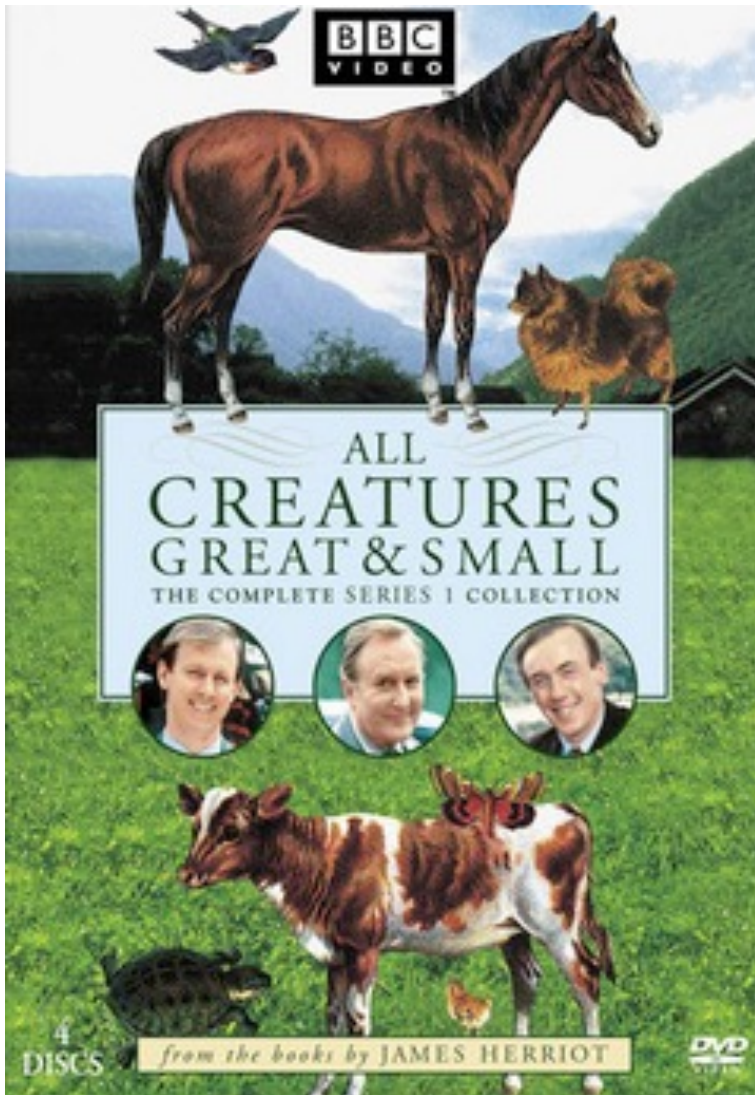


All RNAs great and small



Nascent transcripts

Co-transcriptional and post-transcriptional processes

Gene loops and Rloops

Splicing

3' end formation

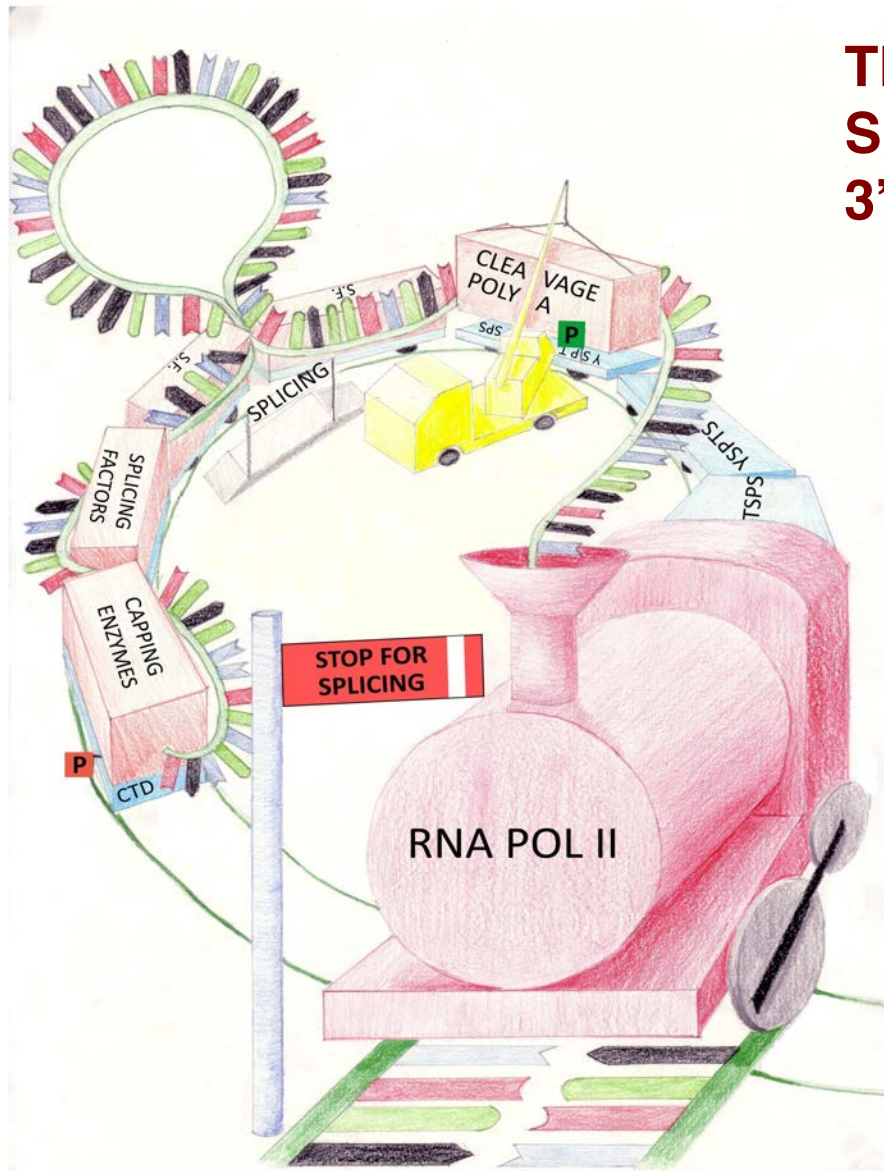
Translation cycle

RNA enzymes and complexes

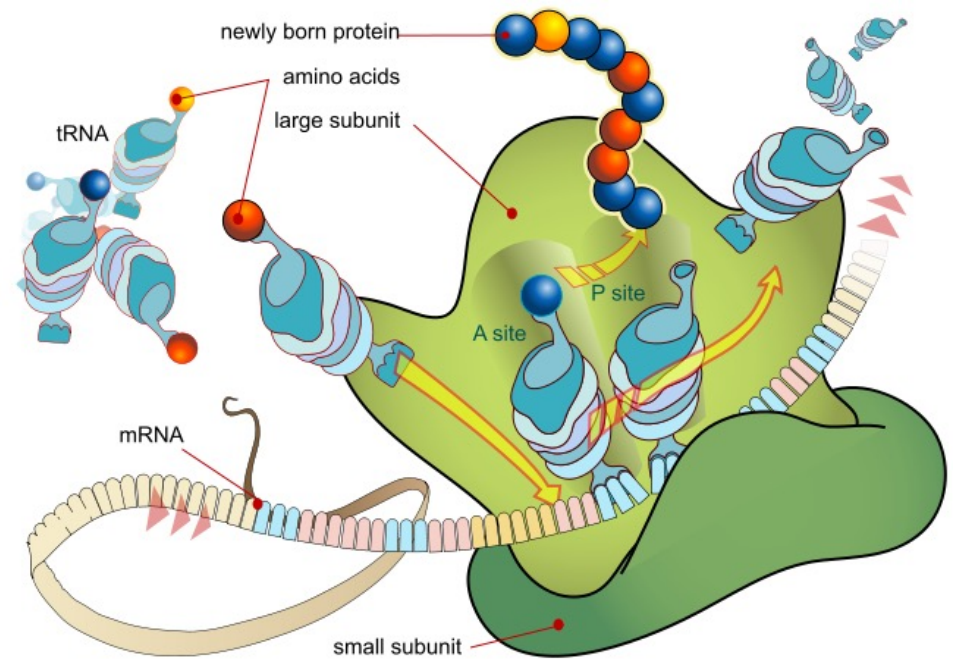


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University of Warsaw

RNA MACHINERIES

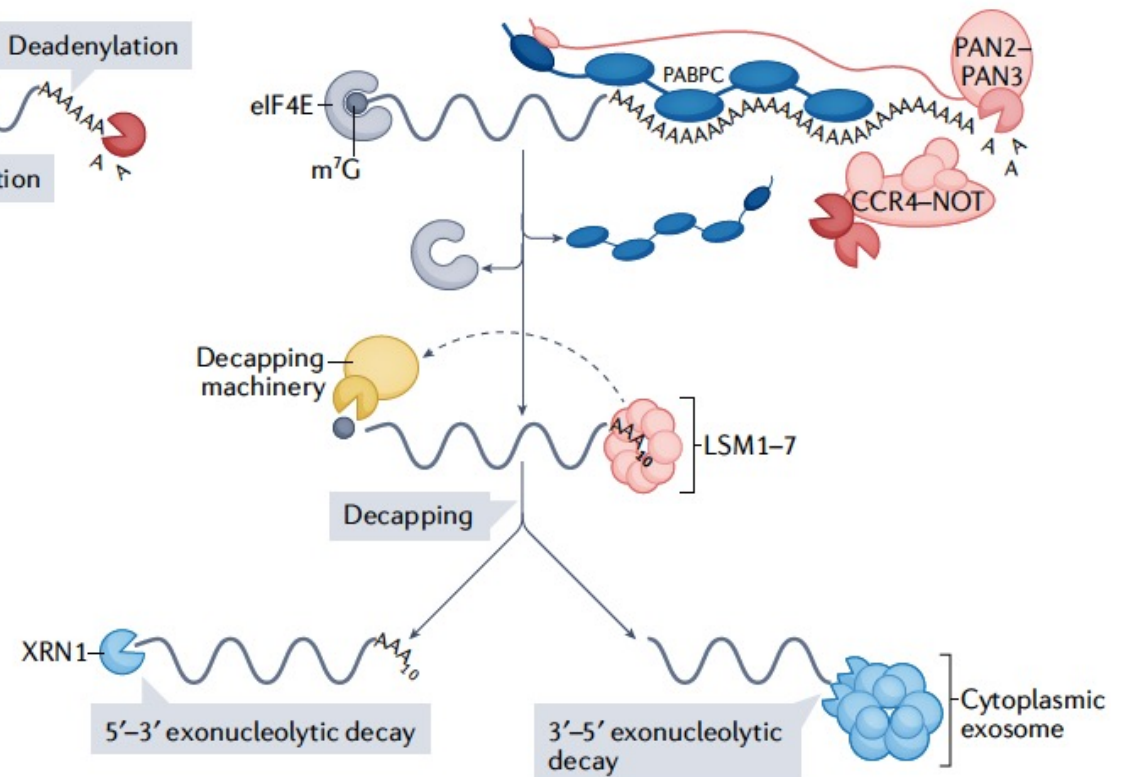
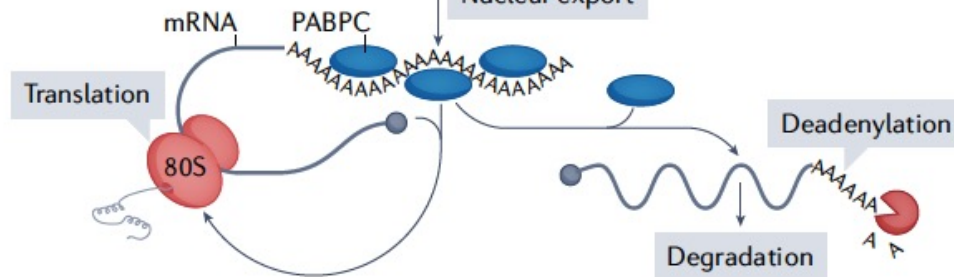
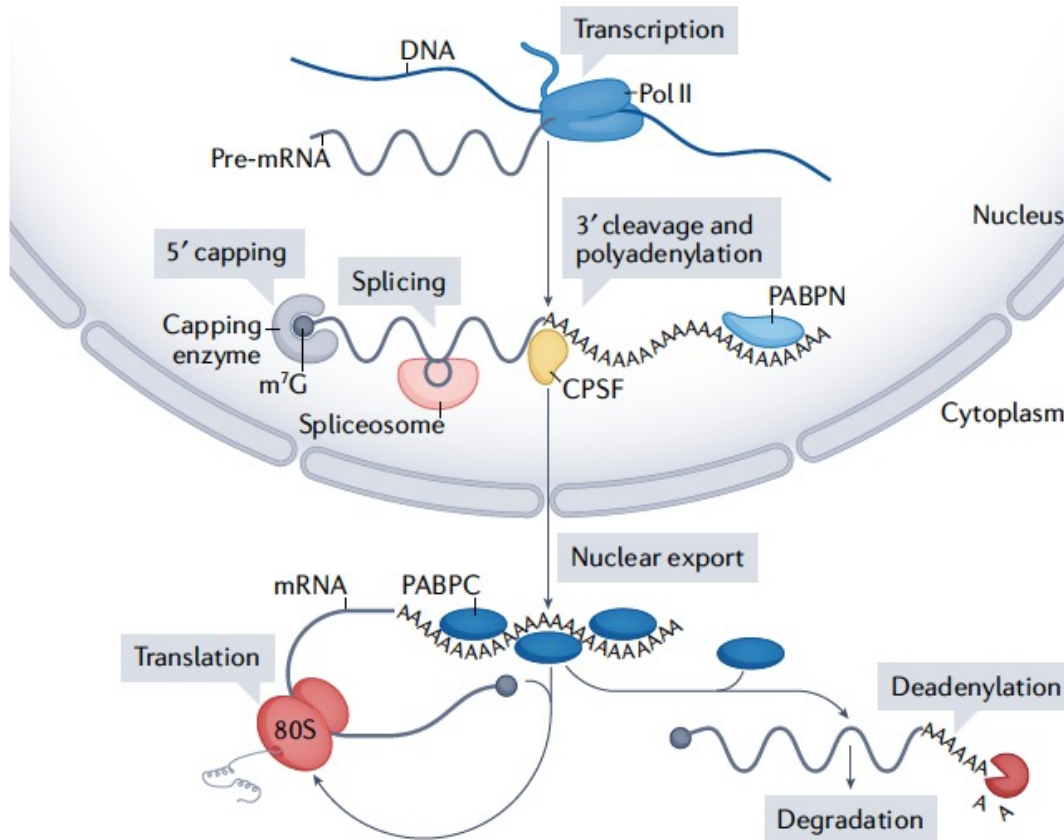


TRANSCRIPTION - RNAP
SPLICING - SPLICEOSOME
3'end FORMATION - CPA

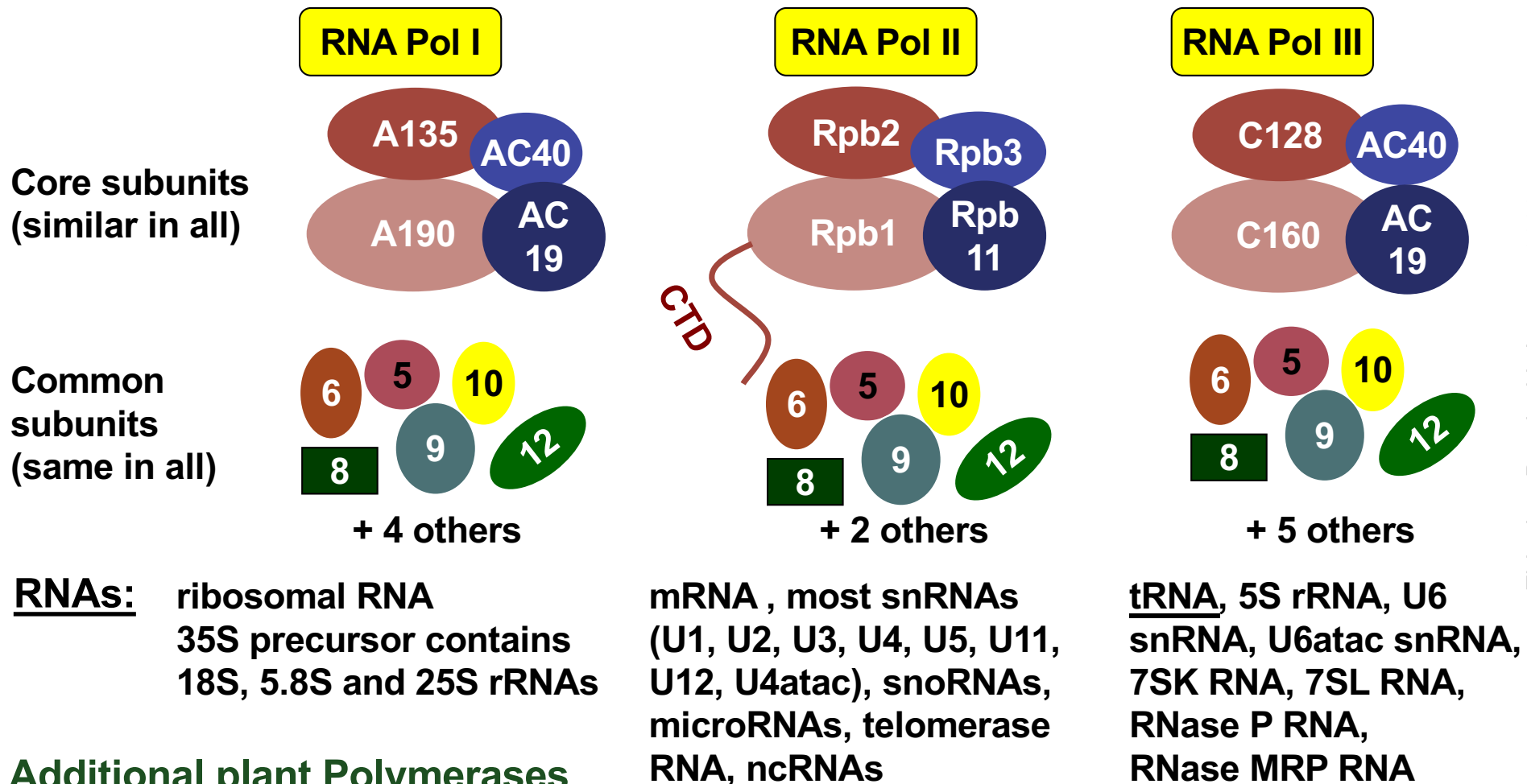


TRANSLATION - RIBOSOME
DEGRADATION

mRNA LIFECYCLE

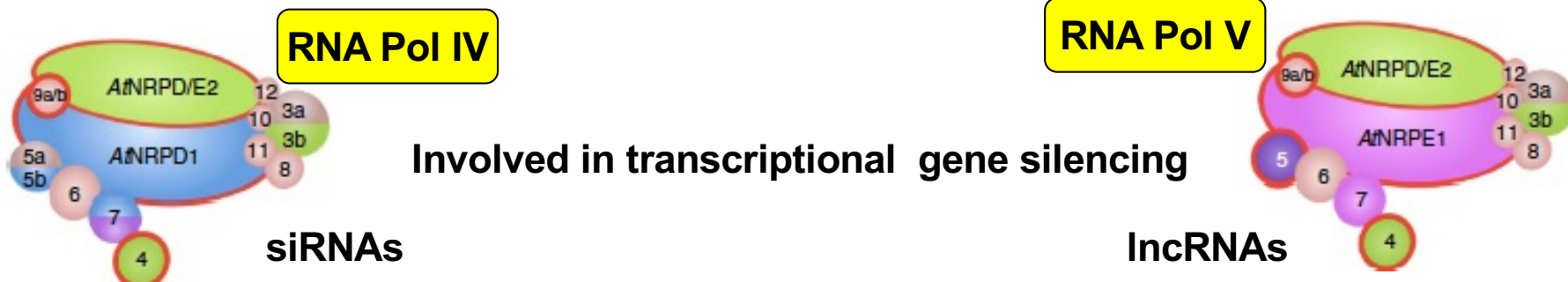


RNA POLYMERASES



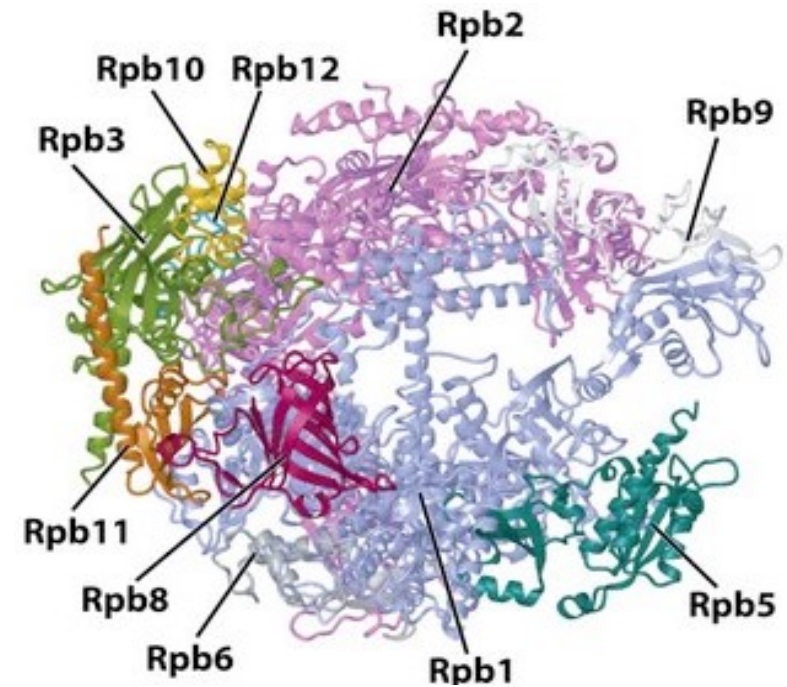
Zbigniew Dominski, lectures 2008

Additional plant Polymerases



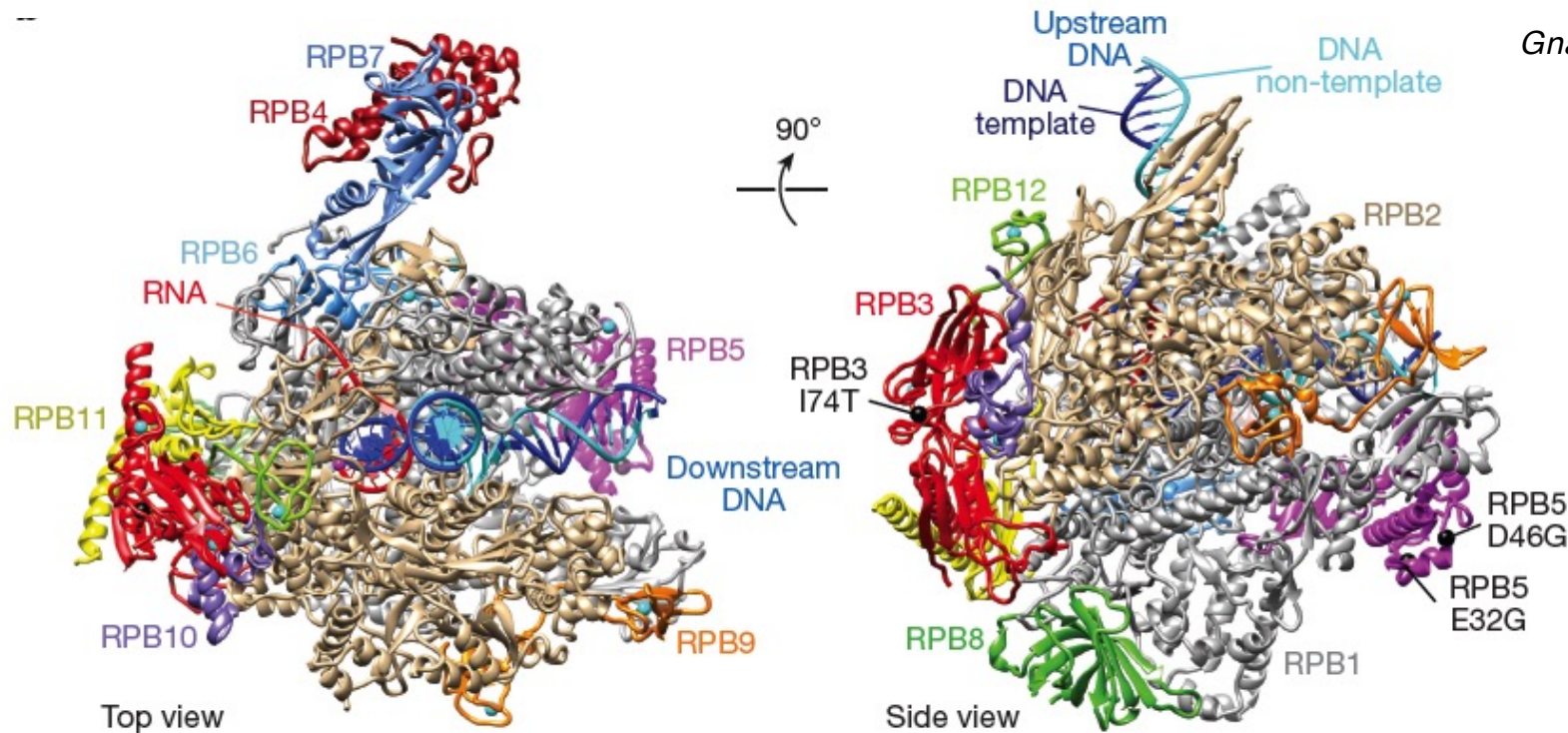
Yeast Pol II

- 12 subunits
- core by specific **Rpb1-3** and **11**
- **Rpb5-6, 8, 10** and **12** - shared by Pol I-III
- specific subcomplex **Rpb4/7** not essential
- associated factors RAP74, RAP30 (TFIIF)



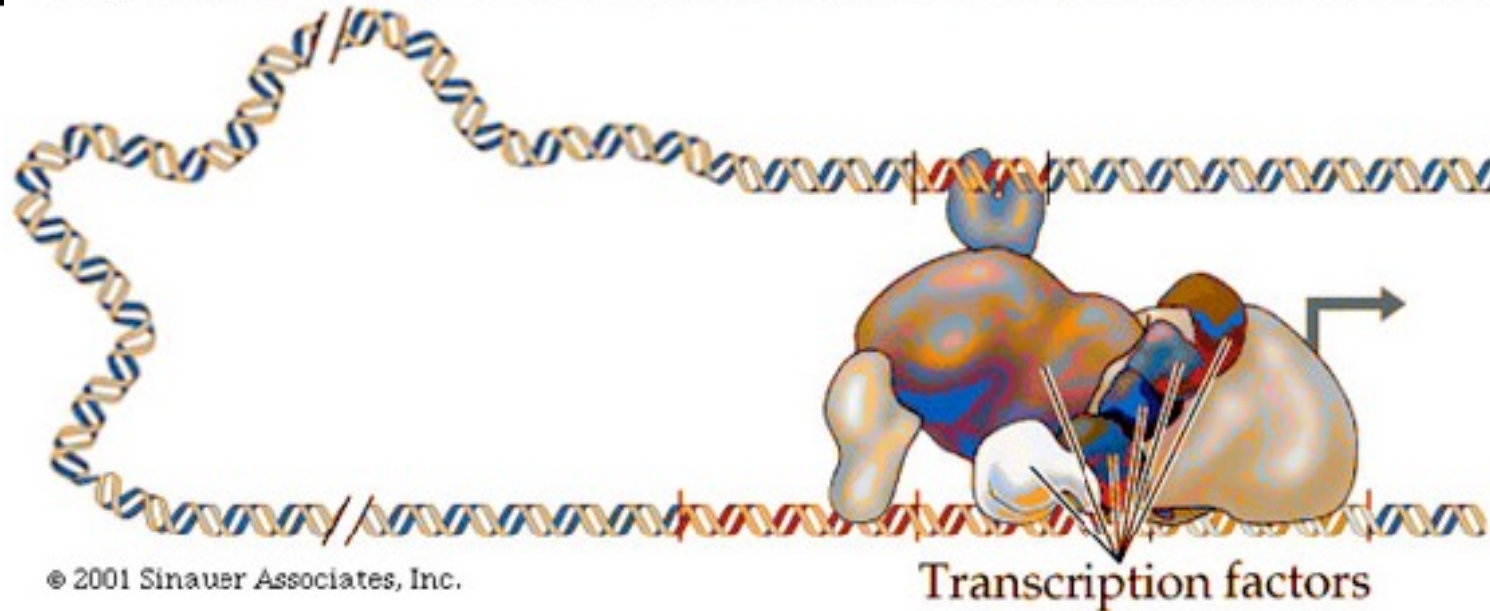
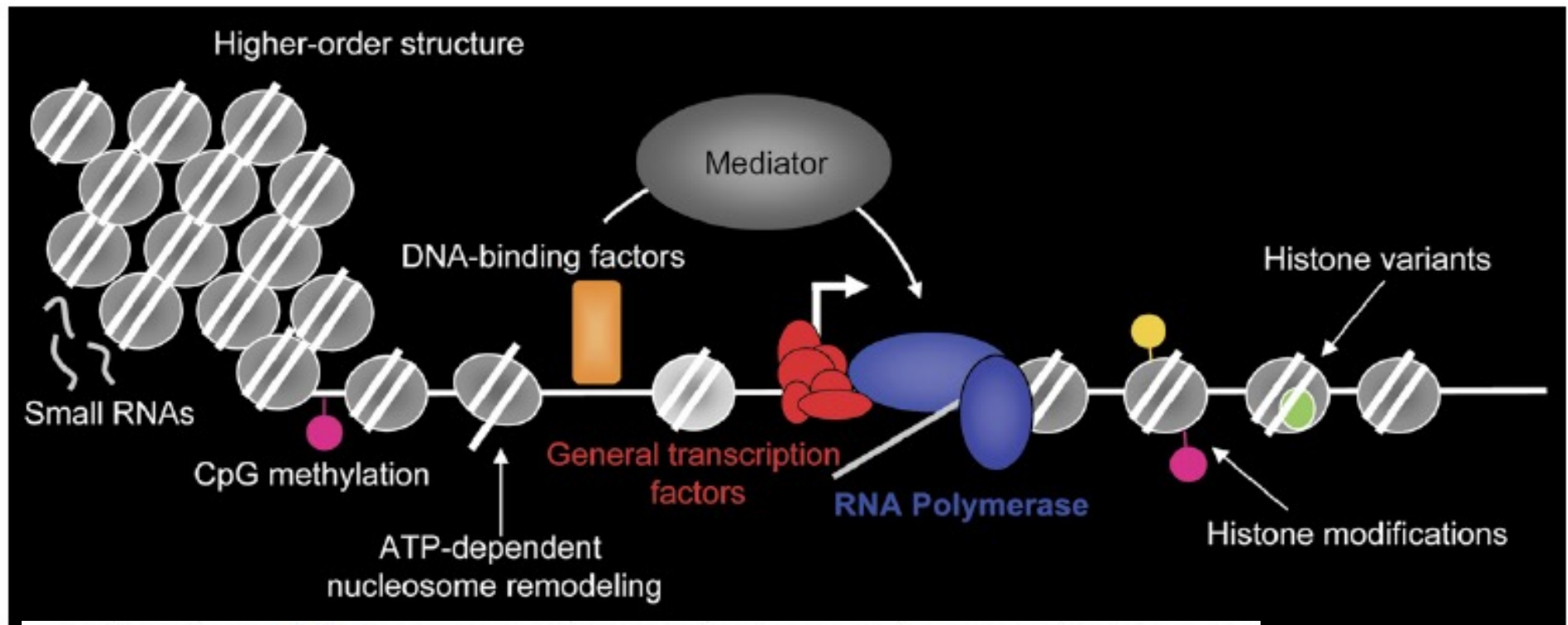
Gnatt et al, Science, 2001

Mammalian Pol II

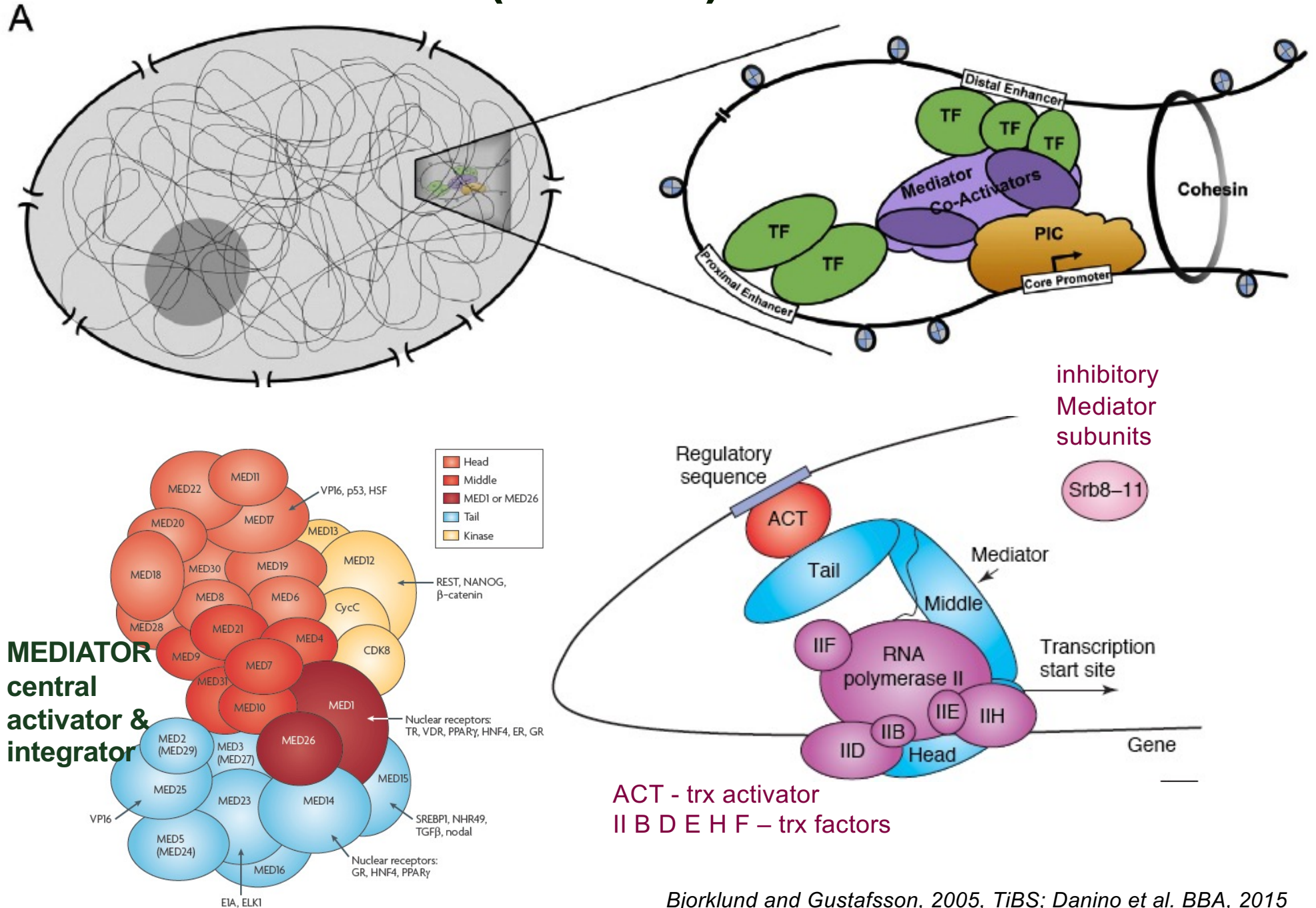


Bernecky et al, 2016, Nature

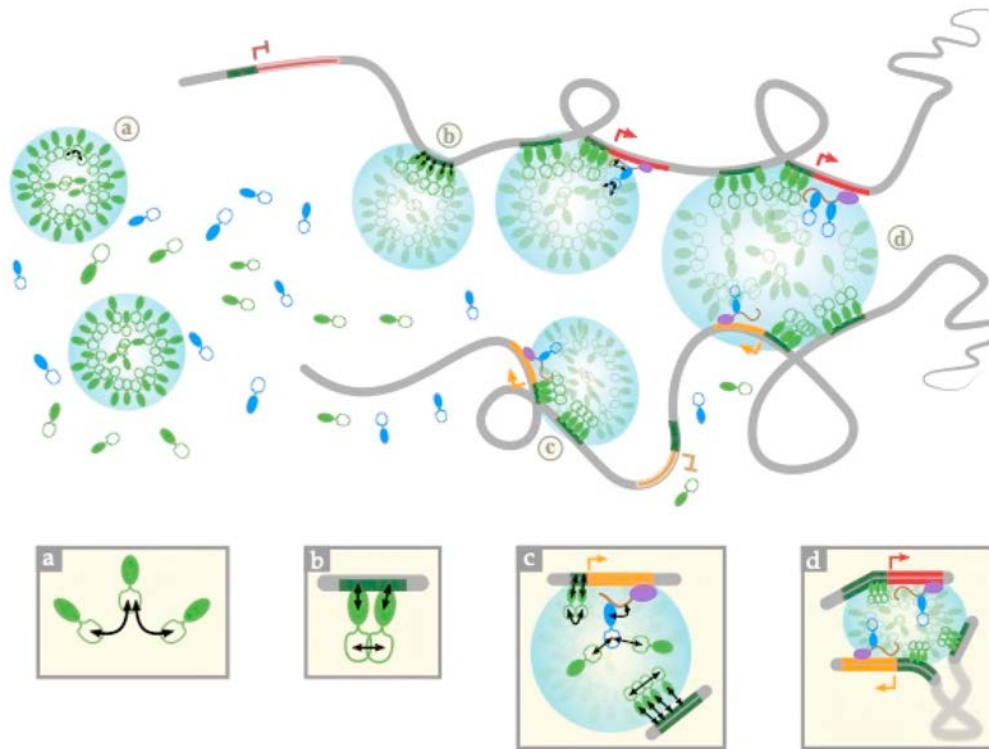
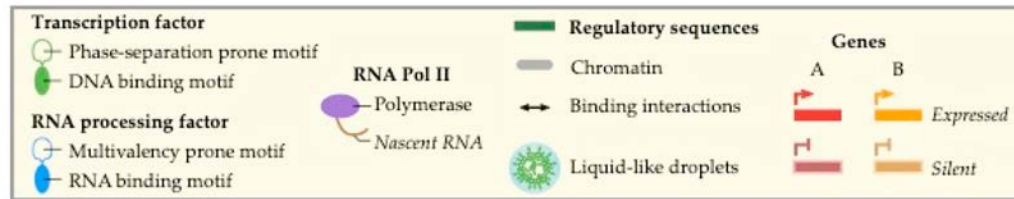
Pol II (RNAPII) in the cell



PoI II (RNAPII) in the cell

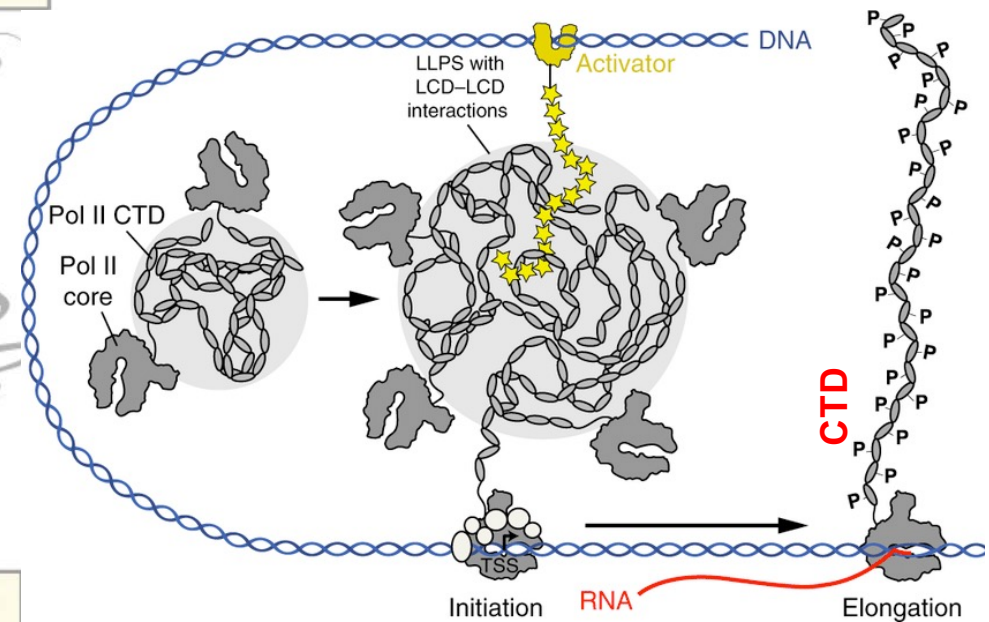


Pol II (RNAPII) in the cell



LLPS, droplets

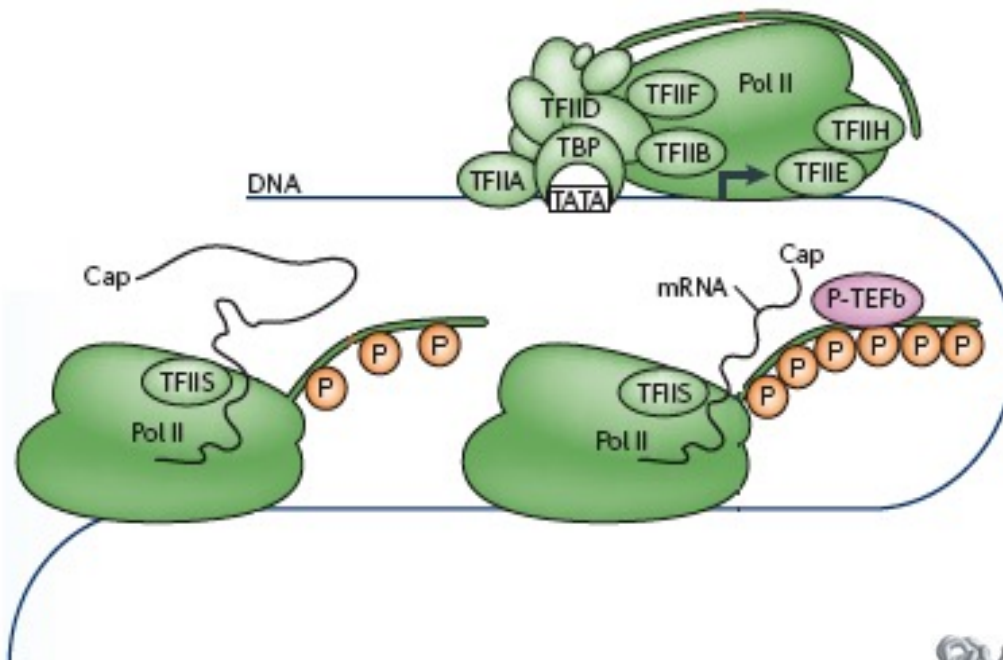
Liquid-liquid phase separation
Transcriptional condensates are formed by phase-separation self-assembly driven by IDR (Intrinsically Disordered Region)-containing proteins (e.g. CTD in Pol II)



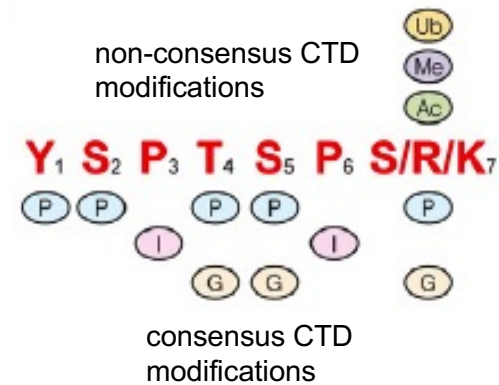
CTD-driven phase separation

Activators recruit/nucleate Pol II hubs near promoters. Initiation-coupled CTD phosphorylation removes individual Pol II enzymes for transcription elongation.

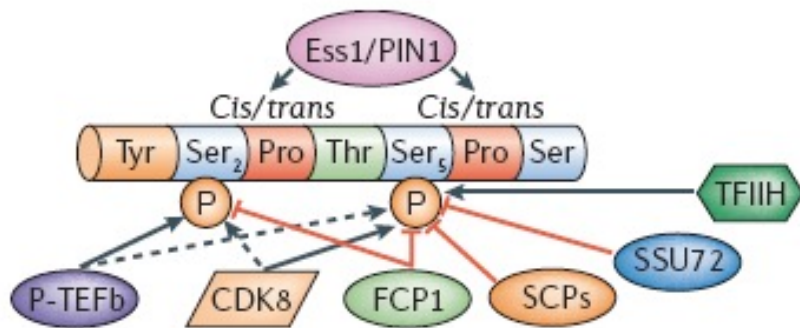
Pol II C-terminal domain (CTD)



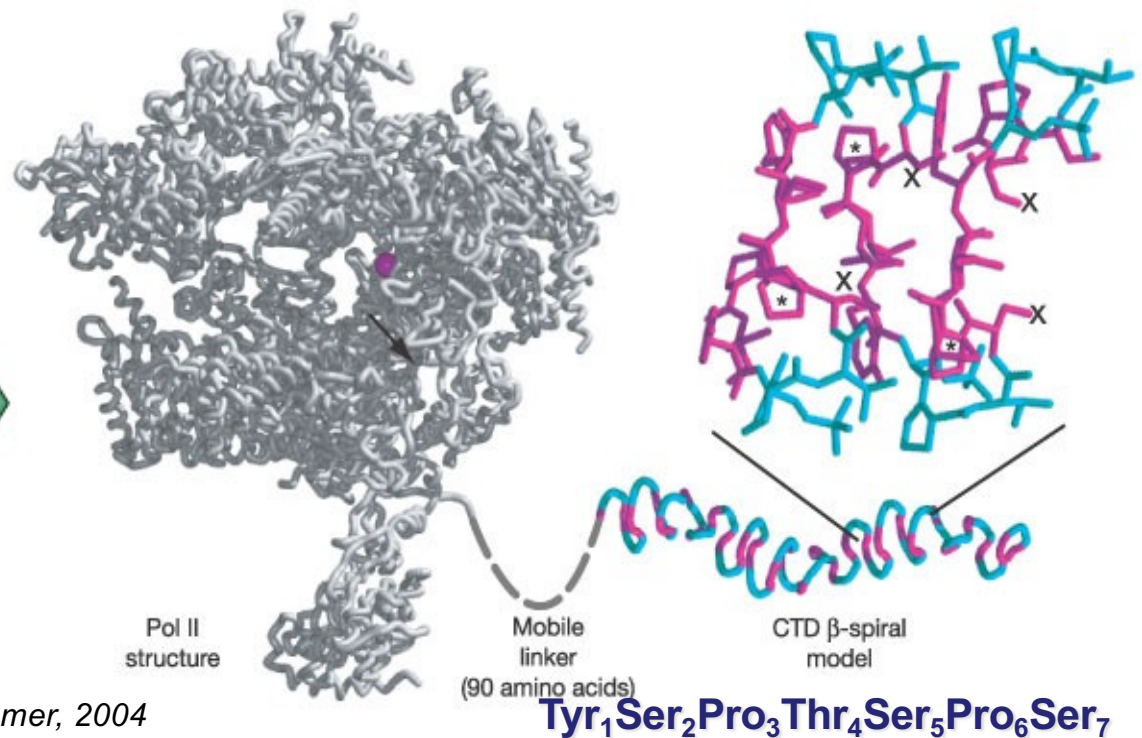
Goodrich and Kugel, Nat. Rev. Mol. Biol., 2006



26 (yeast) - 52 (human) repeats



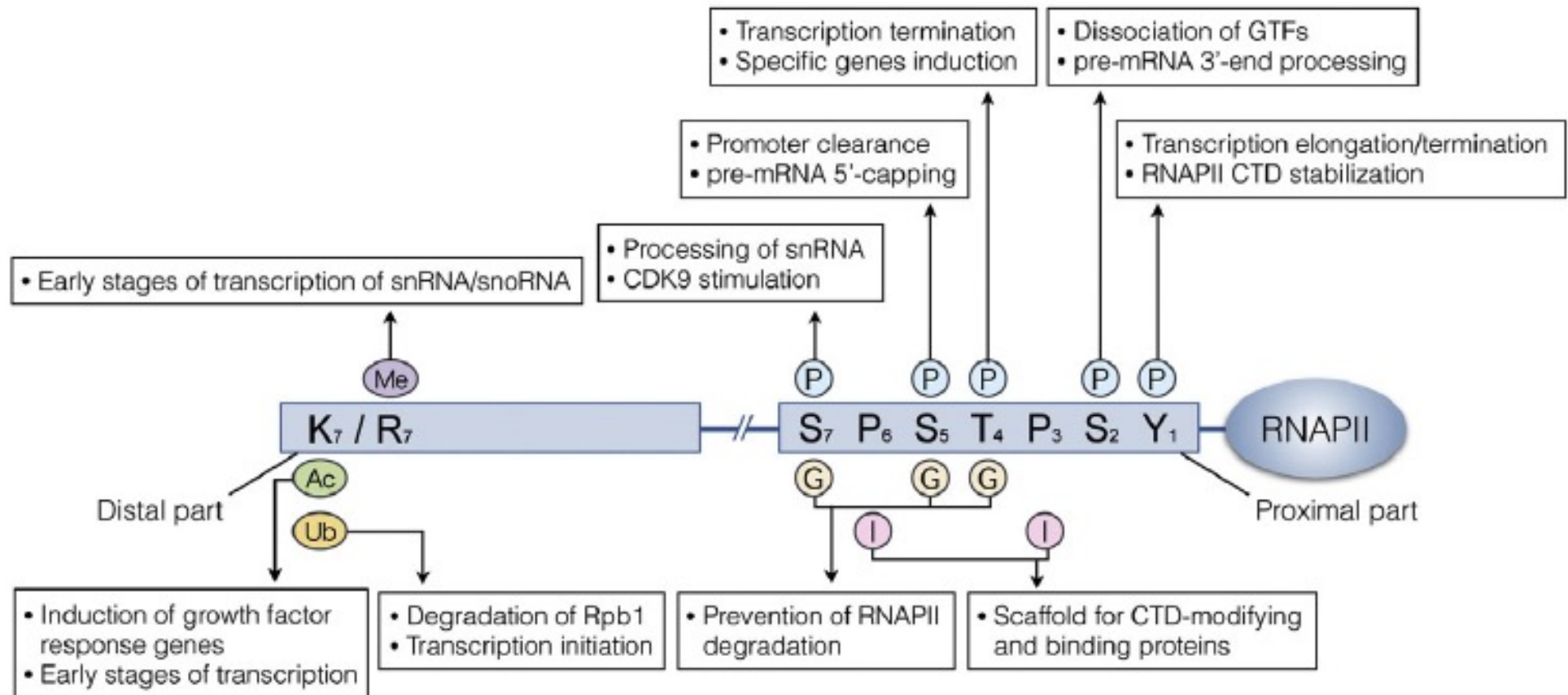
Saunders et al, 2006, Nat.Rev.Mol.Cel.Biol



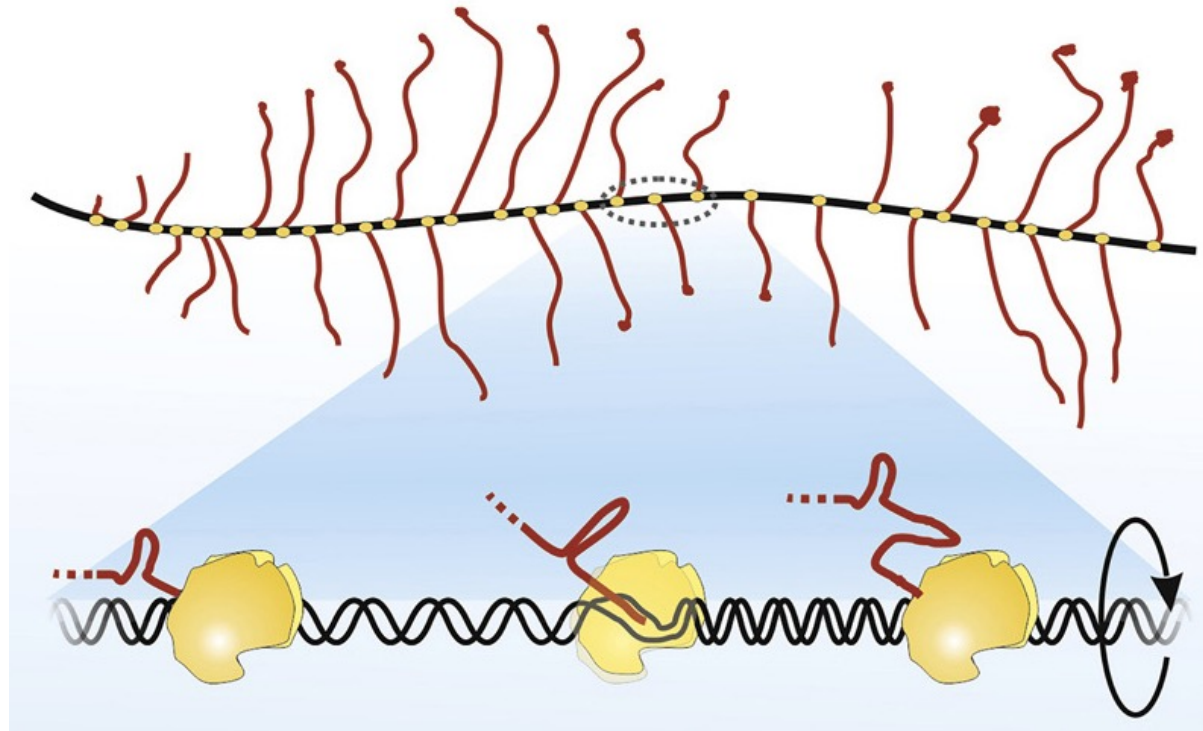
Meinhart and Cramer, 2004

Tyr₁Ser₂Pro₃Thr₄Ser₅Pro₆Ser₇

CTD CODE



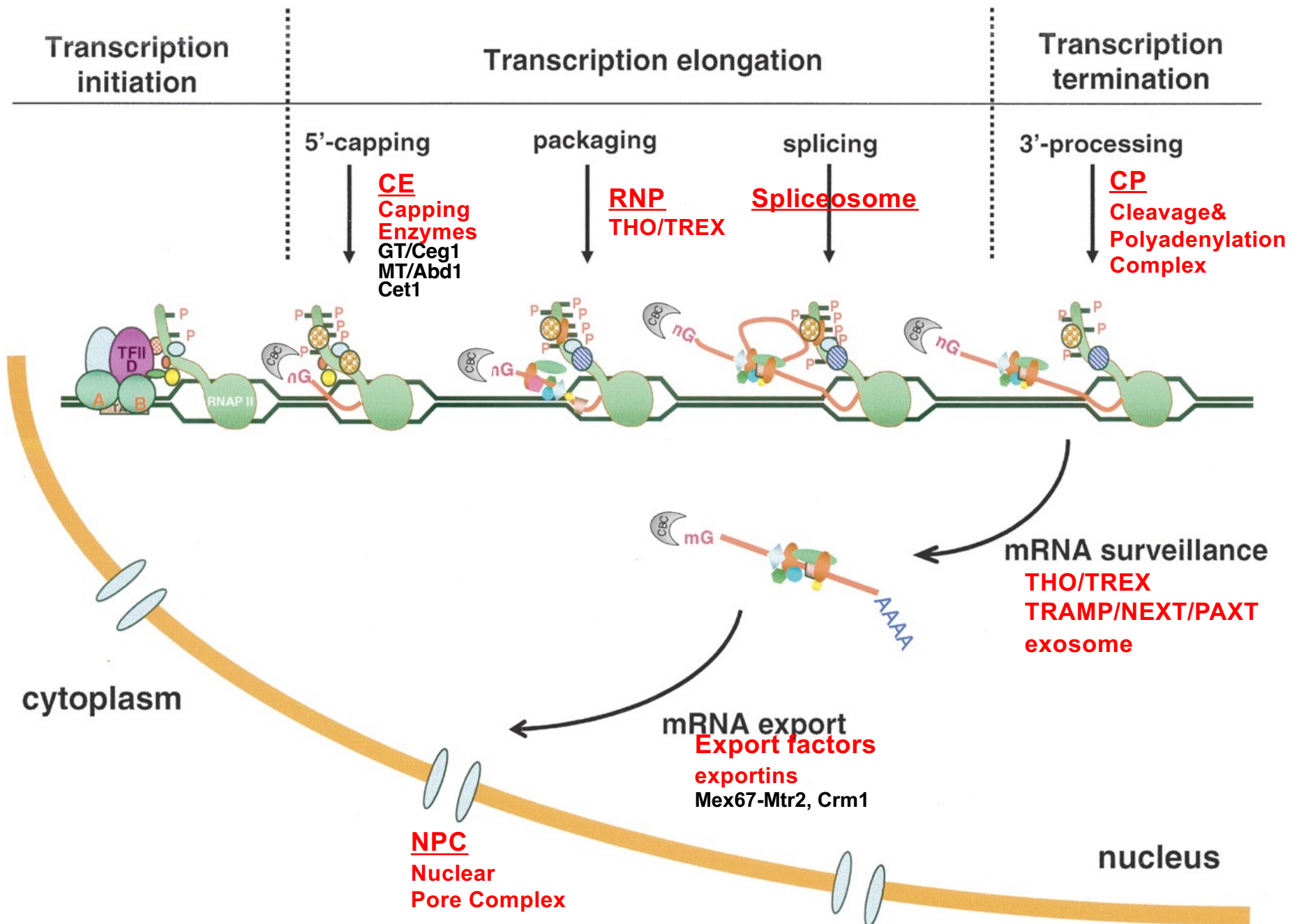
NASCENT TRANSCRIPTS



Nascent transcript = during formation, newly formed, still bound by polymerase

- nascent RNAs couple RNA processing with transcription elongation and chromatin modification
- nascent RNAs modulate binding of proteins to regulatory elements (chromatin)
- regulatory effects of nascent transcripts can be enhanced by gene looping
- high concentrations of nascent RNAs can initiate formation of nuclear bodies
- sometimes the function is conferred by nascent transcription (activity) and not the transcript itself

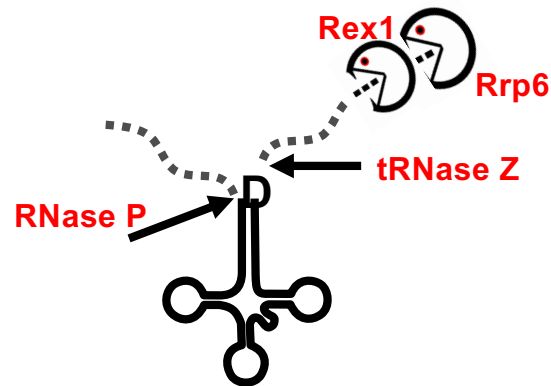
CO-TRANSCRIPTIONAL PROCESSES



POST-TRANSCRIPTIONAL PROCESSES

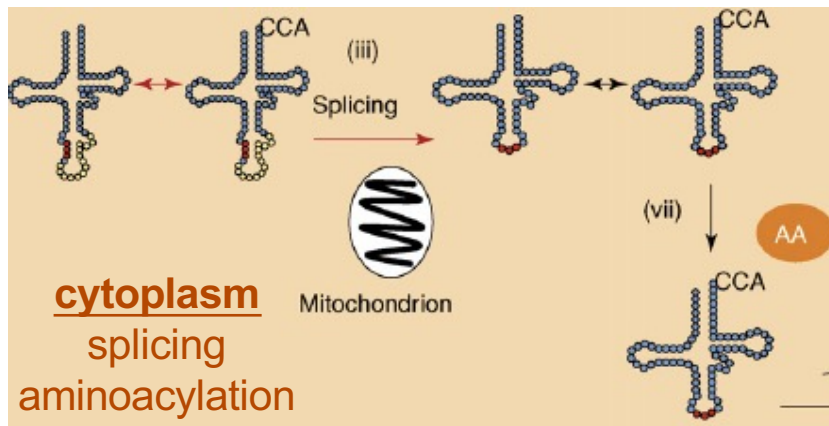
tRNA PROCESSING

- 5' end by RNase P
- 3' end by tRNase Z or
- by exonuclease Rex1 and Rrp6



tRNA SPLICING

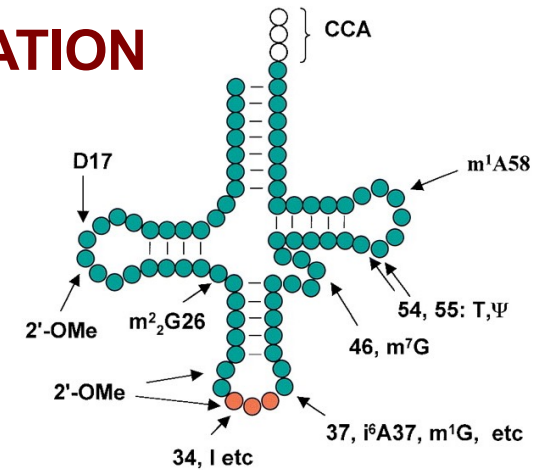
In the cytoplasm on the mitochondrial membrane (YEAST!!)



Hopper and Shaheen, TiBS, 2008

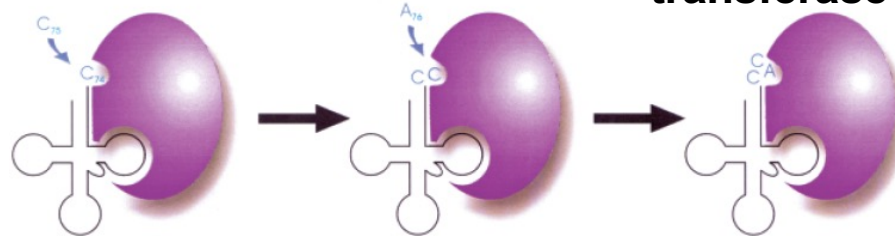
tRNA MODIFICATION

by RNA modifying enzymes

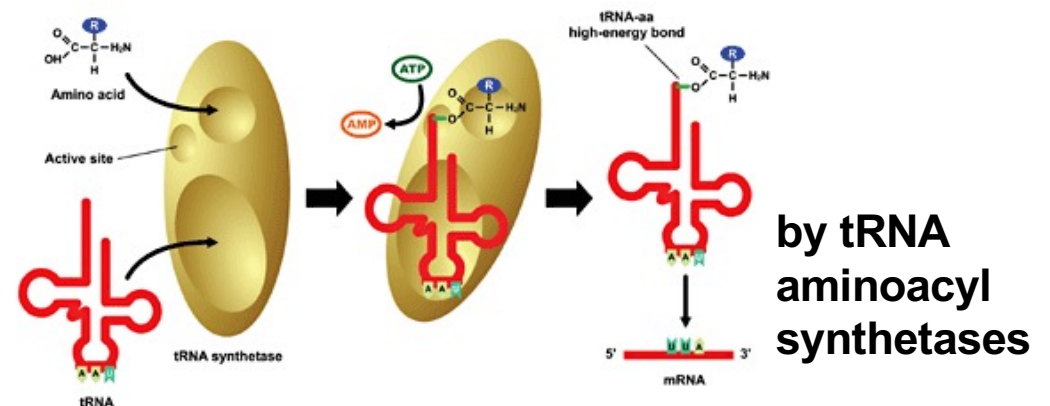


tRNA CCA ADDITION

by tRNA nucleotidyl-transferase



tRNA AMINOACYLATION



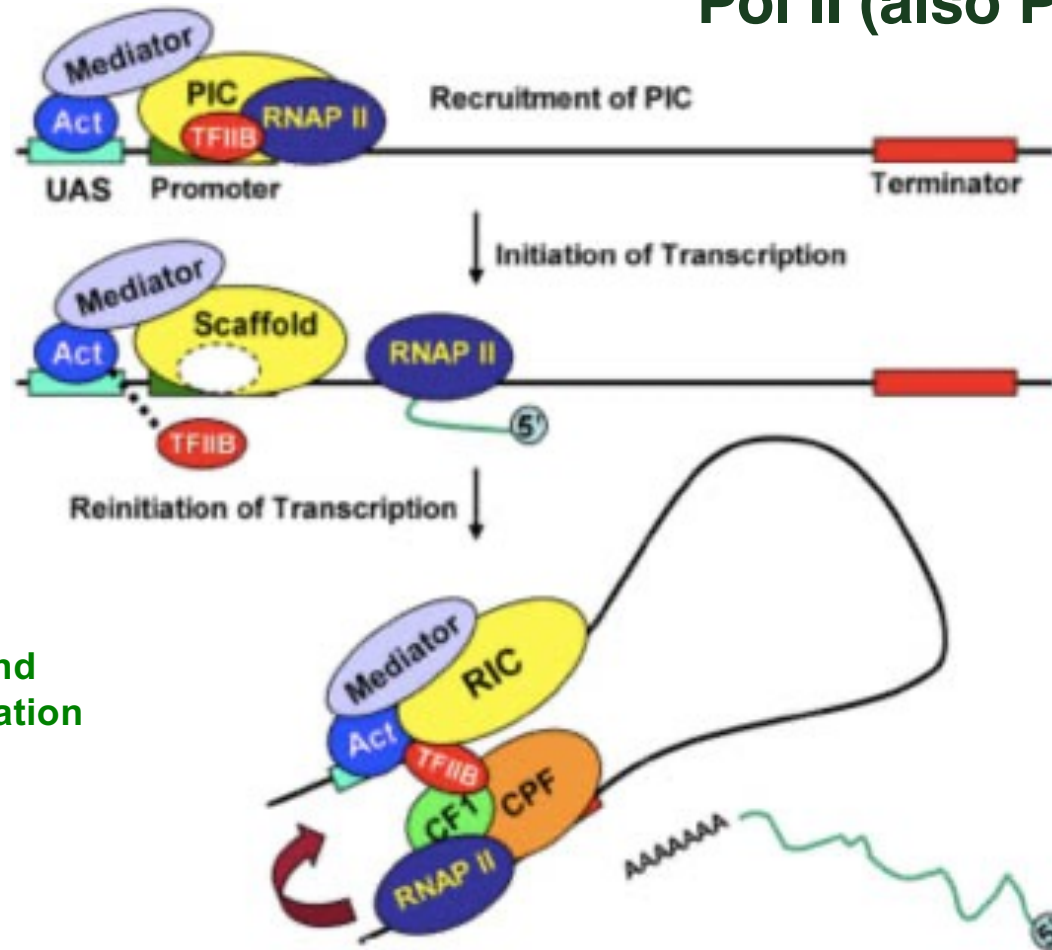
GENE LOOPING

Pol II (also Pol I)

PIC
Preinitiation
Complex

Scaffold
transcription
factors
(TFIID, A, E, H)

CF1, CPF
Cleavage and
Polyadenylation
Complex

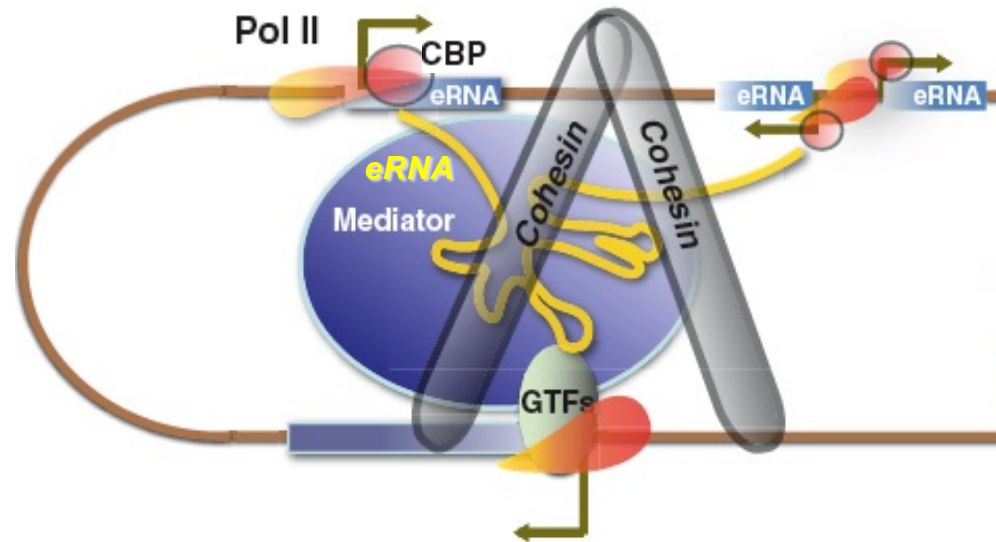
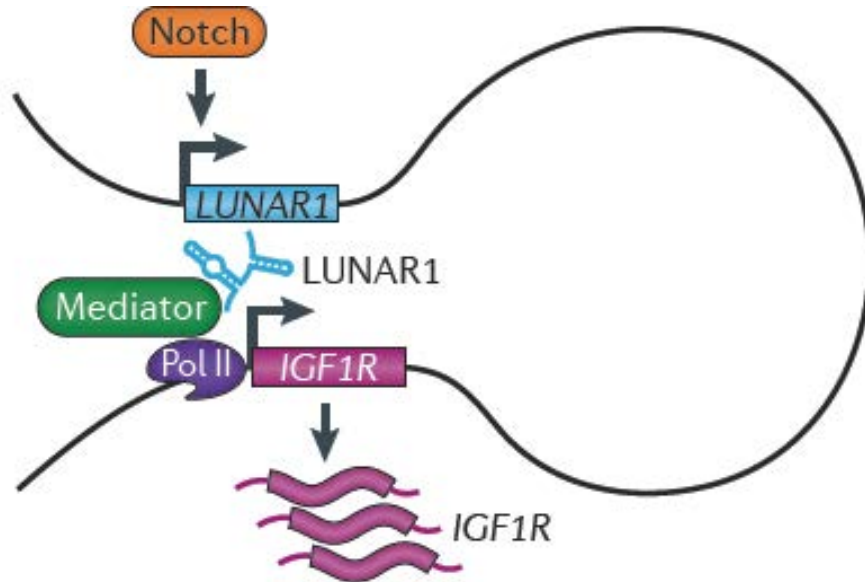


Loop formation requires interaction between factors at the promoter (TFIIB) and terminator (Rna15 from CF1) /in mammals: transcription factors, nuclear receptors, insulators, chromatin remodellers, Polycomb, architectural proteins/

Loop function: facilitation of transcription reinitiation of PolII, but also repression of gene expression (PcG, DNA methylation)

GENE LOOPING

via Mediator and enhancer RNAs (eRNAs)

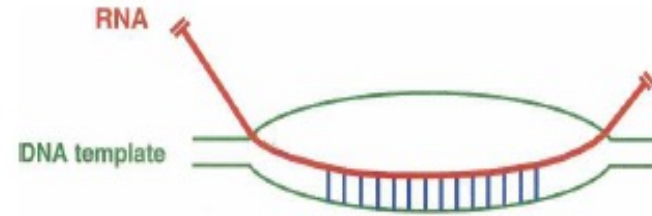


Some eRNAs (e.g. *LUNAR1* near the IGF1R locus) mediate chromosome looping between enhancers and nearby genes via Mediator or MLL protein complexes

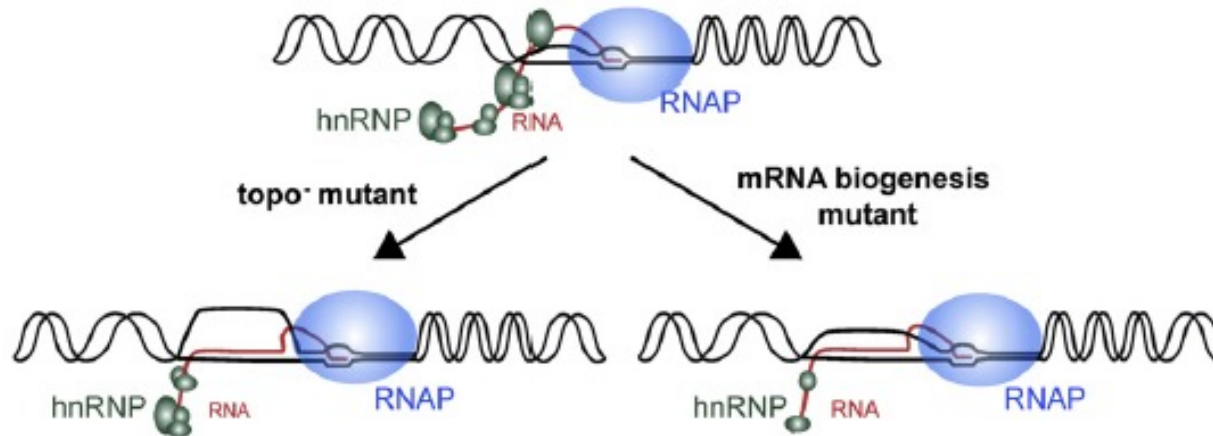
*Quinn and Chang, Nat Rev Genet 2015;
Lai and Shiekhatair, Curr Op Gene Dev 2014*

R-LOOPS

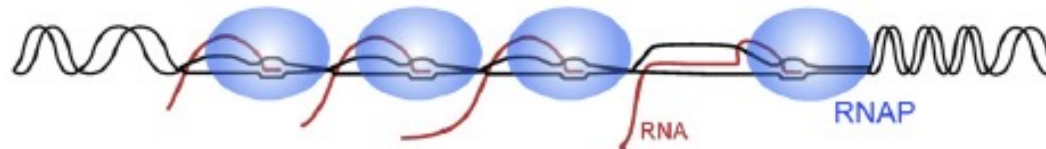
DNA::RNA hybrids formed during transcription before RNP packaging



A Transcription associated R-loop formation



B RNAP roadblock



R-loops

- accumulate in RNP biogenesis mutants (*tho*, *sen1*, mRNA export)
- negative effects: polymerase stalling, termination defects, replication fork stalling, DNA damage, genetic instability
- prevented by topoisomerases, helicase Sen1, THO complex, resolution (cleavage) by RNase H

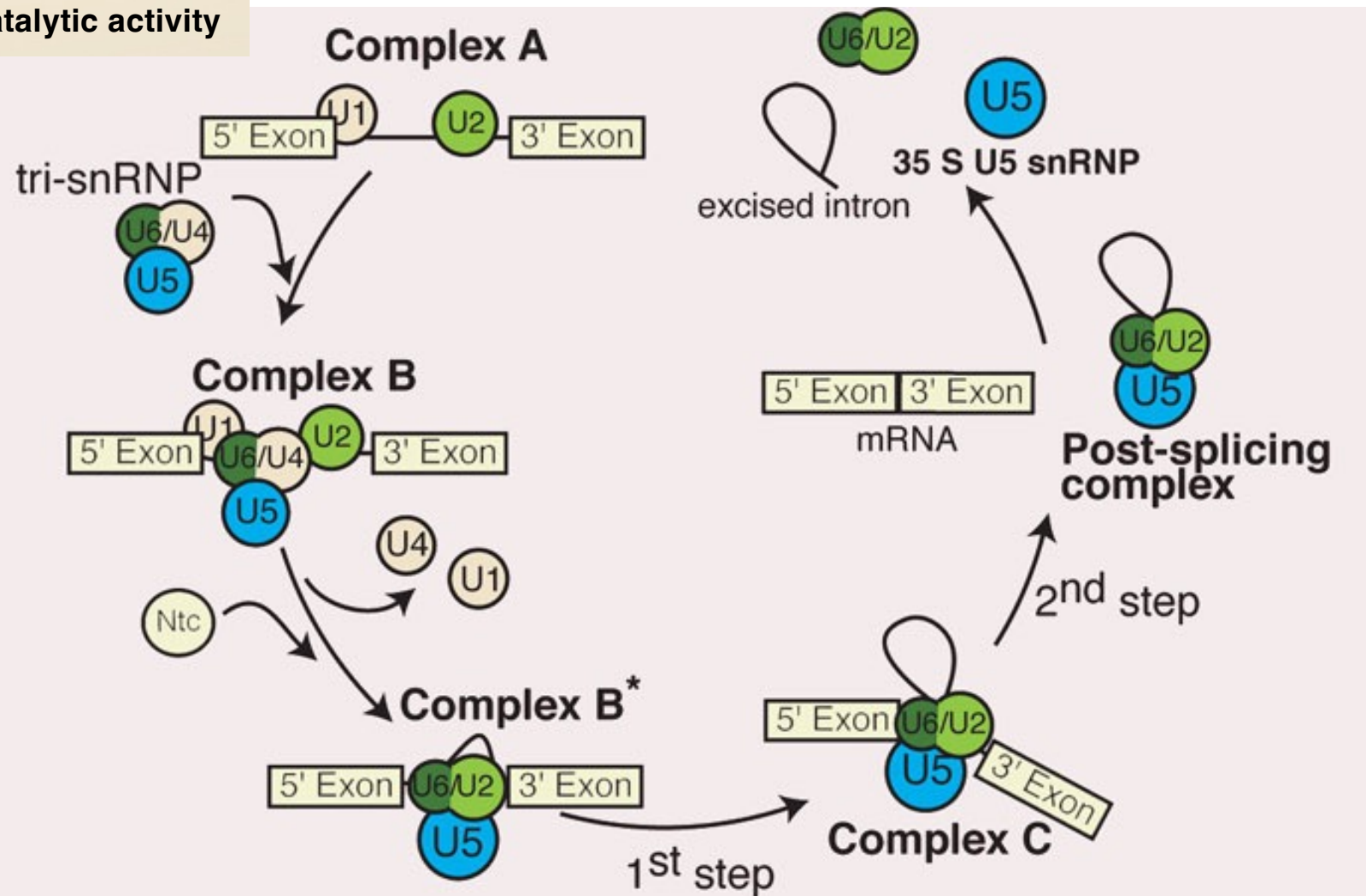
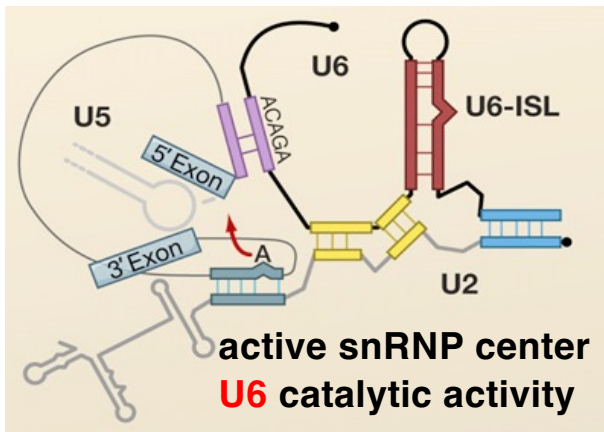
SPLICEOSOME

5 snRNAs: U1, U2, U4, U5, U6

Core Sm or LSM (U6) proteins 1.7 - 3 MDa

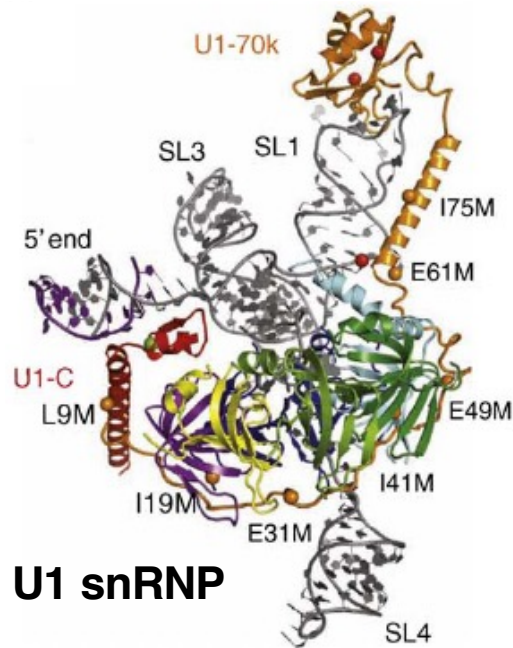
Specific snRNP proteins

Splicing factors



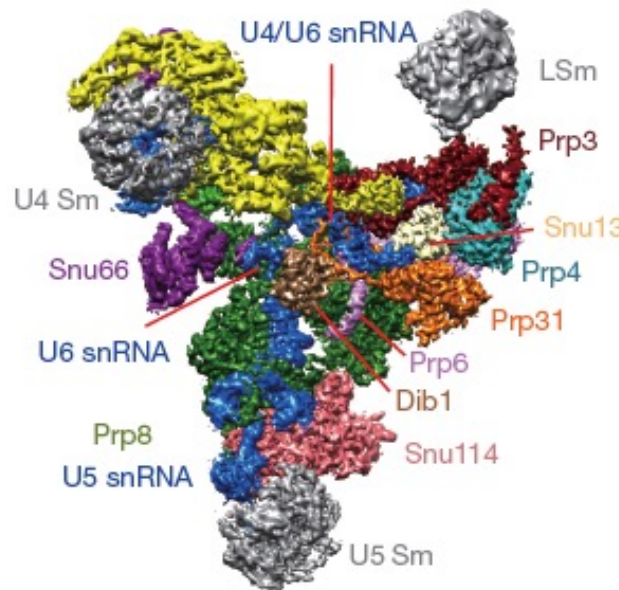
SPLICEOSOME

Cryo- EM



U1 snRNP

Krummel et al, Nature, 2009

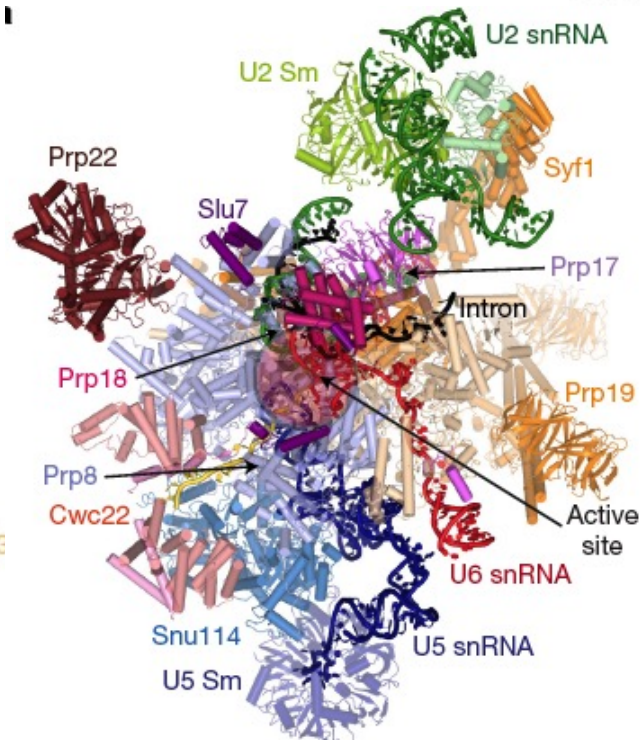


U4/U6.U5 tri-snRNP

Nguyen1, Galej et al, Nature, 2016*

C complex yeast

Galej et al, Nature, 2016



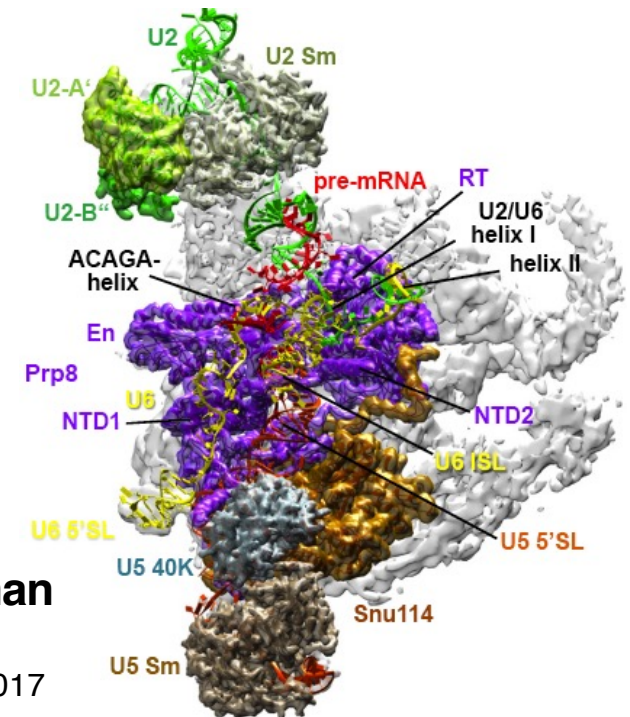
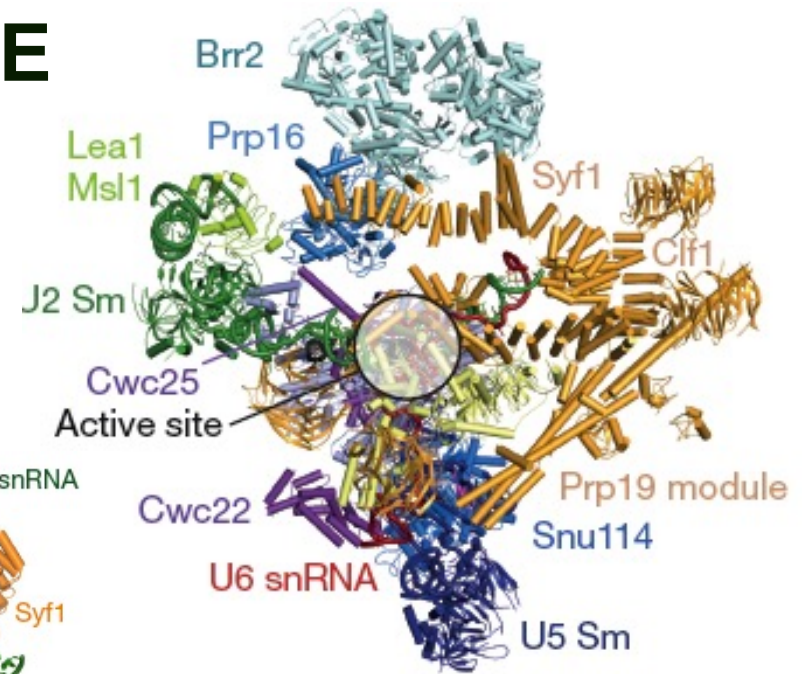
C* complex yeast

Fica et al, Nature, 2017

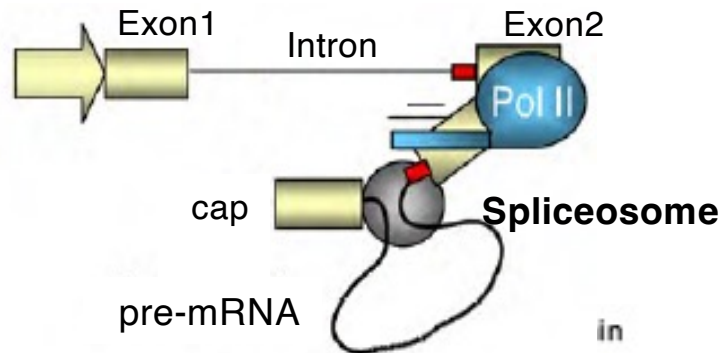
C* complex human

second step

Bertram et al, Nature, 2017

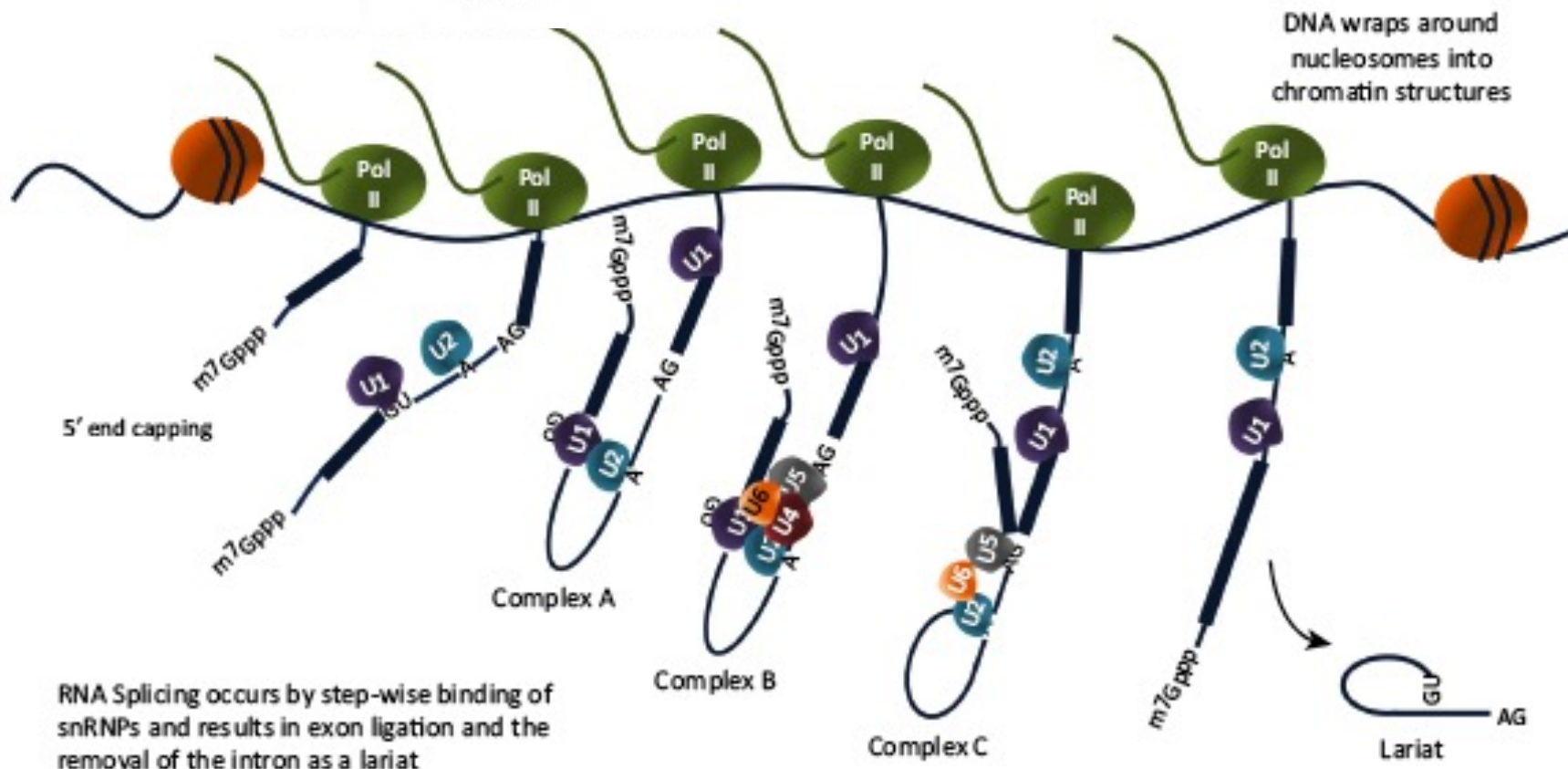


SPLICING: co-transcriptional process



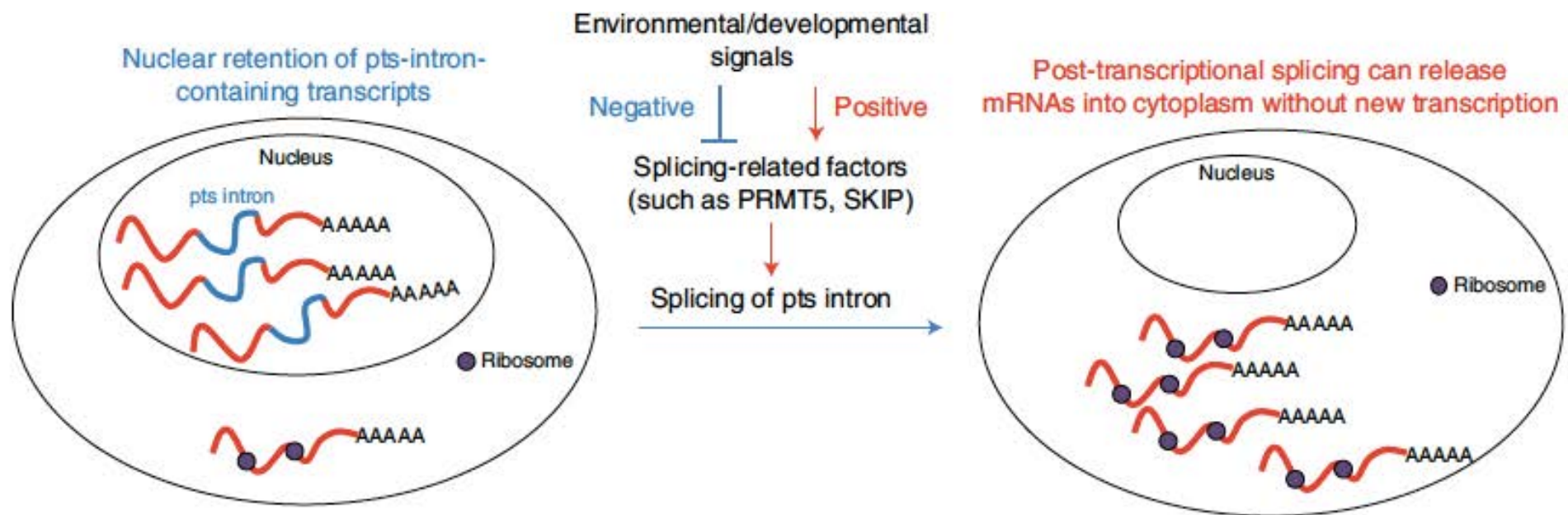
- spliceosome assembly (**Ser5-P**)
- majority of splicing (up to 70-80%)

Munoz et al., TiBS, 2009



Wong et al., TiG, 2014

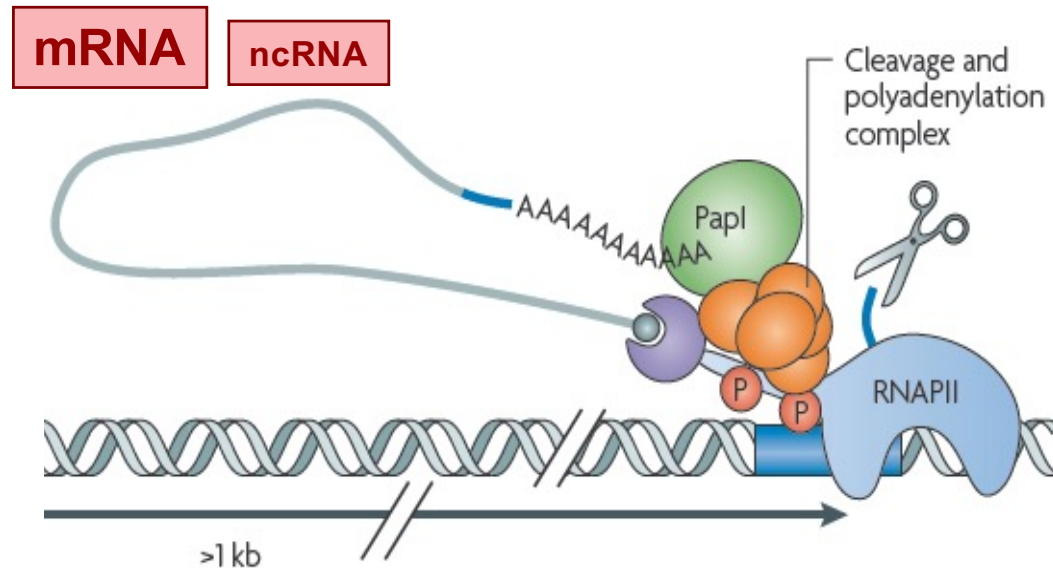
Co-trx vs post-trx splicing



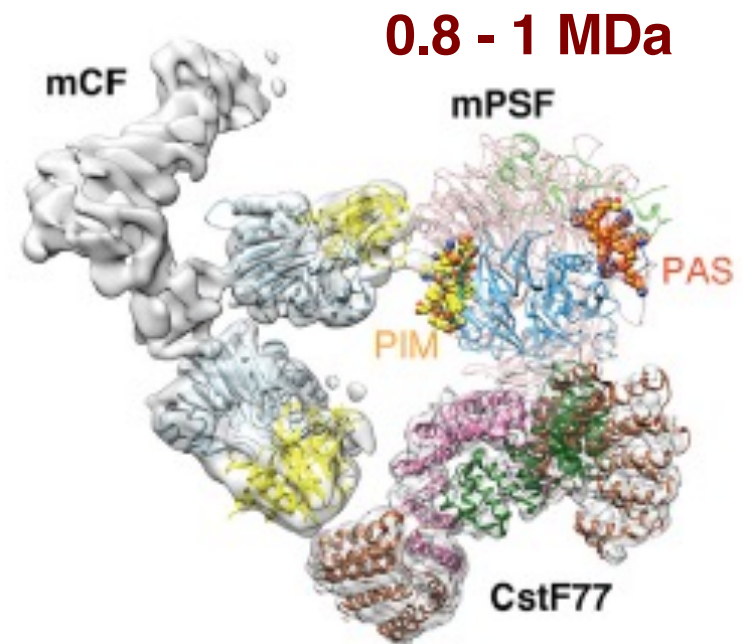
Nanopore-based profiling of chromatin-bound RNA

- Incompletely spliced and polyadenylated transcripts are detected on chromatin
- They are not released and exported to the cytoplasm and undergo post-transcriptional splicing
- Splicing of these introns is regulated in response to various environmental signals
- It represents additional layer of stress-related gene expression reprogramming
- Alternative introns are less efficiently spliced than constitutive introns
- Alternative introns are more often removed post-transcriptionally

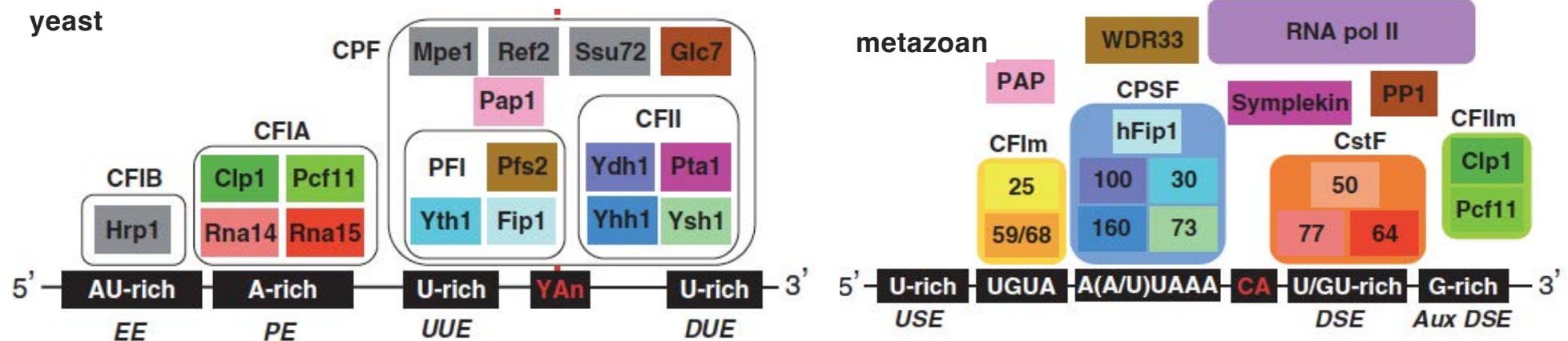
CPA Cleavage and Polyadenylation



Jacquier, Nat. Rev. Genet, 2009



Zhang et al, Mol Cell, 2019

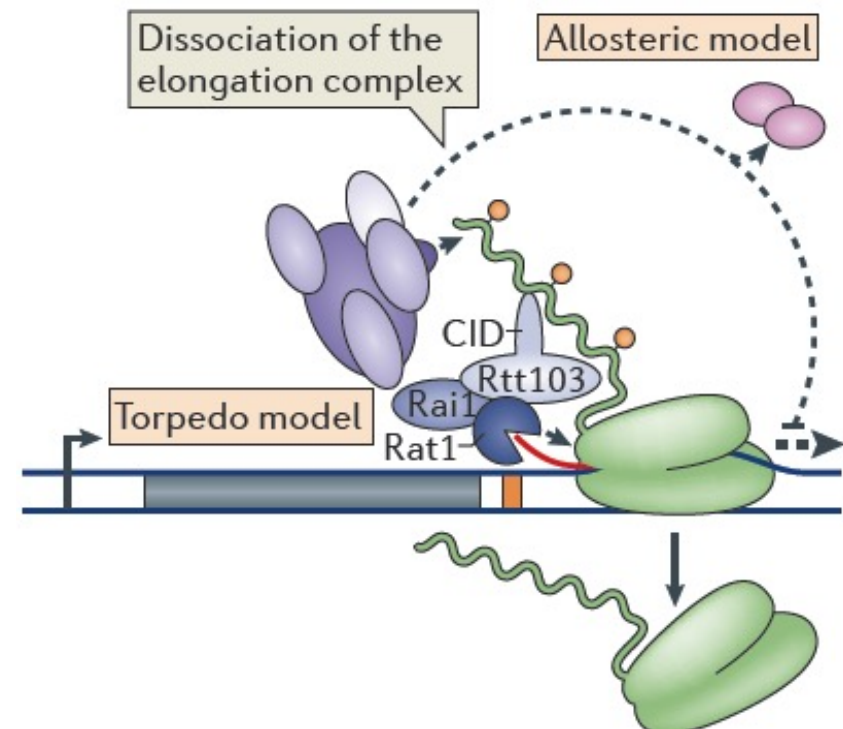
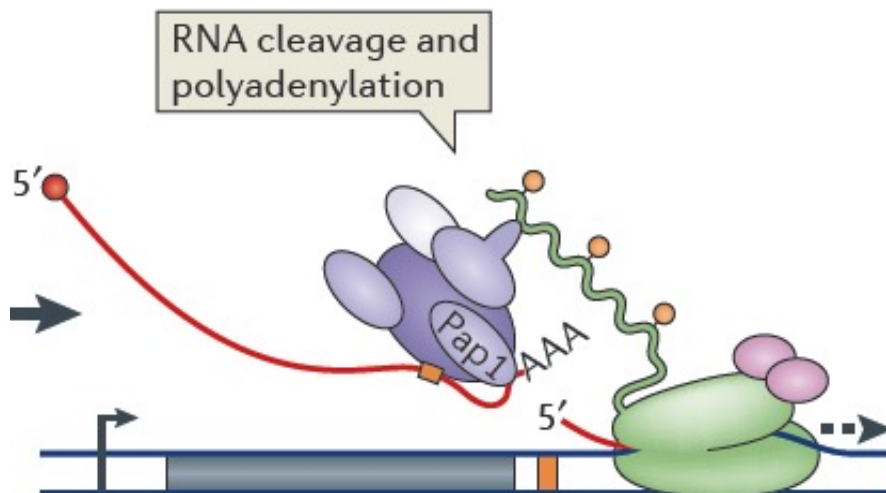
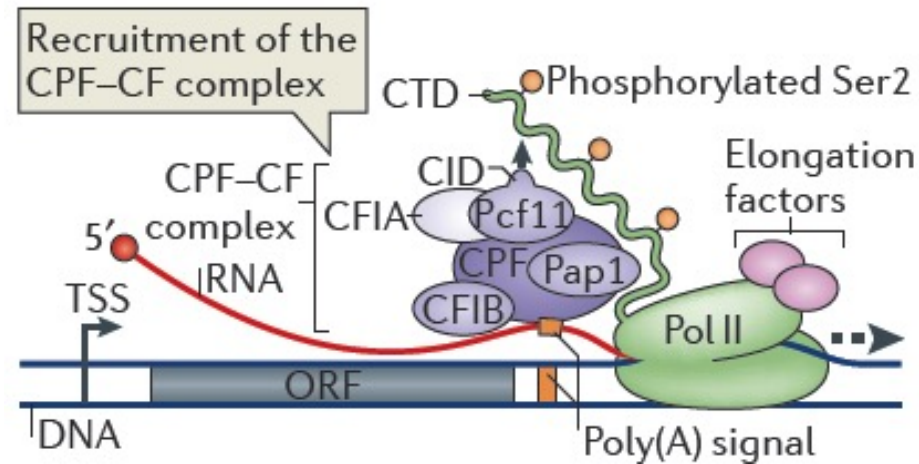


Cleavage by CPSF-73 (human), Brr5/Ysh1 (yeast)

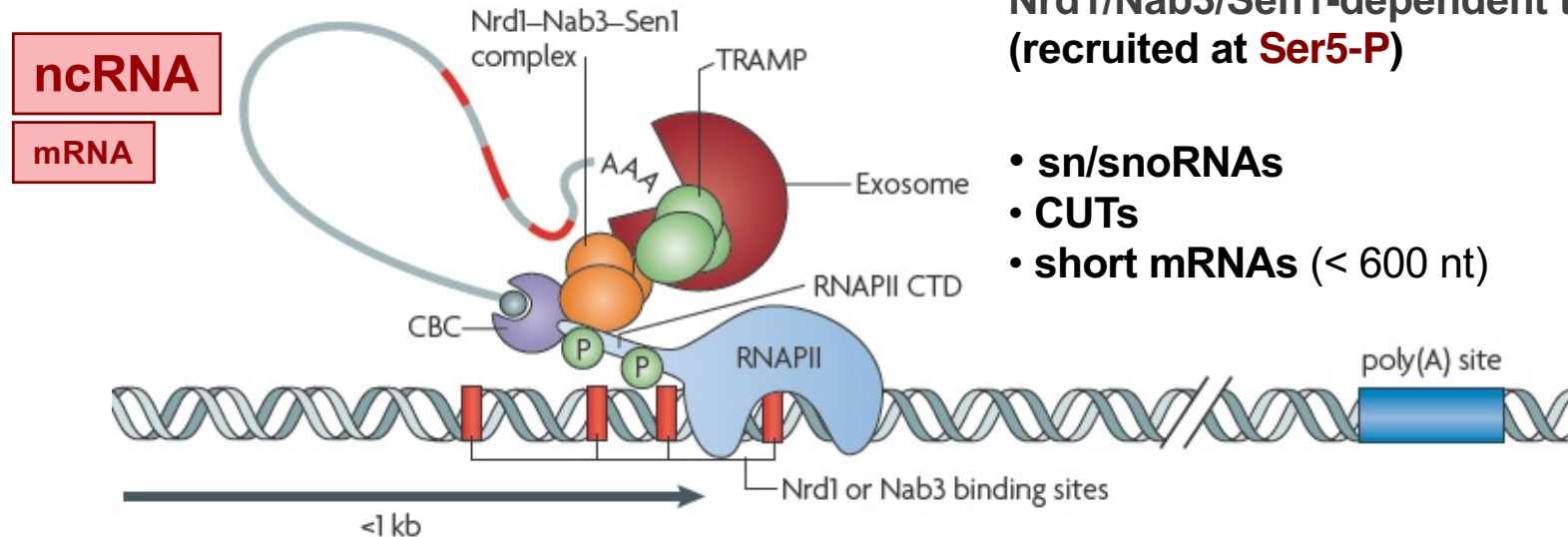
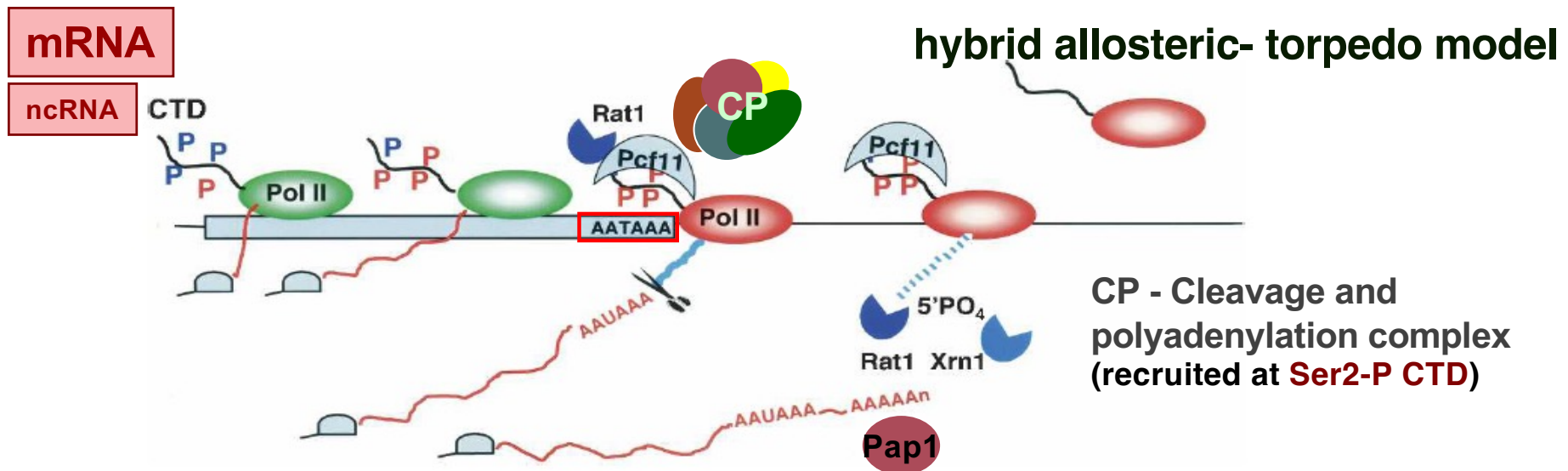
Millevoi and Vagner, NAR, 2008

CPA: mRNA 3' end formation

transcription termination at mRNA genes



POL II TRANSCRIPTION TERMINATION



Lecture on transcription termination by Michał Koper

RIBOSOME

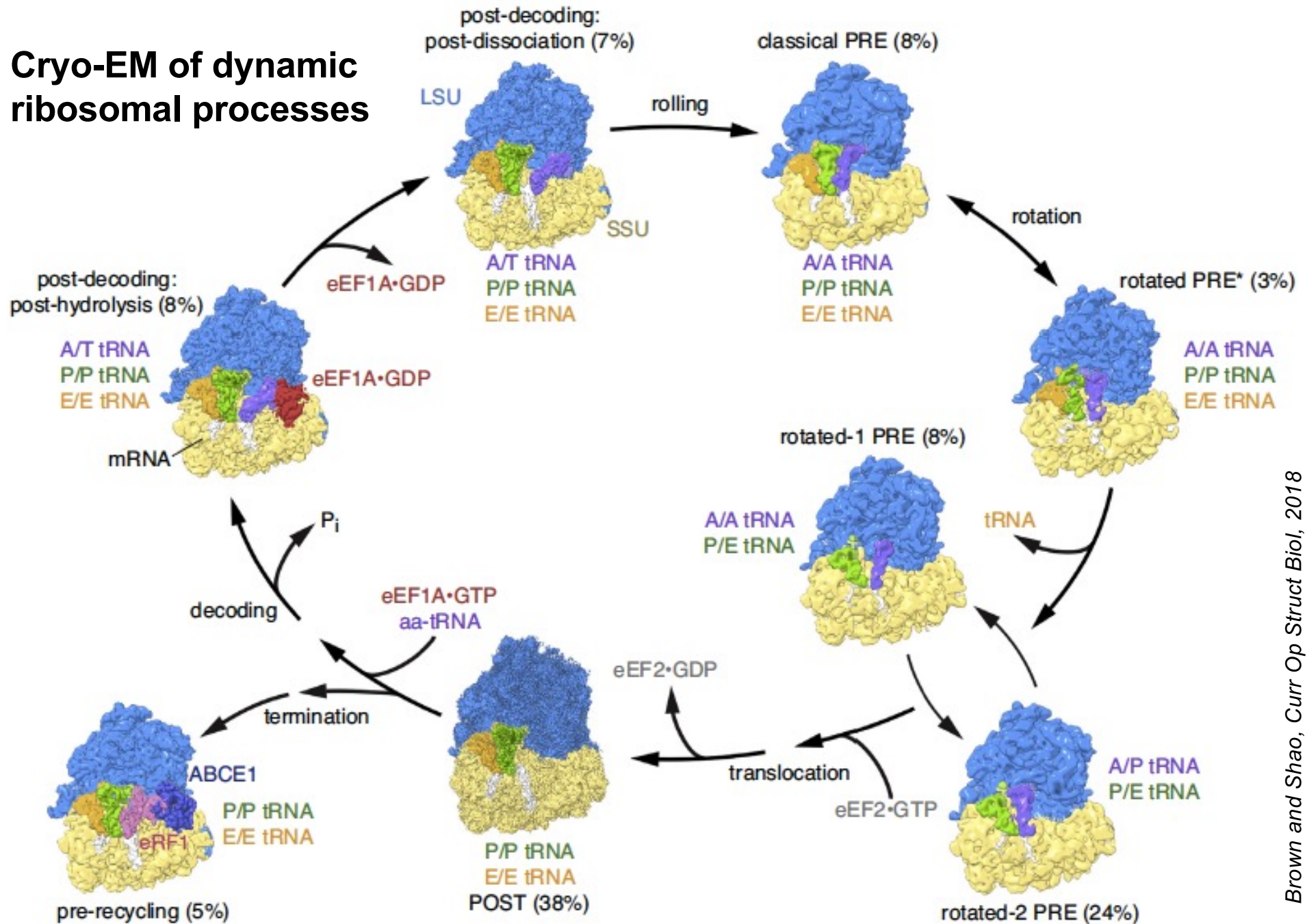
3.3 MDa (yeast) – 4.3 MDa (humans)



- No ribosomal protein with a peptidyl transferase (PT) activity
- Drugs (chloramphenicol) that inhibit PT bind to the 23S rRNA (PT loop)
- Mutations that provide resistance to these drugs map to the PT loop
- Nearly all (99%) of proteins can be stripped from the large subunit and it still retains the PT activity
- Only RNA chains are close enough to the PT center (structure)
- Ribosomal proteins are important for ribosome stability and integrity, but NOT for catalysis

TRANSLATION CYCLE

Cryo-EM of dynamic ribosomal processes



Next lecture

RNA enzymes and complexes

RNA granules and subcellular structures

RNA decay