

diseases

• keratin genetic mutations

↳ blisters in rats

↳ epidermolysis bullosa simplex (Human)

• Amyotrophic lateral sclerosis (ALS)

↳ due to problems in NF, keratins, ABP, MUP

• in cell cytoplasm soft

• nails & hair ↑ S-S ↑ strength

like keratin

\* vimentin & lamins are regulated via phosphorylation at termini → leads to dissociation at ends.

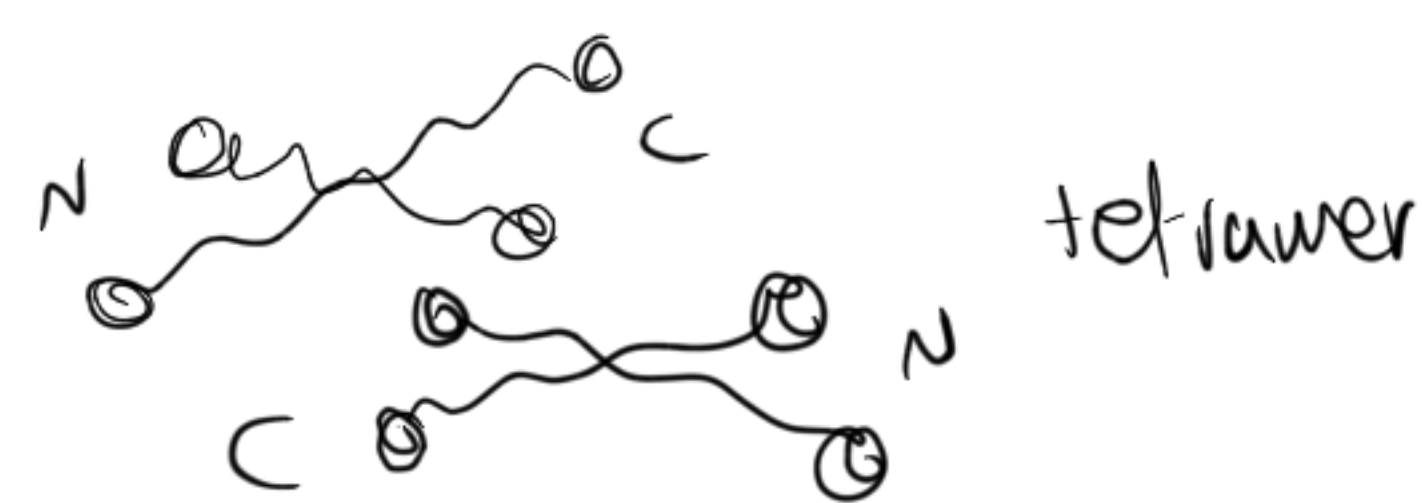
Structure

- Polypeptide
- α-helical center rod
- S & C, N termini → their diff. size

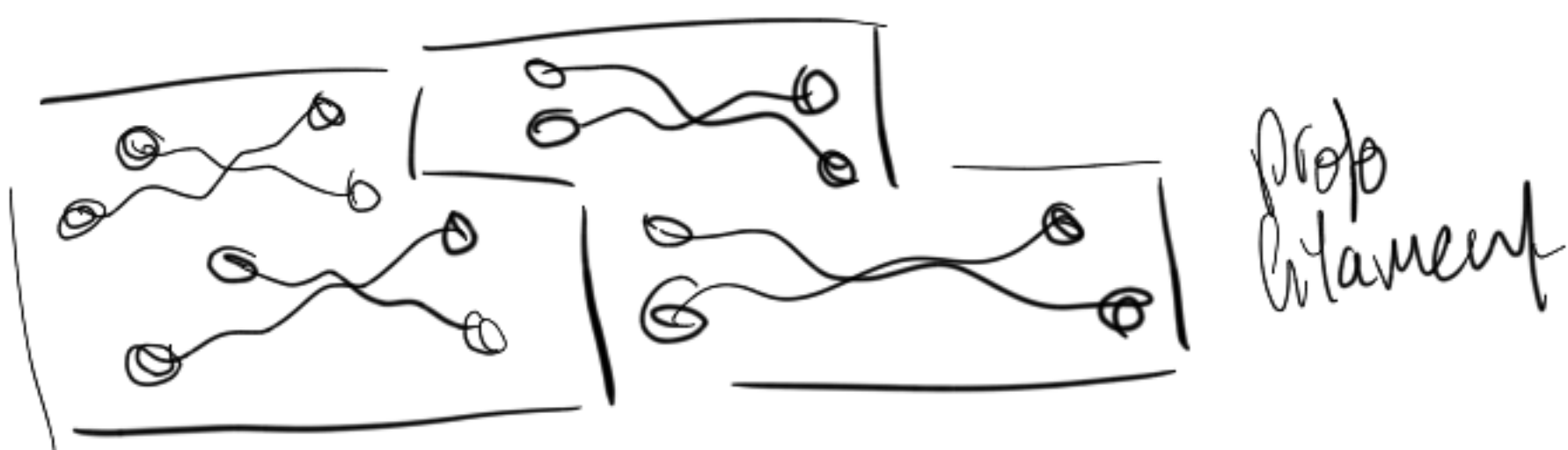
dimerize by coiling



associate in a staggered manner (anti-parallel & not exactly on top of each other)



Associate end-to-end



↓ 8 of them wound together  
filament \* rope-like \* twisted.

cuz intermediate size

Intermediate filaments

• functions

① provides mechanical strength

② Scaffold that integrates AF & MT

③ organize internal structure of cell (mainly nucleus)

• Not dynamic → stable unlike MT & AF

• Types

I acidic epithelium

II basic

III \* vimentin → SM, fibroblasts, WBCs, mesenchyme

III desmin → muscles

IV NF → neuron axon

V \* lamins → Nuclear envelope component

nucleus } \* keratin } anchoring NE  
                  \* vimentin }  
                  \* lamins → component of NE

muscle - desmin connects actin filaments together & to the PM → synchronise contractility

neurons - NF bridge actin filaments & MT together to provide stability

epithelium - desmosomes } IF connect to cadherin-like transmembrane proteins & intermediate binding proteins  
                  • cell-cell }  
                  • like adherens junctions }

- hemidesmosomes } IF connect to integrins (transmembrane receptors) & intermediate fil. binding protein  
• Cell-substratum }  
• like focal adhesion }