

Figure S1: Cryo-EM micrograph, reconstruction and FSC calculation (a): Cryo-EM micrograph of HAd7K/EC23. (b): 2D class averages of HAd7K/EC23. (c, e): Isosurface representations of the cryo-EM 3D reconstructions of HAd7K-(EC23)₃ (c) and HAd7K-(EC23)₂ (e) colored by monomer. (d, f): Fourier Shell Correlation (FSC) curves for HAd7K-(EC23)₃ (d) and HAd7K-(EC23)₂ (f).



Figure S2: Zoomed views of the 3D reconstructions. Views on 3 different areas of HAd7K-(EC23)³ (left column) and HAd7K-(EC23)² (right column). The cryo-EM density is represented in light grey. First row focuses on one β strand in HAd7K, second row on the interaction between EC2 (salmon) and HAd7K (blue) and third row on the interaction between EC3 (salmon) and HAd7K (pink).

Table S1. Summary	v of data	collection	and ato	omic model	statistics
	,	concentor			

Data collection						
Microscope	Krios G3 (Thermo	Fischer Scientific)				
Voltage (kV)	300					
Magnification	215,000x					
Unbinned pixel size	0.325 Å/pixel					
Camera	K2 Summit (Gatan Inc)					
Exposure time	3s					
Number of frames	30					
Total dose (e ⁻ /Ų)	35					
Image processing						
	Had7k-(EC23)2	Had7k-(EC23)3				
EMDB						
Symmetry	C1	C3				
Final number of Particles	97,190	58,219				
Map resolution in Å (FSC 0.143)	3.3	3.1				
Model statistics						
PDB						
Model resolution in Å (FSC 0.5)	3.7	3.4				

Ramachandran favored (%)	95.8	96.0
Ramachandran outliers (%)	0.2	0.8
Rotamer outliers (%)	0	0
C-beta deviations	0	0
Rms on bond lengths	0.0072	0.0063
Rms on bond angles	0.78	0.88
Clashscore	6.95	5.29
Molprobity score	1.67	1.56