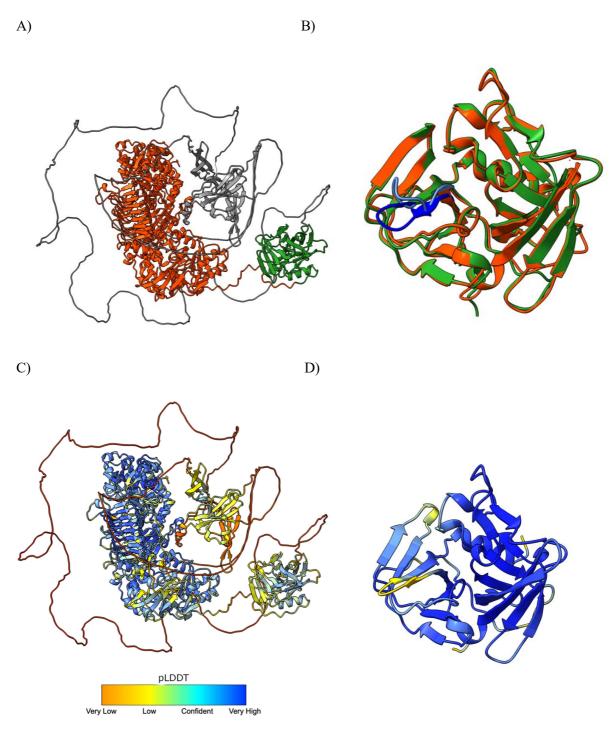


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**Supporting information for article:** 

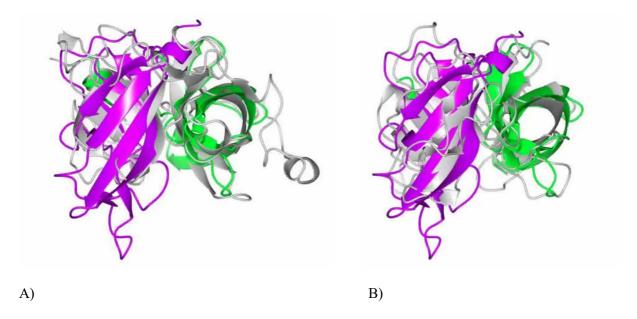
The structure of the Gemella haemolysans M26 IgA1 protease trypsin-like domain

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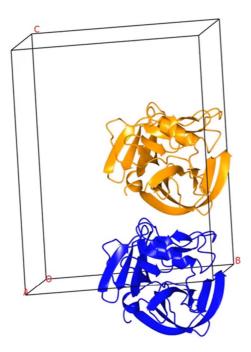


**Figure S1** A comparison of the AlphaFold3-predicted model of *GhIgAP* with the *GhTrp* crystal structure. The Google webserver (https://alphafoldserver.com) for AlphaFold3 was used to generate the full-length model using default settings.A) the complete modelled structure of GhIgAP with the IgAP domain rendered in orange and the trypsin domain in green. The remainder of the N-terminal structure is rendered in grey. B) superimposition of the GhTrp domain from the AlphaFold model (green) with the crystallographic structure (orange). Loop 2 in the modelled and crystallographic structures is rendered in in dark and light blue, respectively. The two structures align well with a Ca RMSD of 1.4Å. The complete GhIgAP (C) and the GhTrp domain (D) AlphaFold3 predicted

structures colored by pLDDT values. The color key corresponds to Very high (pLDDT > 90), Confident (90 > pLDDT > 70), Low (70 > pLDDT > 50), Very low (pLDDT < 50).



**Figure S2** Superimposition of *GhTrp* with previously unsuccessful molecular-replacement search models. The N-terminal (purple) and C-terminal (green) lobes of the GhTrp trypsin-like fold are superimposed with A) the highest sequence identity model at the time (31.2% identity; Cα RMSD of 2.46 Å) and B) the I-TASSER homology model (C $\alpha$  RMSD of 3.03 Å). These search models are shown in gray.



**Figure S3** Asymmetric unit of the *GhTrp* in the context of the unit cell. The two molecules of the GhTrp asymmetric unit are shown in blue and orange. They are related to each other by translational non-crystallographic symmetry along the c-axis of the unit cell.