figure 7b,d,f).

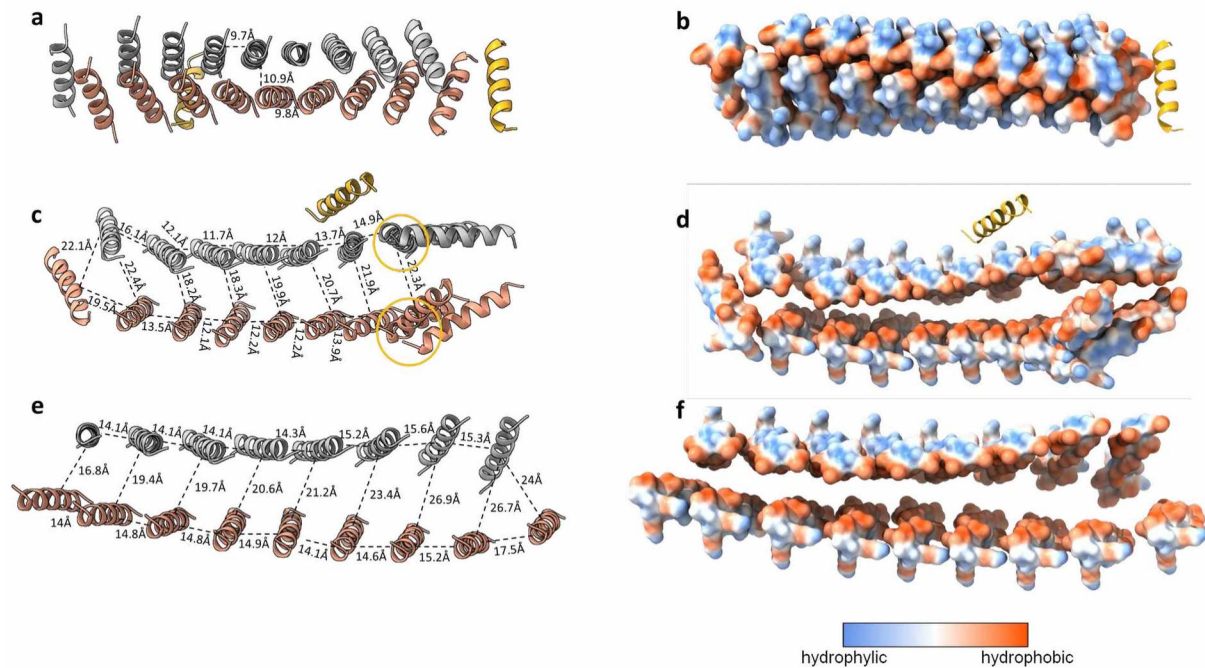


Figure 7. AF2-ColabFold models of amyloid-forming AMPs without experimental structures

(a&b) The 1st ranked model of the 20-mer citropin 1.3. (c&d) The 2nd ranked model of the eicosamer (20-mer) bombinin H4. Yellow circles mark clashes between subunits in panel (c). (e&f) The 3rd ranked model of the 25-mer bombinin H4, with eight sparse subunits removed for clarity. Panels (a&c&e) are colored by sheets, showing inter-sheet and interhelical distances. In gold are subunits that do not participate in the continuous sheets. Panels (b&d&f) are in a surface representation colored by hydrophobicity, as indicated by the scale bar.

Models of human pathological amyloids with antimicrobial activity, A β and IAPP

Human amyloids are notorious for their involvement in neurodegenerative diseases such as Alzheimer's and Parkinson's, in which protein plaques accumulate in the brain. Alzheimer's-associated A β is a 42 (or 40) residue long peptide derived from amyloid precursor protein (57). IAPP is a 37-residue peptide associated with type 2 diabetes, present in various organs, most commonly in the pancreas, and is usually associated with insulin, and involved in glucose metabolism and homeostasis (58). Both IAPP and A β are unstructured proteins that can form cross- β amyloids (59,60). In the presence of anionic bilayer micelles they adopt a

soluble α -helical structure, which is thought to accelerate fibrillation and cross- β formation (59,61–63). IAPP and A β demonstrate antimicrobial activity (32,33,36,64), thereby exhibiting functional similarities with the AMPs. Their characteristics, including their short length, the display of both α -helical and β -rich states, and their antimicrobial activity, make IAPP and A β optimal candidates for assessing the predictions of AF2-ColabFold. We conducted an evaluation of the AF2-ColabFold models of these human amyloids in comparison to the models of the AMPs.

The AF2-ColabFold monomeric model of A β 1-42 is unstructured, as previously reported (60). The pLDDT score of the monomeric A β is lower than that of any of the other monomeric models tested here (Figure S42), perhaps due to its greater length. In the model of decameric (10-mer) A β , most of the subunits are also unstructured, but in the 1st, 2nd and 4th ranked models, some regions are α -helical, and in the 3rd and 5th ranked models some regions form short β -sheets, but most of them consist of clashing β -strands (Figure S43). The models of eicosamer (20-mer) A β showed a similar trend, with some models partially structured as helices (Figure S44).

IAPP is known to be unstructured in solution (59). The AF2-ColabFold monomeric models, with similar pLDDT scores, are α -helical (Figure S45), resembling the helical structure of IAPP in a membrane mimicking environment (61,62). In all decameric (10-mer), pentadecameric (15-mer), and eicosameric (20-mer) models, the subunits are α -helical, with no β -strands detected, while the C-terminal part has a lower pLDDT score, showing unstructured regions. In contrast to the unstructured assembly of the A β models, IAPP predictions show α -helices clustering in various forms, but not as mated sheets (Figure S46-S48).

Cross- α architecture is rare among proteins with experimental structures

The search for structural similarity of different helical assemblies derived from the PSM α 3 cross- α fibril structure, namely arrangements of 4-6 helices in the same or mated sheets, yielded several results, mostly from synthetic sequences or truncations of natural sequences (65–67). Most of the resulting similar structures showed multiple α -helical assemblies of helix-loop-helix bundles, but none of them were continuous sheets or repetitive units forming fibrils (68–71). This supports the rarity of α -helices stacked perpendicular to the fibril axis in the databases.

Secondary structure switchers are rare among proteins with experimental structures

The secondary structure switch in the fibril form, demonstrated at high resolution for uperin 3.5 (14)(15) and suggested for additional amyloid-forming AMPs based on secondary structure measurements (18), may be one of the reasons hindering correct modeling of amyloid fibril structures. We sought to analyze the prevalence of secondary structure switchers among proteins with known experimental structures. Specifically, we searched the PDB for pairs of highly similar protein chains, as described in the Methods section, that contain helical structure in one chain and extended secondary structure in the other chain for at least half of their length.

The analysis yielded only five proteins that have PDB structures with two different secondary structures (Table S2): human A β , prion protein, and glucagon, amphibian uperin 3.5, and bacterial cold shock protein CspA. Except for the latter, the resulting structures are all from proteins that form amyloids, although this feature was not specified in the search parameter. The structure of CspA was determined as a folded protein containing β -sheets and as a cotranslational helical folding intermediate (Figure 8), which explains the differences in structure. For human A β , prion protein, and glucagon, the differences arise from comparing the α -helical form of the protein in its soluble monomeric or oligomeric state, with the cross- β fibril form (examples are provided in Figure 8 and full list of PDBs is given in Table S2). Uperin 3.5 is the only example in the PDB, as far as the analysis could identify, where the same sequence shows both helical and extended conformations in the fibril form. This demonstrates the rarity of secondary structure switchers, at least among proteins with known structures, which may have influenced AlphaFold2 training.

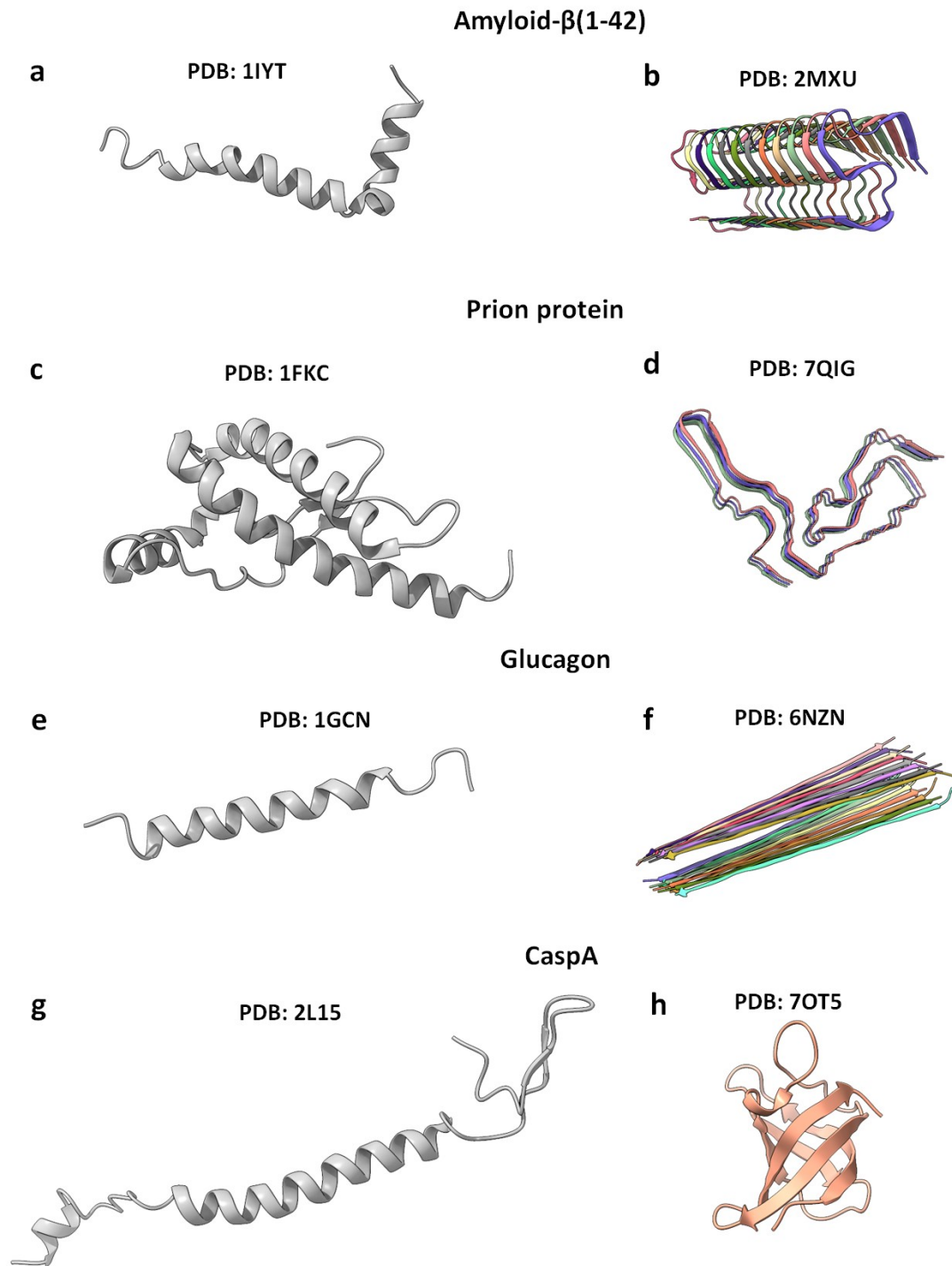


Figure 8. Proteins of known PDB structure observed in two different secondary structures. The structures show α -helical configuration in solution and cross- β in the fibril form of A β 1-42 (PDB 1IYT (72), 2MXU (73)) (a&b), human prion protein (PDB 1FKC (74), 7QIG (75)) (c&d), and glucagon (PDB 1GCN (76), 6NZN (77)) (e&f). The structures of CaspA are shown as a cotranslational helical folding intermediate (PDB 2L15 (78)) (g) or as a folded protein containing β -sheets (PDB 7OT5 (79)) (h) (g&h). (b&d&f) The fibril forms are colored by chain.

Model accuracy and statistics

While discussing the results it is important to take into consideration the estimated prediction accuracy. AF2 provides a per-residue accuracy measure called pIDDT, that represents confidence level on a 0-100 scale, 100 being perfect. AlphaFold assigned pIDDT values in the range of 90 for the monomers, indicative of very high confidence. Notwithstanding, however, values dropped considerably (range of about 20-40) for multimers, indicating much lower confidence. AF2 also provides an estimate of the predicted deviation between each residue of the model from the correct structure after superimposition of one pair of corresponding residues. This estimate, called predicted alignment error (PAE), is reported in Å RMSD. Overall, very low PAE values (near 0) were assigned for predicted superimposition within each of the monomers (intra-monomer) both alone and within the context of the multimers, indicating that the predicted structure of the monomers would superimpose nearly perfectly on the real structure. However, PAE values between monomers (inter-monomer) were rather high, indicating very low confidence in the relative orientations of one monomer with respect to others. Overall, AF2 appears highly confident in the individual monomeric structure but is much less certain about how they pack against each other. AF2 provides another metric known as sequence coverage, which represents the number of sequences per position during the multiple sequence alignment phase. Notably, for IAPP, there are around 900 sequences in the alignment, but more than half of these have less than 50% identity. For A β , aurein 3.3, and PSM α 3, the coverage is approximately 80, 5, and 4 homologous sequences respectively, with three-quarters of these sequences having more than 80% identity. All other proteins tested have a sequence coverage of only 2-3, with 100% identity.

Discussion

Our research focused on functional amyloids with antimicrobial activity, including peptides secreted by a variety of organisms, ranging from prokaryotes to eukaryotes, as well as two human amyloids known to be involved in diseases that also display antimicrobial activity. Although many amyloids play important physiological roles in various organisms and are expected to be subjected to evolutionary selection pressure, they are fundamentally different from globular and membrane proteins, which make up most of the available protein

structures (25). Amyloid-forming proteins and peptides are often unstructured in solution and highly polymorphic when assembled, with their morphology frequently affected by environmental factors such as pH, salts, lipids, metal ions, nucleic acids, post-translational modifications, other proteins, and other factors (80). An extreme case of polymorphism is different secondary structure in highly similar sequences. Our analysis of the PDB has shown that among all protein structures, only one bacterial cold shock protein and four eukaryotic amyloids exhibit complete secondary structure polymorphism, subject to our definition. Such massive switch was observed either when comparing soluble to fibrillar structures in the case of human A β , prion and glucagon, or two fibrillar forms in the case of uperin 3.5 (Table S2 and Figure 8). This presents an extremely small group, a water-drop in the sea of all known structures used for the training of AF and similar methods. Furthermore, the training of the AF2 model is primarily based on low-order oligomers and may be less applicable to higher-order oligomers and fibrils, in addition to the general lack of confidence in oligomers relative to monomers (81). In this study, we examined how these biases in training sets and the inherent polymorphism and self-assembly properties affect amyloid structure prediction using advanced artificial intelligence (AI)-based methods, particularly AlphaFold2 via the ColabFold algorithm (31). While discussing the results it is important to take into consideration the estimated prediction accuracy described in the results section.

A significant observation is that most of the predictions reported here were based on α -helical structures, and did not encompass any β -rich assemblies, even though our test cases included uperin 3.5, aurein 3.3, A β and IAPP, for which cross- β structures were observed experimentally (14,58,82), and such assemblies are a hallmark of amyloid fibrils. Specifically, the AF training set comprised 155,204 PDB entries released prior to August 28, 2019 (30). When we compared these structures with the AmyPro database, a repository of amyloid proteins documented in scientific literature (83), we identified 77 β -rich amyloid structures (including redundant sequences) (0.05% of the training set) and one cross- α structure of PSM α 3 (PDB id 5I55) (0.00064% of the training set). It is difficult to ascertain the extent to which a single or a few dozen structures could influence the AF2 algorithm and its accuracy. However, the growing presence of amyloid polymorphic structures in the PDB offers hope that future algorithms will be able to identify these unique features. Another consideration regarding the accuracy of prediction is the short length of the sequences tested here. Among the 270304 chains of structures in the PDB that were released prior to 28 August 2019, 32,125 (11.9%) of those are shorter than 42 amino acids, which is our longest

tested peptide (A β). Although this proportion is not particularly high, it is certainly not negligible. Another important factor to consider in the analysis is the ratio of helical structures to β -rich structures in the training set. A statistical analysis of the PDB, using the structural classification of proteins (SCOP), revealed that among the structures released before August 28, 2019, there were 18,881 proteins primarily composed of α -helices and 29,123 proteins mainly composed of β -strands (Table S3). Therefore, it is unlikely that the training set would have a disproportionate representation of helices, both in general protein structures and specifically among amyloids. Alternatively, there could be other factors contributing to the significant inclination towards α -helical models in the predicted structures. One plausible explanation for the predicted helical configuration of the multimer could be its influence by the predictions of the monomeric or soluble states, which for some of the cross- β amyloids, are either unstructured or helical. Another potential explanation could be the functional significance inherent in the protein structures, given that the membrane-active form is frequently helical, as is the oligomeric form of several amyloids.

Both PSM α 3 and uperin 3.5 were predicted by AF2 to exhibit cross- α -like mated helical sheets (Figures 1-4), aligning with their observed formation of cross- α fibrils in experimental structures. However, there is a nuance: PSM α 3 predominantly adopts a helical conformation (2), whereas uperin 3.5 tends to form cross- β fibrils unless external factors, such as lipids, induce its helicity (15). Additionally, uperin 3.5 and PSM α 3 exhibit contrasting helical orientations along their sheets. Uperin 3.5 has an antiparallel orientation, while PSM α 3 is parallel. This distinction is further highlighted by their differing thermostabilities. Uperin 3.5 demonstrates greater stability, potentially due to its more stable alternate cross- β conformation (15). Such differences likely originate from their respective sequences, as indicated by the divergent outcomes of our AF2 prediction. It is noteworthy that when multimeric predictions involved over 20 subunits, the predicted PSM α 3 models started to unfold into sparse helices (Figures 3-4, S2-S6). Conversely, uperin 3.5 only began to display fibril-like mated helical sheets when modeled as a 25-mer or larger (Figures 3-4). To comprehensively understand these variations in multimeric order, which impact the assembly and dispersion of helices, further experimental cross- α structures are essential.

In the case of AMPs without known experimental structures, the models did not predict a β -rich form, even though experimental data showed that three out of the four tested AMPs had cross- β structures, and dolabellandin-B2 even displayed a β -rich propensity in

solution (Figures 7 & S21-S41). However, interestingly, the most tightly packed model of mated α -helical sheets that resembled the cross- α configuration among all tested peptides and multimers was the 20-mer citropin 1.3 (Figure 7). This amphibian AMP was shown by biophysical measurements to assume a β -rich fibril configuration but can be induced into an α -helical conformation in the presence of membrane lipids in solution and in the fibril form (18), similar to uperin 3.5 (15).

In addition to the observed α -helical sheets that resemble the cross- α configuration, we also frequently found predictions of α -helices clustered in a globular-like shape, particularly in the 10-mer models of numerous peptides (Figures 5, S5-S6, S9-S11, S13, S18, S20, S22, S24-S25, S27-S30, S33-S36). This phenomenon might be attributed to a bias in the training set, which is predominantly composed of globular proteins. Other recurrent shapes included spiraling helices and doughnut-like assemblies; however, these were often loosely packed and unlikely to form stable assemblies.

We observed an overall limitation in AF2's prediction of amyloid multimers, with an unexpected preference for mated α -helical sheets. Previous studies showed other types of proteins, whose AF models differed from the experimental structures, included intrinsically disordered regions (IDRs) and membrane proteins (86,87). Additionally, limitations in structure prediction were reported for some proteins that bind essential compounds for their structural integrity and function, such as heme groups, zinc ions, and metal ions. Consequently, tools like AlphaFill were developed to enrich AF models with ligands and cofactors (88). The polymorphism of amyloid fibrils might also mirror the challenges in predicting the structural dynamics of proteins undergoing allosteric-induced large conformational changes. AF generates a few models that can potentially sample structures at different locations on the energy landscape, but not necessarily capture all allosteric conformations or provide a low reliability score (86). Another factor leading to low confidence scores of AF2 models is the small number of homologous sequences. To address this issue, a new deep-learning-based algorithm, AlphaFold-Eva, was developed to evaluate AF predictions for proteins with limited homologous sequences by learning geometry information from complex structures (89). A general limitation of AI-methods trained on structural information alone may be related to the 'epigenetic dimension of protein structure,' a term that encompasses all environmental parameters affecting structures beyond the amino acid composition (90). Overall, there is still room for improvement in multimeric predictions

(91), yet future developments in this field are promising, as suggested by a recent report demonstrating that AlphaFold-Multimer successfully predicts homomeric and heteromeric interfaces (81).

In conclusion, the predictions made by AF2-ColabFold-Multimer for amyloid peptides with antimicrobial properties did not align with the more common cross- β amyloid configuration. Instead, they predominantly favored α -helical structures, including paired α -helix sheets with a hydrophobic core, akin to the cross- α configuration. Prior to initiating the structure prediction of amyloidogenic AMPs, we anticipated limitations due to their self-assembling, highly polymorphic nature, short sequence length, and the absence of homologous sequences. Surprisingly, the ability of AF2 to predict cross- α -like configurations was a positive outcome, especially given that the training set included 77 β -rich amyloids compared to a single cross- α structure, and a higher number of all- β proteins compared to all- α proteins (Table S3). Notably, the models did not predict the more common cross- β amyloid configuration, which may suggest that this preference for helices stems from the prediction of the monomeric/soluble state. We hypothesize that the training of AI-based algorithms may enable the identification of sequence-related information associated with evolutionarily driven active states, potentially leading to a more accurate simulation of the membrane-active α -helical form of many of these toxic peptides.

Methods

Structure prediction

To predict the 3-dimensional structures we used the AlphaFold collaborative “jupyter notebook” hosted by google called ColabFold (31), which allows the prediction of protein monomers, homo- and hetero- oligomeric complexes (31). We tested ten different peptides, six with known atomic structure in fibril form deposited in the PDB, and four with no experimental structure, but for which the secondary structure has been investigated using circular dichroism (CD), attenuated total internal reflection Fourier transform infrared spectroscopy (ATR-FTIR), and X-ray fiber diffraction (18). We used the default parameters of ColabFold, and varied the number of subunits, as described in Table 1. To assess the accuracy of each prediction, we employed the following AlphaFold statistics: (a) pLDDT (predicted local-distance difference test), which estimates the confidence of the prediction;

(b) sequence coverage plot, indicating the number of sequences per position during the multiple sequence alignment step; and (c) predicted alignment error (PAE), representing the expected positional error when aligning the true and predicted structures for each residue. These evaluation metrics are presented for each AlphaFold prediction we conducted in Supplementary Figures S1-S48.

Structure visualization and analysis

Visualization and analysis of the predicted and known structures were performed using UCSF ChimeraX version 1.4 (92). Interhelical and inter-sheet distances were calculated by first determining the center of mass of each subunit using the ‘measure center’ command, and then using the ‘distance’ command between selected centers of mass. To quantify the compactness of the packing, we compared the solvent accessible surface area (SASA) buried within the fibril, calculated using the ‘measure buriedarea’ command in ChimeraX. We selected a subunit located at the center of the cross- α assembly and measured the area buried within the multimer complex. The percentage of buried area was calculated from the total surface area of the selected chain, rounded to one decimal place. Values for all SASA and interhelical and inter-sheet distances are presented in Table S1.

Search for structures resembling the cross- α configuration of PSM α 3

Different helical packings derived from the PSM α 3 cross- α structure (arrangement of 4-6 helices in the same or mated sheets) were searched for structural similarity using TopSearch (93), which allows structure alignment against all PDB entries released before April 18, 2018 (93).

Searching the PDB for identical sequences with different secondary structures

All entries in wwPDB were downloaded from the PDB archive in mmCIF format. Entries were filtered for the following criteria: polypeptide(L) chain type, at least 70% of residues resolved in the atomic structure, and at least half of the modeled residues assumed a helical or extended secondary structure, as determined by the publisher of the structure, or automatically assigned by DSSP (94). Each chain that passed the filtering was classified, and 66985 helical chains and 9409 extended chains were found. Each extended chain was matched to the helical chains using Diamond (95) BLASTP with minimum sequence identity of 80% and minimum coverage of 80% for both chains. 74 matches were found, including one pair of CspA, two pairs of uperin 3.5, 14 pairs of A β , 14 pairs of glucagon, and 43 pairs

of human prion. The exact pipeline and code are available at <https://github.com/GabiAxel/pdb-similar-sequence-different-secondary-structure>. The results are given in Table S2.

Acknowledgments

We acknowledge research support from the Israel Science Foundation (Grants No. 2111/20 (M.L.) and 1764/21 (N.B.-T.)) and the Cure Alzheimer's Fund (M.L.). N.B.-T.'s research is supported in part by the Abraham E. Kazan Chair in Structural Biology, Tel Aviv University. We would like to acknowledge the use of OpenAI's GPT-3 model and DeepL write to improve the quality of writing in this study.

References

1. Huan Y, Kong Q, Mou H, Yi H. Antimicrobial Peptides: Classification, Design, Application and Research Progress in Multiple Fields. *Front Microbiol* [Internet]. 2020 Oct 16;11:582779. Available from: <http://dx.doi.org/10.3389/fmicb.2020.582779>
2. Tayeb-Fligelman E, Tabachnikov O, Moshe A, Goldshmidt-Tran O, Sawaya MR, Coquelle N, et al. The cytotoxic *Staphylococcus aureus* PSM α 3 reveals a cross- α amyloid-like fibril. *Science* [Internet]. 2017 Feb 24;355(6327):831–3. Available from: <http://dx.doi.org/10.1126/science.aaf4901>
3. Tayeb-Fligelman E, Salinas N, Tabachnikov O, Landau M. *Staphylococcus aureus* PSM α 3 Cross- α Fibril Polymorphism and Determinants of Cytotoxicity. *Structure* [Internet]. 2020 Mar 3;28(3):301–13.e6. Available from: <http://dx.doi.org/10.1016/j.str.2019.12.006>
4. Salinas N, Colletier JP, Moshe A, Landau M. Extreme amyloid polymorphism in *Staphylococcus aureus* virulent PSM α peptides. *Nat Commun* [Internet]. 2018 Aug 29;9(1):3512. Available from: <http://dx.doi.org/10.1038/s41467-018-05490-0>
5. Zhao H, Sood R, Jutila A, Bose S, Fimland G, Nissen-Meyer J, et al. Interaction of the antimicrobial peptide pheromone Plantaricin A with model membranes: implications for a novel mechanism of action. *Biochim Biophys Acta* [Internet]. 2006 Sep;1758(9):1461–74. Available from: <http://dx.doi.org/10.1016/j.bbamem.2006.03.037>
6. Sayegh RSR, Batista I de FC, Melo RL de, Riske KA, Daffre S, Montich G, et al. Longipin: An Amyloid Antimicrobial Peptide from the Harvestman *Acutisoma longipes* (Arachnida: Opiliones) with Preferential Affinity for Anionic Vesicles. *PLoS One* [Internet]. 2016 Dec 20;11(12):e0167953. Available from: <http://dx.doi.org/10.1371/journal.pone.0167953>
7. Wang M, Huang M, Zhang J, Ma Y, Li S, Wang J. A novel secretion and online-cleavage strategy for production of cecropin A in *Escherichia coli*. *Sci Rep* [Internet]. 2017 Aug 4;7(1):7368. Available from: <http://dx.doi.org/10.1038/s41598-017-07411-5>

8. Lyu Y, Fitriyanti M, Narsimhan G. Nucleation and growth of pores in 1,2-Dimyristoyl-sn-glycero-3-phosphocholine (DMPC) / cholesterol bilayer by antimicrobial peptides melittin, its mutants and cecropin P1. *Colloids Surf B Biointerfaces* [Internet]. 2019 Jan 1;173:121–7. Available from: <http://dx.doi.org/10.1016/j.colsurfb.2018.09.049>
9. Singh PK, Ghosh D, Tewari D, Mohite GM, Carvalho E, Jha NN, et al. Cytotoxic helix-rich oligomer formation by melittin and pancreatic polypeptide. *PLoS One* [Internet]. 2015 Mar 24;10(3):e0120346. Available from: <http://dx.doi.org/10.1371/journal.pone.0120346>
10. Auvynet C, El Amri C, Lacombe C, Bruston F, Bourdais J, Nicolas P, et al. Structural requirements for antimicrobial versus chemoattractant activities for dermaseptin S9. *FEBS J* [Internet]. 2008 Aug;275(16):4134–51. Available from: <http://dx.doi.org/10.1111/j.1742-4658.2008.06554.x>
11. Juhl DW, Glattard E, Lointier M, Bampilis P, Bechinger B. The Reversible Non-covalent Aggregation Into Fibers of PGLa and Magainin 2 Preserves Their Antimicrobial Activity and Synergism. *Front Cell Infect Microbiol* [Internet]. 2020 Sep 30;10:526459. Available from: <http://dx.doi.org/10.3389/fcimb.2020.526459>
12. Wang G, Narayana JL, Mishra B, Zhang Y, Wang F, Wang C, et al. Design of Antimicrobial Peptides: Progress Made with Human Cathelicidin LL-37. *Adv Exp Med Biol* [Internet]. 2019;1117:215–40. Available from: http://dx.doi.org/10.1007/978-981-13-3588-4_12
13. Domanov YA, Kinnunen PKJ. Antimicrobial peptides temporins B and L induce formation of tubular lipid protrusions from supported phospholipid bilayers. *Biophys J* [Internet]. 2006 Dec 15;91(12):4427–39. Available from: <http://dx.doi.org/10.1529/biophysj.106.091702>
14. Bücker R, Seuring C, Cazey C, Veith K, García-Alai M, Grünwald K, et al. The Cryo-EM structures of two amphibian antimicrobial cross- β amyloid fibrils. *Nat Commun* [Internet]. 2022 Jul 27;13(1):4356. Available from: <http://dx.doi.org/10.1038/s41467-022-32039-z>
15. Salinas N, Tayeb-Fligelman E, Sammito MD, Bloch D, Jelinek R, Noy D, et al. The amphibian antimicrobial peptide uperin 3.5 is a cross- α /cross- β chameleon functional amyloid. *Proc Natl Acad Sci U S A* [Internet]. 2021 Jan 19;118(3). Available from: <http://dx.doi.org/10.1073/pnas.2014442118>
16. Zhao H, Jutila A, Nurminen T, Wickström SA, Keski-Oja J, Kinnunen PKJ. Binding of endostatin to phosphatidylserine-containing membranes and formation of amyloid-like fibers. *Biochemistry* [Internet]. 2005 Mar 1;44(8):2857–63. Available from: <http://dx.doi.org/10.1021/bi048510j>
17. Engelberg Y, Landau M. The Human LL-37(17-29) antimicrobial peptide reveals a functional supramolecular structure [Internet]. Vol. 11, *Nature Communications*. 2020. Available from: <http://dx.doi.org/10.1038/s41467-020-17736-x>
18. Ragonis-Bachar P, Rayan B, Barnea E, Engelberg Y, Upcher A, Landau M. Natural Antimicrobial Peptides Self-assemble as α/β Chameleon Amyloids. *Biomacromolecules* [Internet]. 2022 Sep 12;23(9):3713–27. Available from: <http://dx.doi.org/10.1021/acs.biomac.2c00582>
19. Gour S, Kumar V, Singh A, Gadhave K, Goyal P, Pandey J, et al. Mammalian antimicrobial peptide protegrin-4 self assembles and forms amyloid-like aggregates: Assessment of its functional relevance. *J Pept Sci* [Internet]. 2019 Mar;25(3):e3151. Available from: <http://dx.doi.org/10.1002/psc.3151>
20. Wang CK, King GJ, Conibear AC, Ramos MC, Chaousis S, Henriques ST, et al. Mirror Images

of Antimicrobial Peptides Provide Reflections on Their Functions and Amyloidogenic Properties. *J Am Chem Soc* [Internet]. 2016 May 4;138(17):5706–13. Available from: <http://dx.doi.org/10.1021/jacs.6b02575>

21. Baltutis V, O’Leary PD, Martin LL. Self-Assembly of Linear, Natural Antimicrobial Peptides: An Evolutionary Perspective. *Chempluschem* [Internet]. 2022 Oct 5;e202200240. Available from: <http://dx.doi.org/10.1002/cplu.202200240>
22. Eisenberg DS, Sawaya MR. Structural Studies of Amyloid Proteins at the Molecular Level [Internet]. Vol. 86, Annual Review of Biochemistry. 2017. p. 69–95. Available from: <http://dx.doi.org/10.1146/annurev-biochem-061516-045104>
23. Ragonis-Bachar P, Landau M. Functional and pathological amyloid structures in the eyes of 2020 cryo-EM. *Curr Opin Struct Biol* [Internet]. 2021 Jun;68:184–93. Available from: <http://dx.doi.org/10.1016/j.sbi.2021.01.006>
24. Chiti F, Dobson CM. Protein misfolding, functional amyloid, and human disease. *Annu Rev Biochem* [Internet]. 2006;75:333–66. Available from: <http://dx.doi.org/10.1146/annurev.biochem.75.101304.123901>
25. Pinheiro F, Santos J, Ventura S. AlphaFold and the amyloid landscape. *J Mol Biol* [Internet]. 2021 Oct 1;433(20):167059. Available from: <http://dx.doi.org/10.1016/j.jmb.2021.167059>
26. Zielinski M, Röder C, Schröder GF. Challenges in sample preparation and structure determination of amyloids by cryo-EM. *J Biol Chem* [Internet]. 2021 Aug;297(2):100938. Available from: <http://dx.doi.org/10.1016/j.jbc.2021.100938>
27. Serpell LC, Radford SE, Otzen DE. AlphaFold: A Special Issue and A Special Time for Protein Science. *J Mol Biol* [Internet]. 2021 Oct 1;433(20):167231. Available from: <http://dx.doi.org/10.1016/j.jmb.2021.167231>
28. Integrative Structural Biology in the Era of Accurate Structure Prediction. *J Mol Biol* [Internet]. 2021 Oct 1 [cited 2022 Nov 28];433(20):167127. Available from: <http://dx.doi.org/10.1016/j.jmb.2021.167127>
29. Zhang O, Haghighatlari M, Li J, Liu ZH, Namini A, Teixeira JMC, et al. Learning to evolve structural ensembles of unfolded and disordered proteins using experimental solution data. *J Chem Phys* [Internet]. 2023 May 7;158(17). Available from: <http://dx.doi.org/10.1063/5.0141474>
30. Jumper J, Evans R, Pritzel A, Green T, Figurnov M, Ronneberger O, et al. Highly accurate protein structure prediction with AlphaFold. *Nature* [Internet]. 2021 Aug;596(7873):583–9. Available from: <http://dx.doi.org/10.1038/s41586-021-03819-2>
31. Mirdita M, Schütze K, Moriwaki Y, Heo L, Ovchinnikov S, Steinegger M. ColabFold: making protein folding accessible to all. *Nat Methods* [Internet]. 2022 Jun;19(6):679–82. Available from: <http://dx.doi.org/10.1038/s41592-022-01488-1>
32. Kumar DKV, Choi SH, Washicosky KJ, Eimer WA, Tucker S, Ghofrani J, et al. Amyloid- β peptide protects against microbial infection in mouse and worm models of Alzheimer’s disease. *Sci Transl Med* [Internet]. 2016 May 25;8(340):340ra72. Available from: <http://dx.doi.org/10.1126/scitranslmed.aaf1059>
33. Soscia SJ, Kirby JE, Washicosky KJ, Tucker SM, Ingelsson M, Hyman B, et al. The Alzheimer’s disease-associated amyloid beta-protein is an antimicrobial peptide. *PLoS One* [Internet]. 2010 Mar 3;5(3):e9505. Available from:

<http://dx.doi.org/10.1371/journal.pone.0009505>

34. Bourgade K, Garneau H, Giroux G, Le Page AY, Bocti C, Dupuis G, et al. β -Amyloid peptides display protective activity against the human Alzheimer's disease-associated herpes simplex virus-1. *Biogerontology* [Internet]. 2015 Feb;16(1):85–98. Available from: <http://dx.doi.org/10.1007/s10522-014-9538-8>
35. White MR, Kandel R, Tripathi S, Condon D, Qi L, Taubenberger J, et al. Alzheimer's Associated β -Amyloid Protein Inhibits Influenza A Virus and Modulates Viral Interactions with Phagocytes [Internet]. Vol. 9, PLoS ONE. 2014. p. e101364. Available from: <http://dx.doi.org/10.1371/journal.pone.0101364>
36. Wang L, Liu Q, Chen JC, Cui YX, Zhou B, Chen YX, et al. Antimicrobial activity of human islet amyloid polypeptides: an insight into amyloid peptides' connection with antimicrobial peptides. *Biol Chem* [Internet]. 2012 Jul;393(7):641–6. Available from: <http://dx.doi.org/10.1515/hsz-2012-0107>
37. Kreutzberger MAB, Wang S, Beltran LC, Tuachi A, Zuo X, Egelman EH, et al. Phenol-soluble modulins PSM α 3 and PSM β 2 form nanotubes that are cross- α amyloids. *Proc Natl Acad Sci U S A* [Internet]. 2022 May 17;119(20):e2121586119. Available from: <http://dx.doi.org/10.1073/pnas.2121586119>
38. Zhang SQ, Huang H, Yang J, Kratochvil HT, Lolicato M, Liu Y, et al. Designed peptides that assemble into cross- α amyloid-like structures. *Nat Chem Biol* [Internet]. 2018 Sep;14(9):870–5. Available from: <http://dx.doi.org/10.1038/s41589-018-0105-5>
39. Calabrese AN, Liu Y, Wang T, Musgrave IF, Pukala TL, Tabor RF, et al. The Amyloid Fibril-Forming Properties of the Amphibian Antimicrobial Peptide Uperin 3.5. *Chembiochem* [Internet]. 2016 Feb 2;17(3):239–46. Available from: <http://dx.doi.org/10.1002/cbic.201500518>
40. Martin LL, Kubeil C, Piantavigna S, Tikkoo T, Gray NP, John T, et al. Amyloid aggregation and membrane activity of the antimicrobial peptide uperin 3.5 [Internet]. Vol. 110, Peptide Science. 2018. p. e24052. Available from: <http://dx.doi.org/10.1002/pep2.24052>
41. John T, Dealey TJA, Gray NP, Patil NA, Hossain MA, Abel B, et al. The Kinetics of Amyloid Fibrillar Aggregation of Uperin 3.5 Is Directed by the Peptide's Secondary Structure. *Biochemistry* [Internet]. 2019 Sep 3;58(35):3656–68. Available from: <http://dx.doi.org/10.1021/acs.biochem.9b00536>
42. John T, Greene GW, Patil NA, Dealey TJA, Hossain MA, Abel B, et al. Adsorption of Amyloidogenic Peptides to Functionalized Surfaces Is Biased by Charge and Hydrophilicity. *Langmuir* [Internet]. 2019 Nov 12;35(45):14522–31. Available from: <http://dx.doi.org/10.1021/acs.langmuir.9b02063>
43. Luo F, Gui X, Zhou H, Gu J, Li Y, Liu X, et al. Atomic structures of FUS LC domain segments reveal bases for reversible amyloid fibril formation. *Nat Struct Mol Biol* [Internet]. 2018 Apr;25(4):341–6. Available from: <http://dx.doi.org/10.1038/s41594-018-0050-8>
44. Murray DT, Kato M, Lin Y, Thurber KR, Hung I, McKnight SL, et al. Structure of FUS Protein Fibrils and Its Relevance to Self-Assembly and Phase Separation of Low-Complexity Domains. *Cell* [Internet]. 2017 Oct 19;171(3):615–27.e16. Available from: <http://dx.doi.org/10.1016/j.cell.2017.08.048>
45. Lu J, Cao Q, Hughes MP, Sawaya MR, Boyer DR, Cascio D, et al. CryoEM structure of the low-complexity domain of hnRNPA2 and its conversion to pathogenic amyloid. *Nat Commun* [Internet]. 2020 Aug 14;11(1):4090. Available from: <http://dx.doi.org/10.1038/s41467-020->

46. Cao Q, Boyer DR, Sawaya MR, Ge P, Eisenberg DS. Cryo-EM structures of four polymorphic TDP-43 amyloid cores. *Nat Struct Mol Biol* [Internet]. 2019 Jul;26(7):619–27. Available from: <http://dx.doi.org/10.1038/s41594-019-0248-4>
47. Lee M, Ghosh U, Thurber KR, Kato M, Tycko R. Molecular structure and interactions within amyloid-like fibrils formed by a low-complexity protein sequence from FUS. *Nat Commun* [Internet]. 2020 Nov 12;11(1):5735. Available from: <http://dx.doi.org/10.1038/s41467-020-19512-3>
48. Hughes MP, Sawaya MR, Boyer DR, Goldschmidt L, Rodriguez JA, Cascio D, et al. Atomic structures of low-complexity protein segments reveal kinked β sheets that assemble networks. *Science* [Internet]. 2018 Feb 9;359(6376):698–701. Available from: <http://dx.doi.org/10.1126/science.aan6398>
49. Xhindoli D, Pacor S, Benincasa M, Scocchi M, Gennaro R, Tossi A. The human cathelicidin LL-37--A pore-forming antibacterial peptide and host-cell modulator. *Biochim Biophys Acta* [Internet]. 2016 Mar;1858(3):546–66. Available from: <http://dx.doi.org/10.1016/j.bbamem.2015.11.003>
50. Sood R, Domanov Y, Pietiäinen M, Kontinen VP, Kinnunen PKJ. Binding of LL-37 to model biomembranes: insight into target vs host cell recognition. *Biochim Biophys Acta* [Internet]. 2008 Apr;1778(4):983–96. Available from: <http://dx.doi.org/10.1016/j.bbamem.2007.11.016>
51. Shahmiri M, Enciso M, Adda CG, Smith BJ, Perugini MA, Mechler A. Membrane Core-Specific Antimicrobial Action of Cathelicidin LL-37 Peptide Switches Between Pore and Nanofibre Formation. *Sci Rep* [Internet]. 2016 Nov 30;6:38184. Available from: <http://dx.doi.org/10.1038/srep38184>
52. Rajasekaran G, Kim EY, Shin SY. LL-37-derived membrane-active FK-13 analogs possessing cell selectivity, anti-biofilm activity and synergy with chloramphenicol and anti-inflammatory activity. *Biochim Biophys Acta Biomembr* [Internet]. 2017 May;1859(5):722–33. Available from: <http://dx.doi.org/10.1016/j.bbamem.2017.01.037>
53. Wegener KL, Wabnitz PA, Carver JA, Bowie JH, Chia BC, Wallace JC, et al. Host defence peptides from the skin glands of the Australian blue mountains tree-frog *Litoria citropa*. Solution structure of the antibacterial peptide citropin 1.1. *Eur J Biochem* [Internet]. 1999 Oct;265(2):627–37. Available from: <http://dx.doi.org/10.1046/j.1432-1327.1999.00750.x>
54. Mignogna G, Simmaco M, Kreil G, Barra D. Antibacterial and haemolytic peptides containing D-alloisoleucine from the skin of *Bombina variegata*. *EMBO J* [Internet]. 1993 Dec;12(12):4829–32. Available from: <http://dx.doi.org/10.1002/j.1460-2075.1993.tb06172.x>
55. Asoodeh A, Azam AG, Chamani J. Identification and Characterization of Novel Antibacterial Peptides from Skin Secretions of *Euphlyctis cyanophlyctis* [Internet]. Vol. 18, *International Journal of Peptide Research and Therapeutics*. 2012. p. 107–15. Available from: <http://dx.doi.org/10.1007/s10989-011-9284-6>
56. Iijima R, Kisugi J, Yamazaki M. A novel antimicrobial peptide from the sea hare *Dolabella auricularia*. *Dev Comp Immunol* [Internet]. 2003 Apr;27(4):305–11. Available from: [http://dx.doi.org/10.1016/s0145-305x\(02\)00105-2](http://dx.doi.org/10.1016/s0145-305x(02)00105-2)
57. Chen GF, Xu TH, Yan Y, Zhou YR, Jiang Y, Melcher K, et al. Amyloid beta: structure, biology and structure-based therapeutic development. *Acta Pharmacol Sin* [Internet]. 2017 Sep;38(9):1205–35. Available from: <http://dx.doi.org/10.1038/aps.2017.28>

58. Röder C, Kupreichyk T, Gremer L, Schäfer LU, Pothula KR, Ravelli RBG, et al. Cryo-EM structure of islet amyloid polypeptide fibrils reveals similarities with amyloid- β fibrils. *Nat Struct Mol Biol* [Internet]. 2020 Jul;27(7):660–7. Available from: <http://dx.doi.org/10.1038/s41594-020-0442-4>
59. Williamson JA, Loria JP, Miranker AD. Helix stabilization precedes aqueous and bilayer-catalyzed fiber formation in islet amyloid polypeptide. *J Mol Biol* [Internet]. 2009 Oct 23;393(2):383–96. Available from: <http://dx.doi.org/10.1016/j.jmb.2009.07.077>
60. Zhang S, Iwata K, Lachenmann MJ, Peng JW, Li S, Stimson ER, et al. The Alzheimer's peptide α adopts a collapsed coil structure in water. *J Struct Biol* [Internet]. 2000 Jun;130(2-3):130–41. Available from: <http://dx.doi.org/10.1006/jsbi.2000.4288>
61. Kirkitadze MD, Condrón MM, Teplow DB. Identification and characterization of key kinetic intermediates in amyloid β -protein fibrillogenesis. *J Mol Biol* [Internet]. 2001 Oct 5;312(5):1103–19. Available from: <http://dx.doi.org/10.1006/jmbi.2001.4970>
62. Nanga RPR, Brender JR, Vivekanandan S, Ramamoorthy A. Structure and membrane orientation of IAPP in its natively amidated form at physiological pH in a membrane environment. *Biochim Biophys Acta* [Internet]. 2011 Oct;1808(10):2337–42. Available from: <http://dx.doi.org/10.1016/j.bbamem.2011.06.012>
63. Shao H, Jao S, Ma K, Zagorski MG. Solution structures of micelle-bound amyloid β -(1-40) and β -(1-42) peptides of Alzheimer's disease. *J Mol Biol* [Internet]. 1999 Jan 15;285(2):755–73. Available from: <http://dx.doi.org/10.1006/jmbi.1998.2348>
64. White MR, Kandel R, Tripathi S, Condon D, Qi L, Taubenberger J, et al. Alzheimer's Associated β -Amyloid Protein Inhibits Influenza A Virus and Modulates Viral Interactions with Phagocytes [Internet]. Vol. 9, PLoS ONE. 2014. p. e101364. Available from: <http://dx.doi.org/10.1371/journal.pone.0101364>
65. Brunette TJ, Parmeggiani F, Huang PS, Bhabha G, Ekiert DC, Tsutakawa SE, et al. Exploring the repeat protein universe through computational protein design. *Nature* [Internet]. 2015 Dec 24;528(7583):580–4. Available from: <http://dx.doi.org/10.1038/nature16162>
66. Fallas JA, Ueda G, Sheffler W, Nguyen V, McNamara DE, Sankaran B, et al. Computational design of self-assembling cyclic protein homo-oligomers. *Nat Chem* [Internet]. 2017 Apr;9(4):353–60. Available from: <http://dx.doi.org/10.1038/nchem.2673>
67. Xiang K, Nagaike T, Xiang S, Kilic T, Beh MM, Manley JL, et al. Crystal structure of the human symplekin-Ssu72-CTD phosphopeptide complex. *Nature* [Internet]. 2010 Oct 7;467(7316):729–33. Available from: <http://dx.doi.org/10.1038/nature09391>
68. Baradaran R, Berrisford JM, Minhas GS, Sazanov LA. Crystal structure of the entire respiratory complex I. *Nature* [Internet]. 2013 Feb 28;494(7438):443–8. Available from: <http://www.nature.com/articles/nature11871>
69. Lucas M, Gershlick DC, Vidaurrezaga A, Rojas AL, Bonifacino JS, Hierro A. Structural Mechanism for Cargo Recognition by the Retromer Complex. *Cell* [Internet]. 2016 Dec 1;167(6):1623–35.e14. Available from: <http://dx.doi.org/10.1016/j.cell.2016.10.056>
70. Sun S, Gao S, Kondabagil K, Xiang Y, Rossmann MG, Rao VB. Structure and function of the small terminase component of the DNA packaging machine in T4-like bacteriophages. *Proc Natl Acad Sci U S A* [Internet]. 2012 Jan 17;109(3):817–22. Available from: <http://dx.doi.org/10.1073/pnas.1110224109>

71. Subramanian R, Ti SC, Tan L, Darst SA, Kapoor TM. Marking and measuring single microtubules by PRC1 and kinesin-4. *Cell* [Internet]. 2013 Jul 18;154(2):377–90. Available from: <http://dx.doi.org/10.1016/j.cell.2013.06.021>
72. Crescenzi O, Tomaselli S, Guerrini R, Salvadori S, D'Ursi AM, Temussi PA, et al. Solution structure of the Alzheimer amyloid beta-peptide (1-42) in an apolar microenvironment. Similarity with a virus fusion domain. *Eur J Biochem* [Internet]. 2002 Nov;269(22):5642–8. Available from: <http://dx.doi.org/10.1046/j.1432-1033.2002.03271.x>
73. Xiao Y, Ma B, McElheny D, Parthasarathy S, Long F, Hoshi M, et al. A β (1-42) fibril structure illuminates self-recognition and replication of amyloid in Alzheimer's disease. *Nat Struct Mol Biol* [Internet]. 2015 Jun;22(6):499–505. Available from: <http://dx.doi.org/10.1038/nsmb.2991>
74. Zhang Y, Swietnicki W, Zagorski MG, Surewicz WK, Sönnichsen FD. Solution structure of the E200K variant of human prion protein. Implications for the mechanism of pathogenesis in familial prion diseases. *J Biol Chem* [Internet]. 2000 Oct 27;275(43):33650–4. Available from: <http://dx.doi.org/10.1074/jbc.C000483200>
75. Manka SW, Zhang W, Wenborn A, Betts J, Joiner S, Saibil HR, et al. 2.7 Å cryo-EM structure of ex vivo RML prion fibrils. *Nat Commun* [Internet]. 2022 Jul 13;13(1):4004. Available from: <http://dx.doi.org/10.1038/s41467-022-30457-7>
76. Sasaki K, Dockerill S, Adamiak DA, Tickle IJ, Blundell T. X-ray analysis of glucagon and its relationship to receptor binding. *Nature* [Internet]. 1975 Oct 30;257(5529):751–7. Available from: <http://dx.doi.org/10.1038/257751a0>
77. Gelenter MD, Smith KJ, Liao SY, Mandala VS, Dregni AJ, Lamm MS, et al. The peptide hormone glucagon forms amyloid fibrils with two coexisting β -strand conformations. *Nat Struct Mol Biol* [Internet]. 2019 Jul;26(7):592–8. Available from: <http://dx.doi.org/10.1038/s41594-019-0238-6>
78. Tang Y, Schneider WM, Shen Y, Raman S, Inouye M, Baker D, et al. Fully automated high-quality NMR structure determination of small (2)H-enriched proteins. *J Struct Funct Genomics* [Internet]. 2010 Dec;11(4):223–32. Available from: <http://dx.doi.org/10.1007/s10969-010-9095-6>
79. Agirrezabala X, Samatova E, Macher M, Liutkute M, Maiti M, Gil-Carton D, et al. A switch from α -helical to β -strand conformation during co-translational protein folding. *EMBO J* [Internet]. 2022 Feb 15;41(4):e109175. Available from: <http://dx.doi.org/10.15252/embj.2021109175>
80. Sheng J, Olrichs NK, Gadella BM, Kaloyanova DV, Helms JB. Regulation of Functional Protein Aggregation by Multiple Factors: Implications for the Amyloidogenic Behavior of the CAP Superfamily Proteins. *Int J Mol Sci* [Internet]. 2020 Sep 7 [cited 2022 Nov 28];21(18):6530. Available from: <https://www.mdpi.com/1422-0067/21/18/6530>
81. Evans R, O'Neill M, Pritzel A, Antropova N, Senior A, Green T, et al. Protein complex prediction with AlphaFold-Multimer [Internet]. *bioRxiv*. 2022 [cited 2023 Aug 27]. p. 2021.10.04.463034. Available from: <https://www.biorxiv.org/content/10.1101/2021.10.04.463034v2>
82. Kollmer M, Close W, Funk L, Rasmussen J, Bsoul A, Schierhorn A, et al. Cryo-EM structure and polymorphism of A β amyloid fibrils purified from Alzheimer's brain tissue. *Nat Commun* [Internet]. 2019 Oct 29;10(1):4760. Available from: <http://dx.doi.org/10.1038/s41467-019-12683-8>

83. Varadi M, De Baets G, Vranken WF, Tompa P, Pancsa R. AmyPro: a database of proteins with validated amyloidogenic regions. *Nucleic Acids Res* [Internet]. 2018 [cited 2021 Jul 24];46(D1):D387–92. Available from: <https://doi.org/10.1093/nar/gkx950>
84. Cracchiolo OM, Edun DN, Betti VM, Goldberg JM, Serrano AL. Cross- α/β polymorphism of PSM α 3 fibrils. *Proc Natl Acad Sci U S A* [Internet]. 2022 Feb 1;119(5). Available from: <http://dx.doi.org/10.1073/pnas.2114923119>
85. Zaman M, Andreasen M. Modulating Kinetics of the Amyloid-Like Aggregation of *S. aureus* Phenol-Soluble Modulins by Changes in pH. *Microorganisms* [Internet]. 2021 Jan 7;9(1). Available from: <http://dx.doi.org/10.3390/microorganisms9010117>
86. Nussinov R, Zhang M, Liu Y, Jang H. AlphaFold, Artificial Intelligence (AI), and Allostery. *J Phys Chem B* [Internet]. 2022 Sep 1;126(34):6372–83. Available from: <http://dx.doi.org/10.1021/acs.jpcb.2c04346>
87. Del Alamo D, Sala D, Mchaourab HS, Meiler J. Sampling alternative conformational states of transporters and receptors with AlphaFold2. *Elife* [Internet]. 2022 Mar 3;11. Available from: <http://dx.doi.org/10.7554/eLife.75751>
88. Hekkelman ML, de Vries I, Joosten RP, Perrakis A. AlphaFill: enriching AlphaFold models with ligands and cofactors. *Nat Methods* [Internet]. 2023 Feb;20(2):205–13. Available from: <http://dx.doi.org/10.1038/s41592-022-01685-y>
89. Xiong H, Han L, Wang Y, Chai P. Evaluating the Reliability of AlphaFold 2 for Unknown Complex Structures with Deep Learning [Internet]. *bioRxiv*. 2022 [cited 2023 Aug 27]. p. 2022.07.08.499384. Available from: <https://www.biorxiv.org/content/10.1101/2022.07.08.499384v1.abstract>
90. Azzaz F, Yahi N, Chahinian H, Fantini J. The Epigenetic Dimension of Protein Structure Is an Intrinsic Weakness of the AlphaFold Program. *Biomolecules* [Internet]. 2022 Oct 20;12(10). Available from: <http://dx.doi.org/10.3390/biom12101527>
91. Zhu W, Shenoy A, Kundrotas P, Elofsson A. Evaluation of AlphaFold-Multimer prediction on multi-chain protein complexes. *Bioinformatics* [Internet]. 2023 Jul 1;39(7). Available from: <http://dx.doi.org/10.1093/bioinformatics/btad424>
92. Pettersen EF, Goddard TD, Huang CC, Meng EC, Couch GS, Croll TI, et al. UCSF ChimeraX: Structure visualization for researchers, educators, and developers. *Protein Sci* [Internet]. 2021 Jan;30(1):70–82. Available from: <http://dx.doi.org/10.1002/pro.3943>
93. Structure-Based Characterization of Multiprotein Complexes. *Structure* [Internet]. 2014 Jul 8 [cited 2023 Mar 8];22(7):1063–70. Available from: <http://dx.doi.org/10.1016/j.str.2014.05.005>
94. Kabsch W, Sander C. Dictionary of protein secondary structure: pattern recognition of hydrogen-bonded and geometrical features. *Biopolymers* [Internet]. 1983 Dec;22(12):2577–637. Available from: <http://dx.doi.org/10.1002/bip.360221211>
95. Buchfink B, Reuter K, Drost HG. Sensitive protein alignments at tree-of-life scale using DIAMOND. *Nat Methods* [Internet]. 2021 Apr 7 [cited 2023 Mar 8];18(4):366–8. Available from: <https://www.nature.com/articles/s41592-021-01101-x>