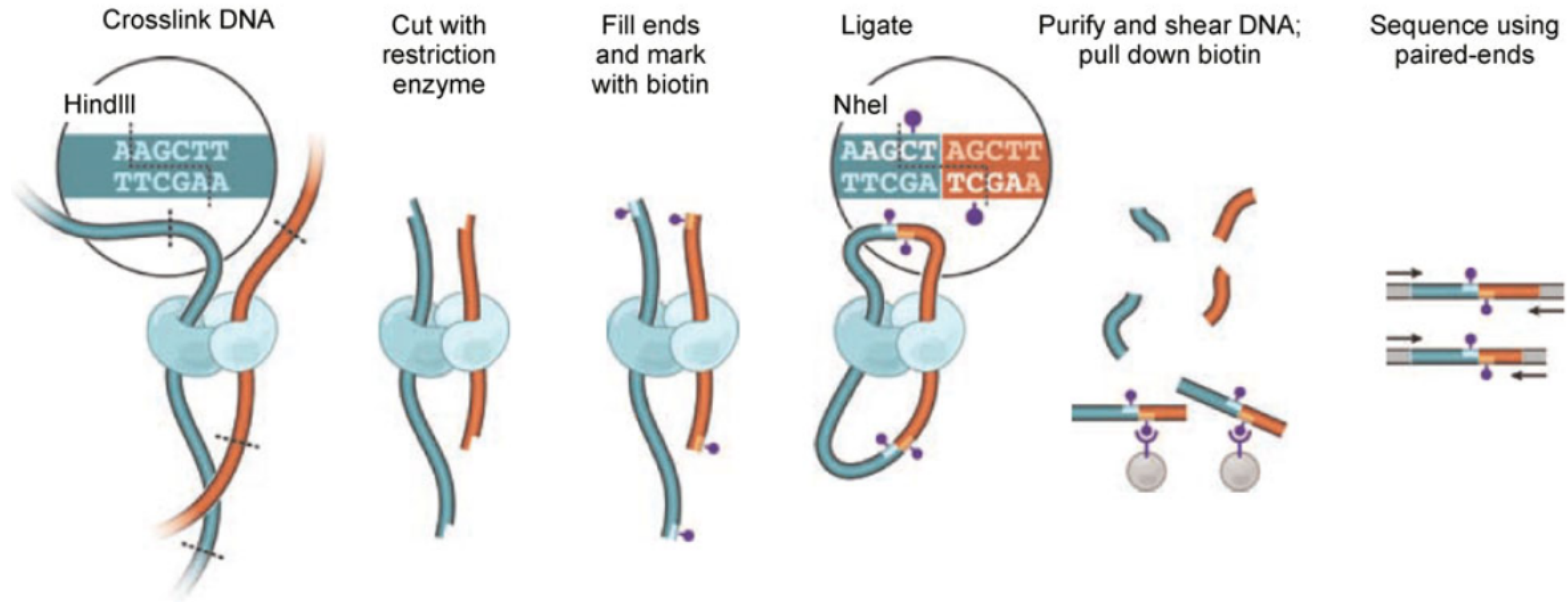


Hi-C contact matrices: filtering

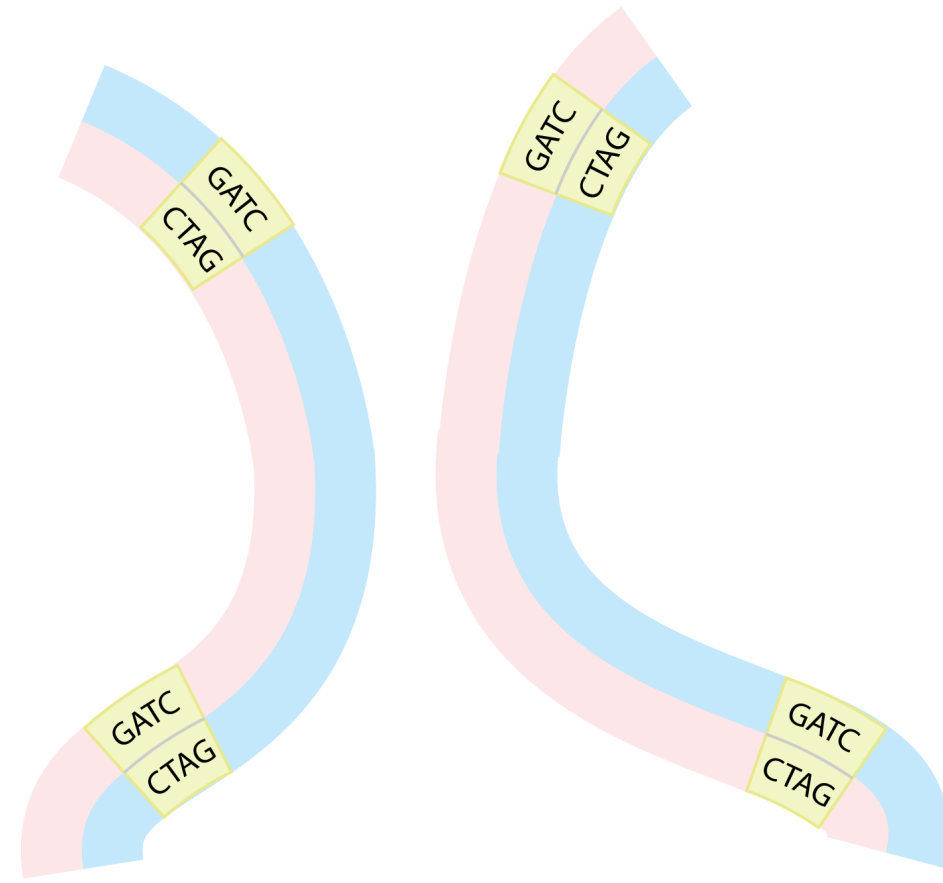
**Marco Di Stefano, David Castillo & Marc A.
Marti-Renom**
Structural Genomics Group (CNAG-CRG)



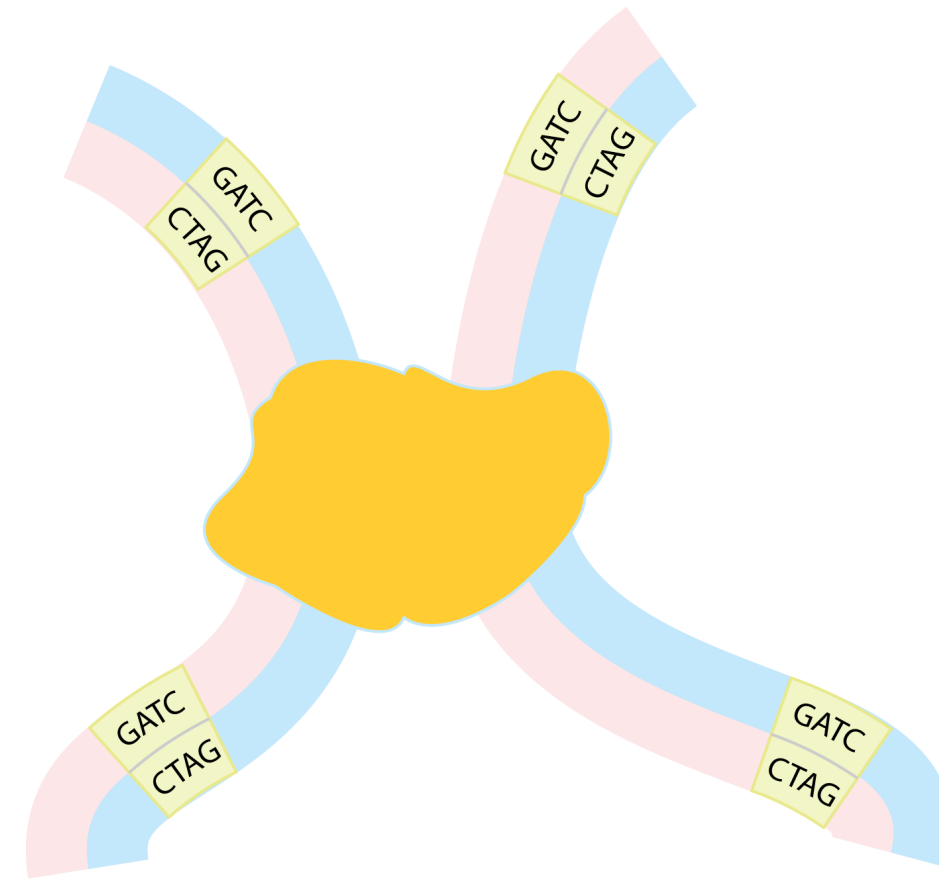
Hi-C experiment



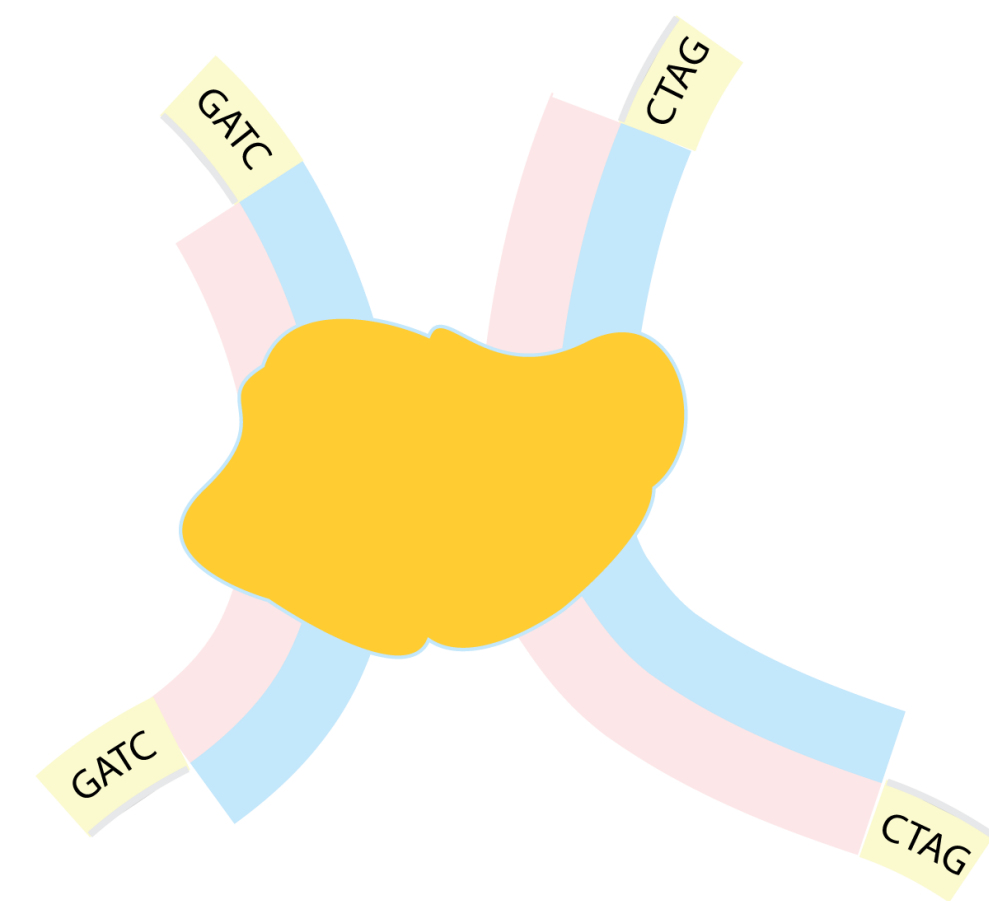
Hi-C experiment



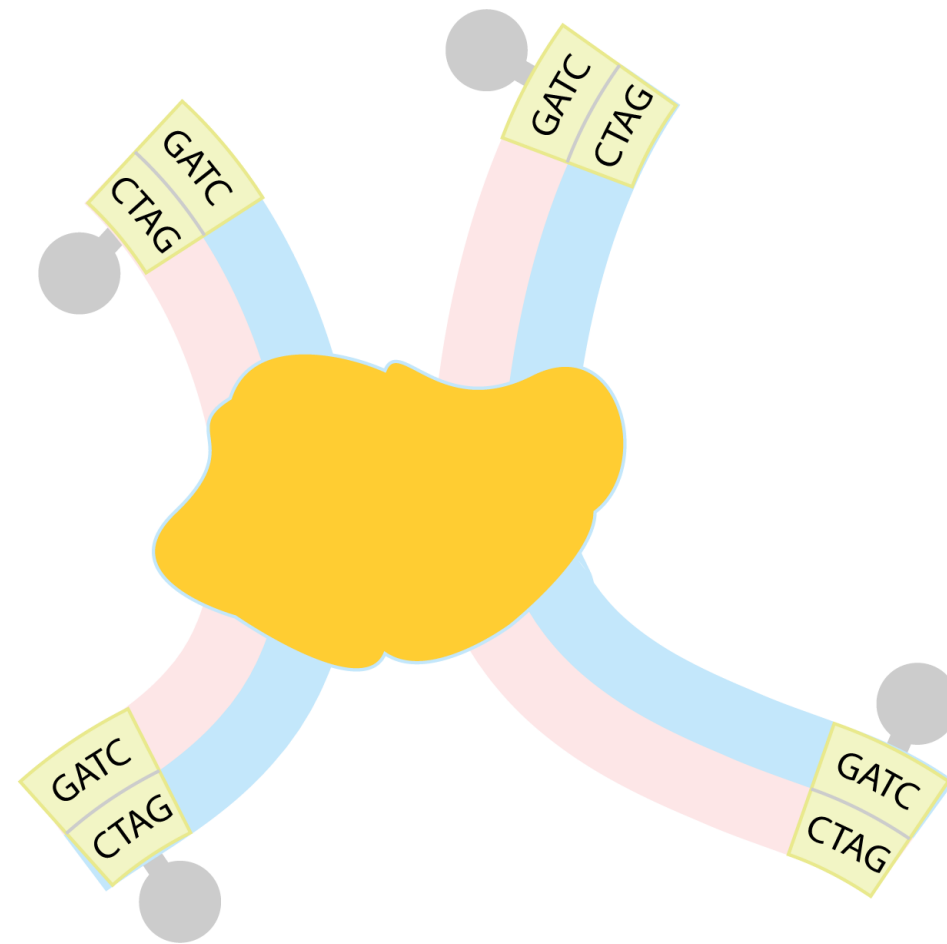
Two pieces of chromatin
are close in 3D



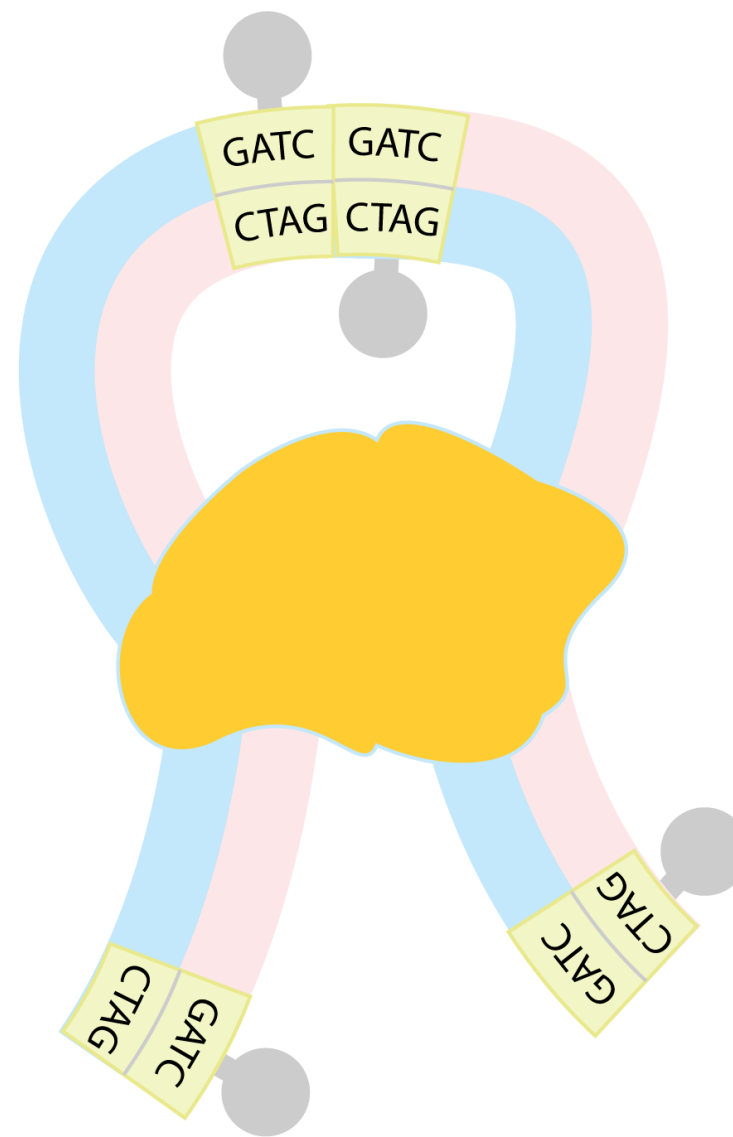
Cross-linking



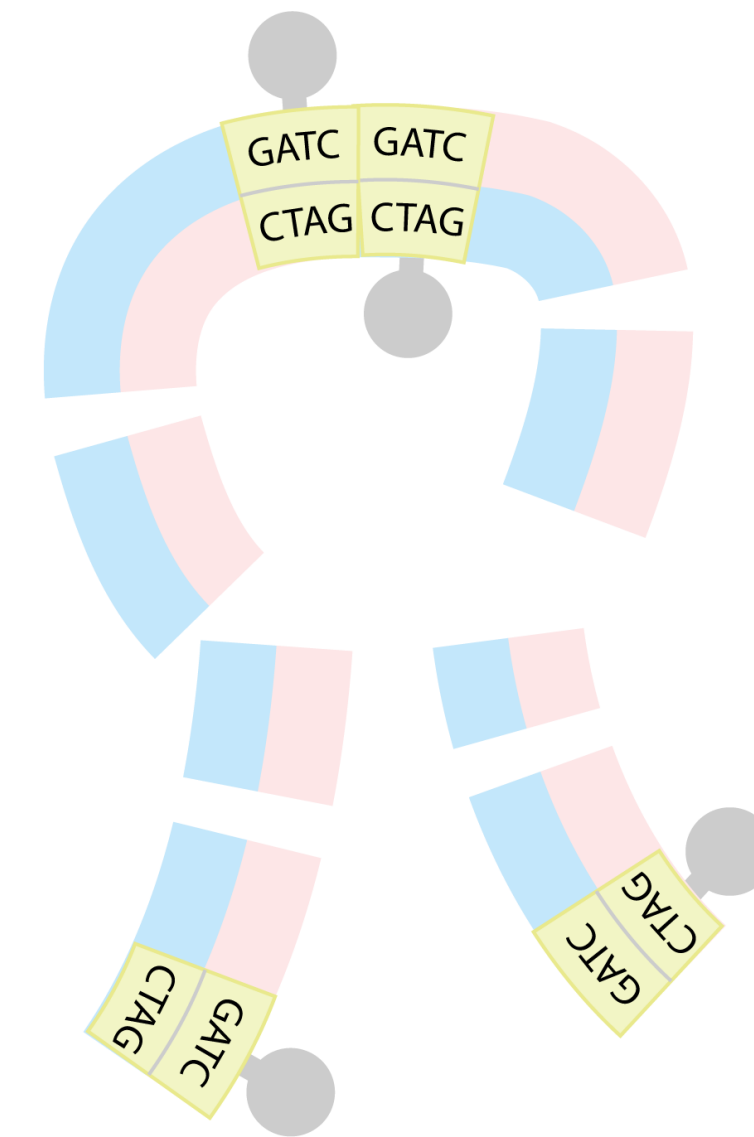
Digestion



Biotinylation

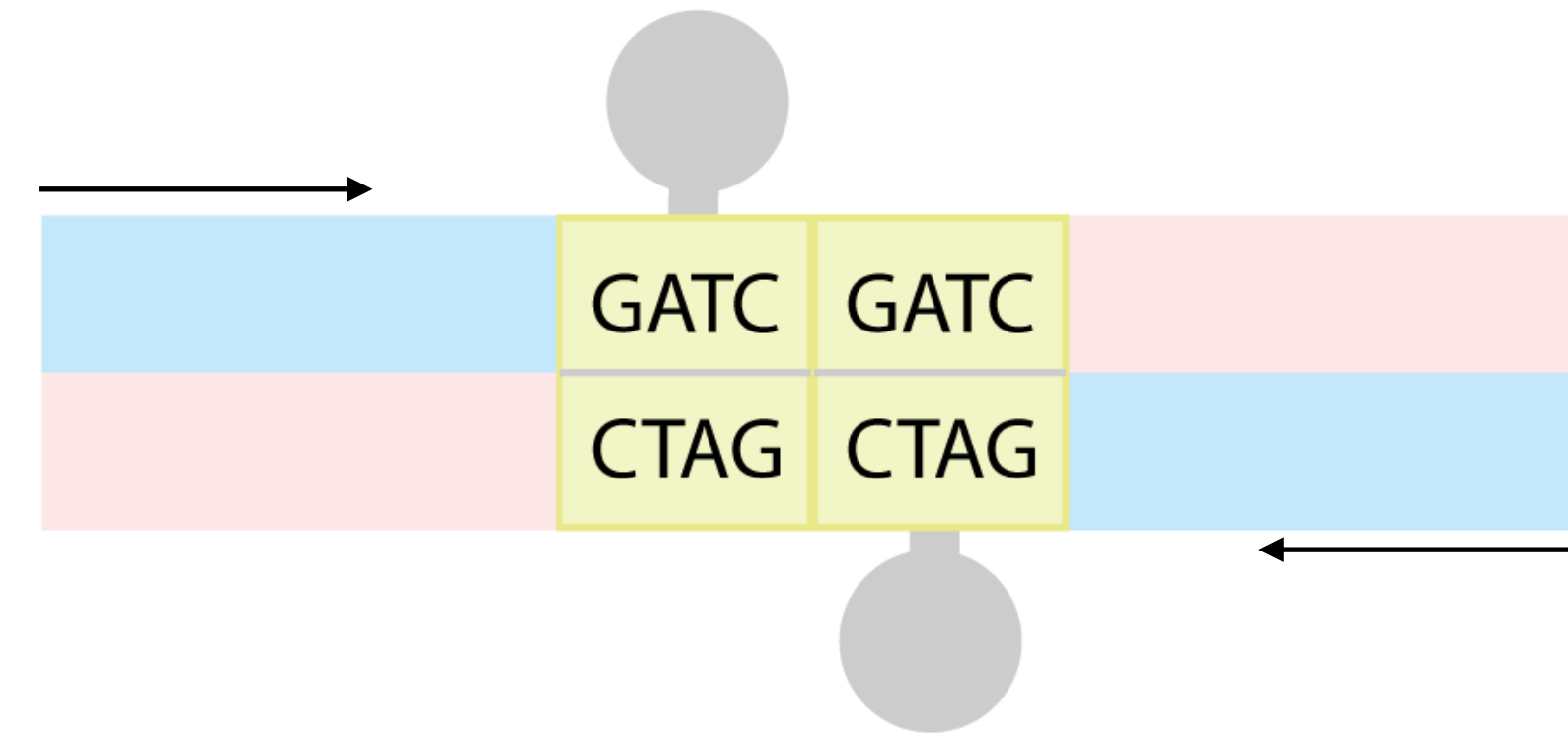
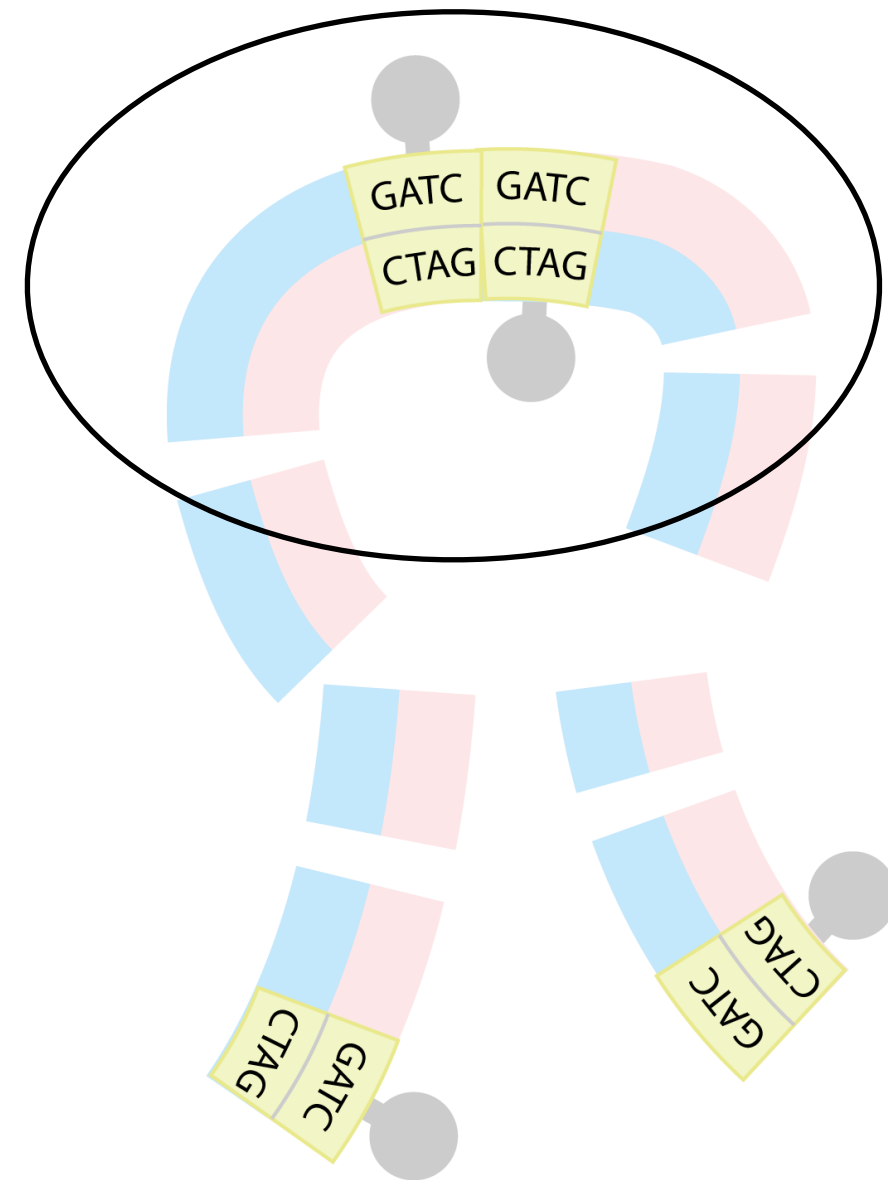


Ligation

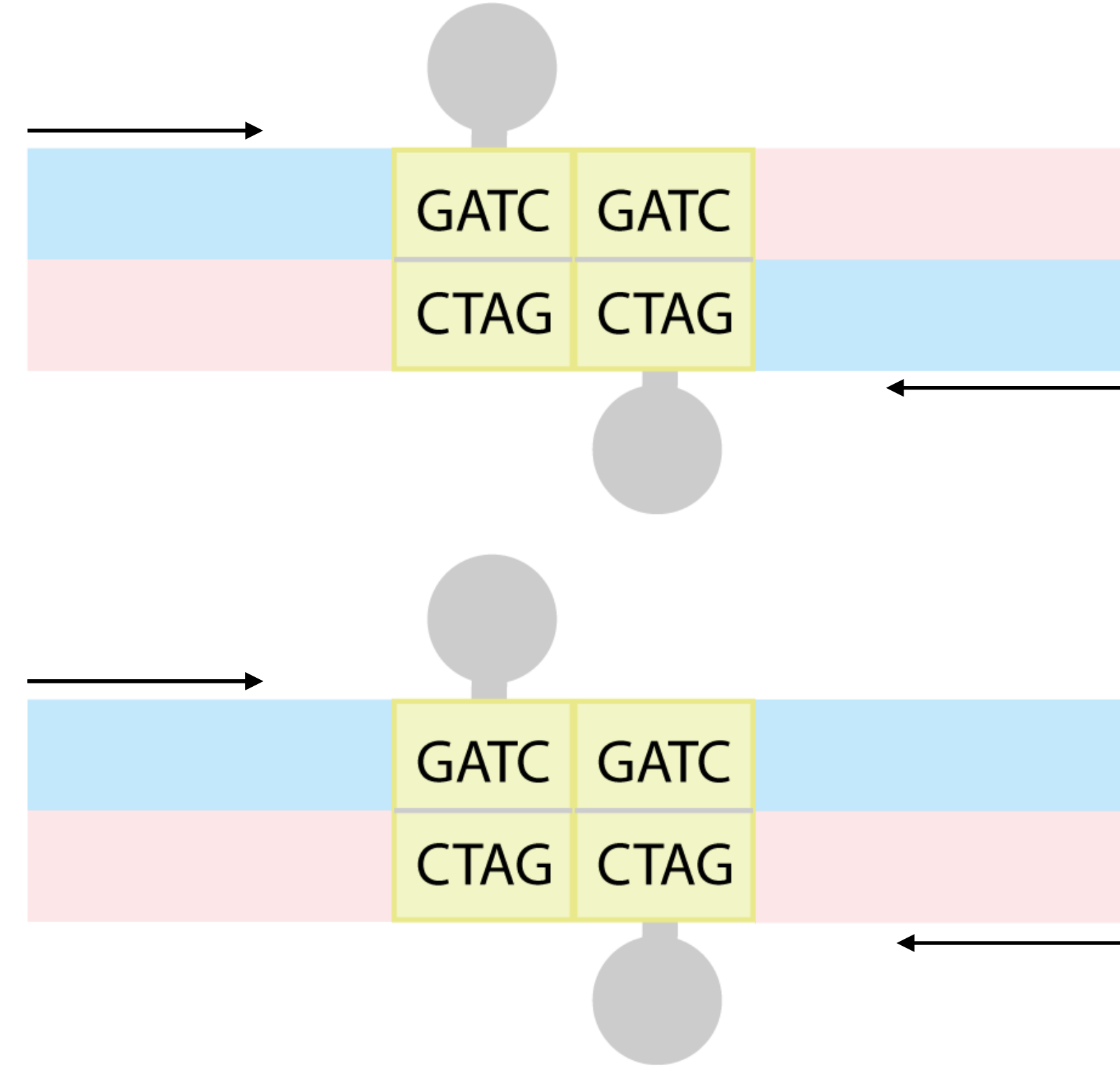
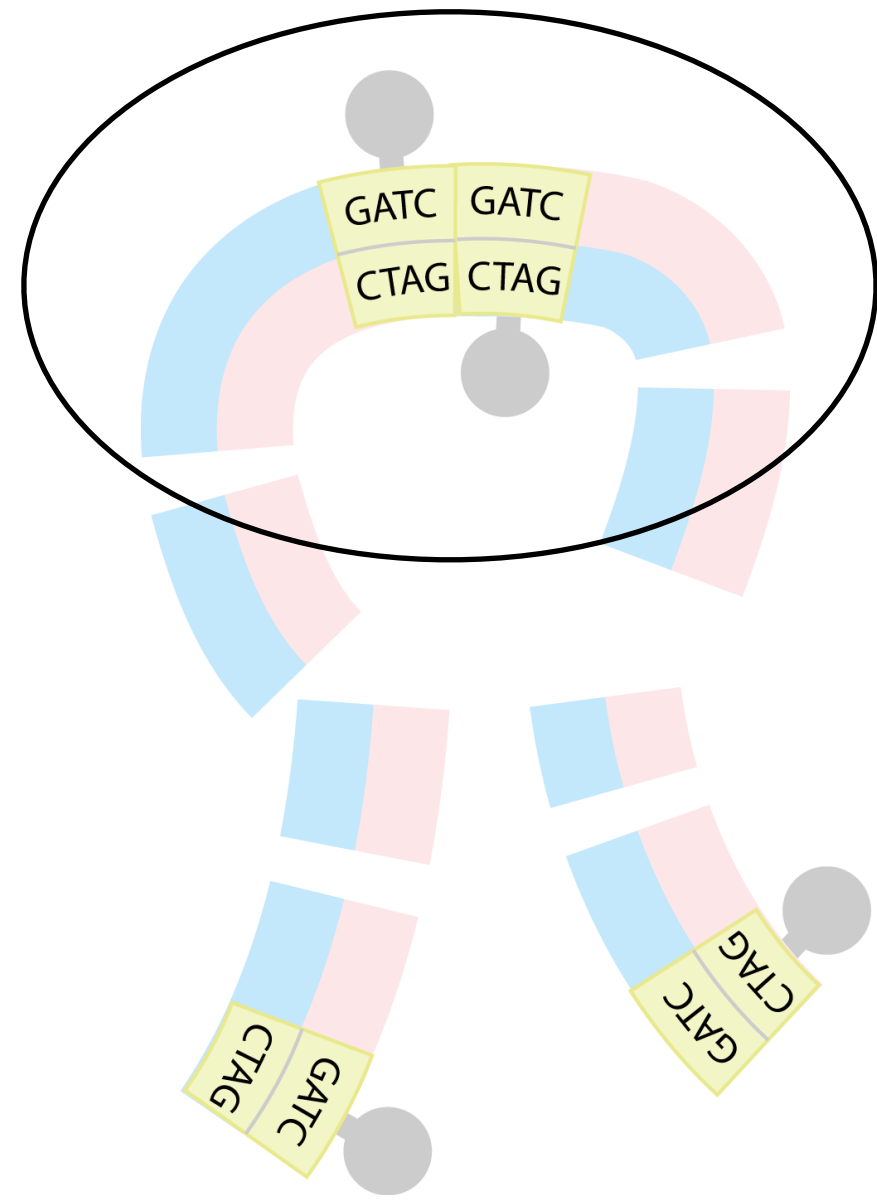


Shearing

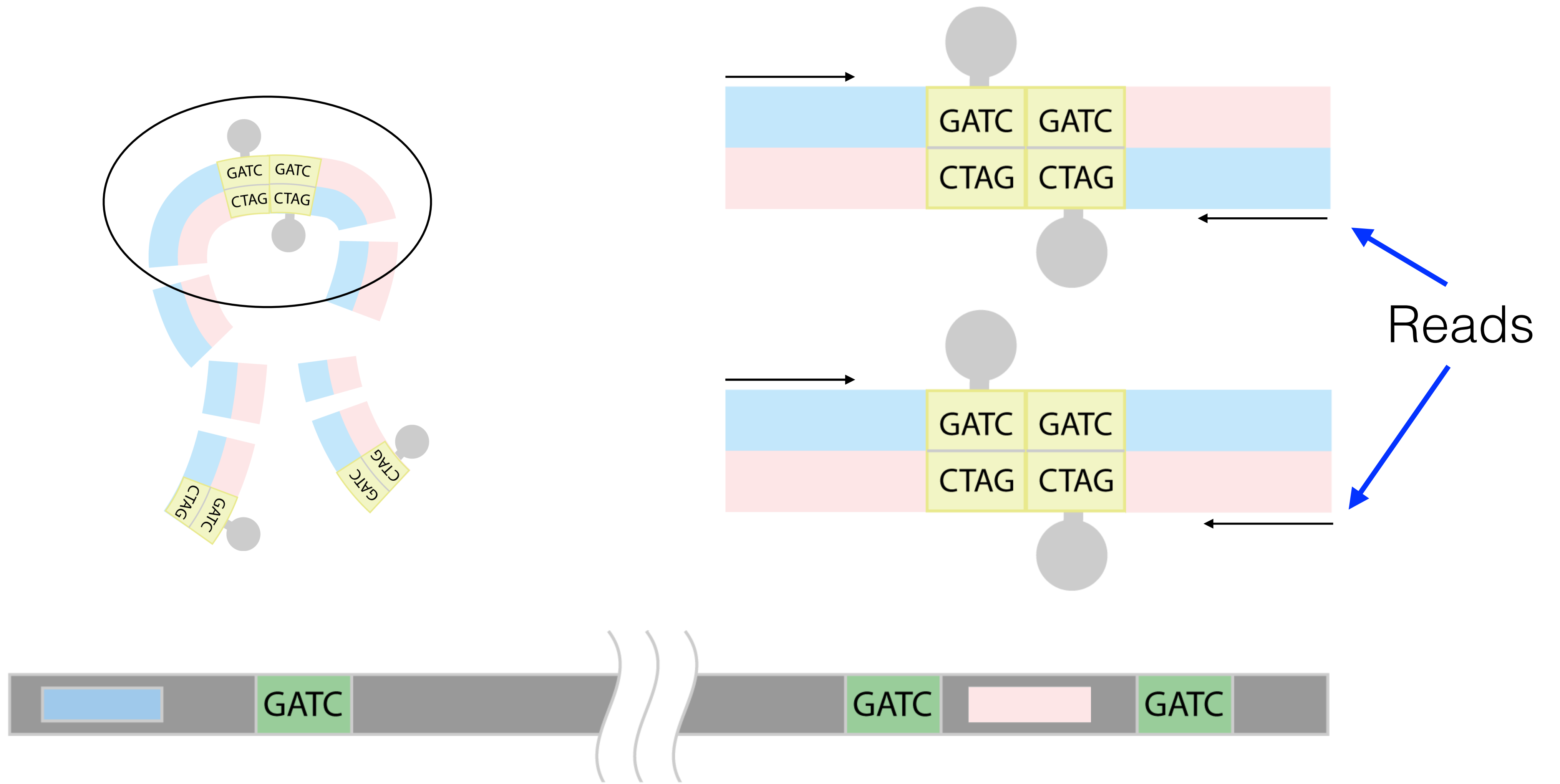
Valid pair



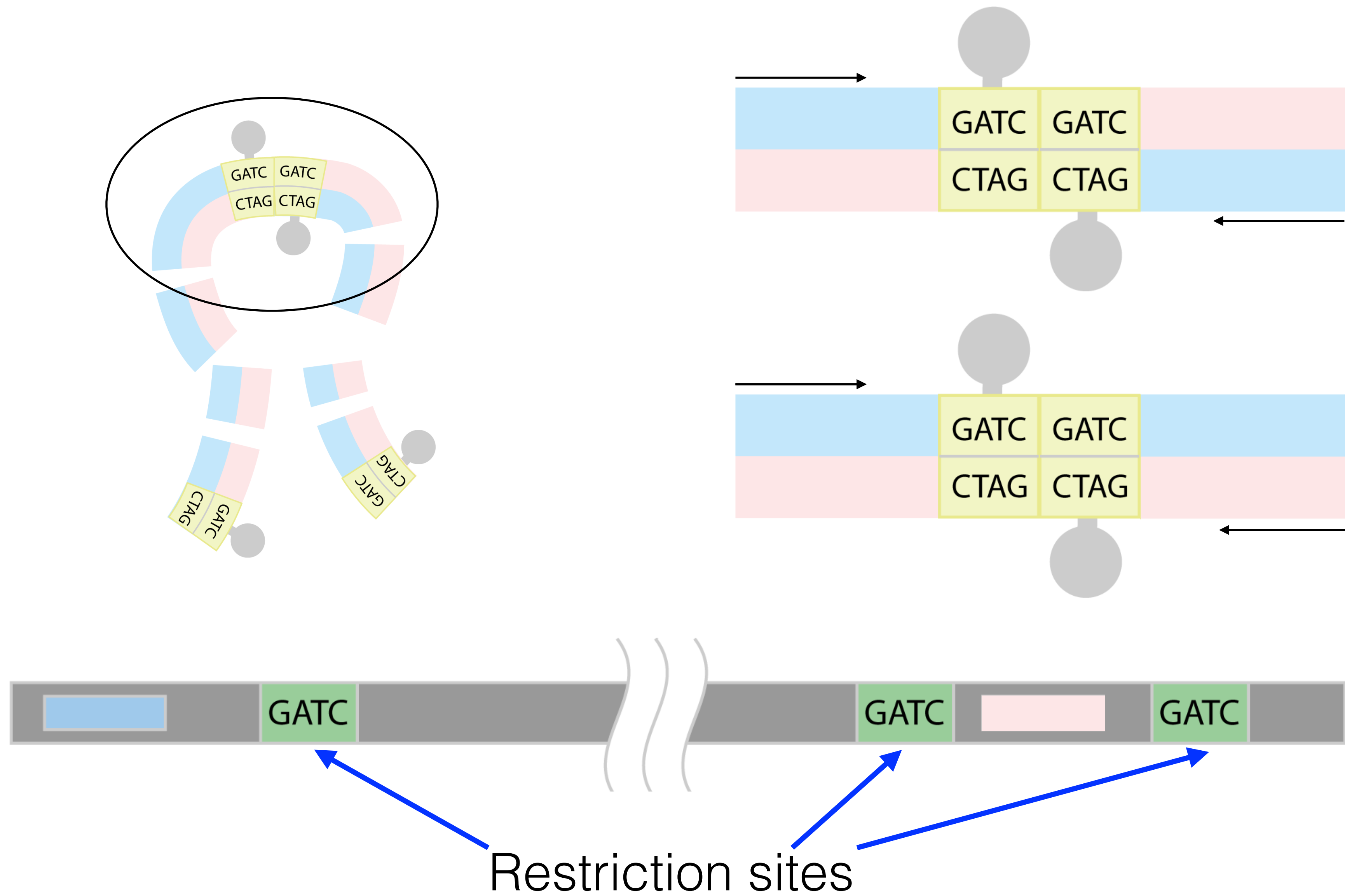
Valid pair (~36% of the mapped read-pairs)



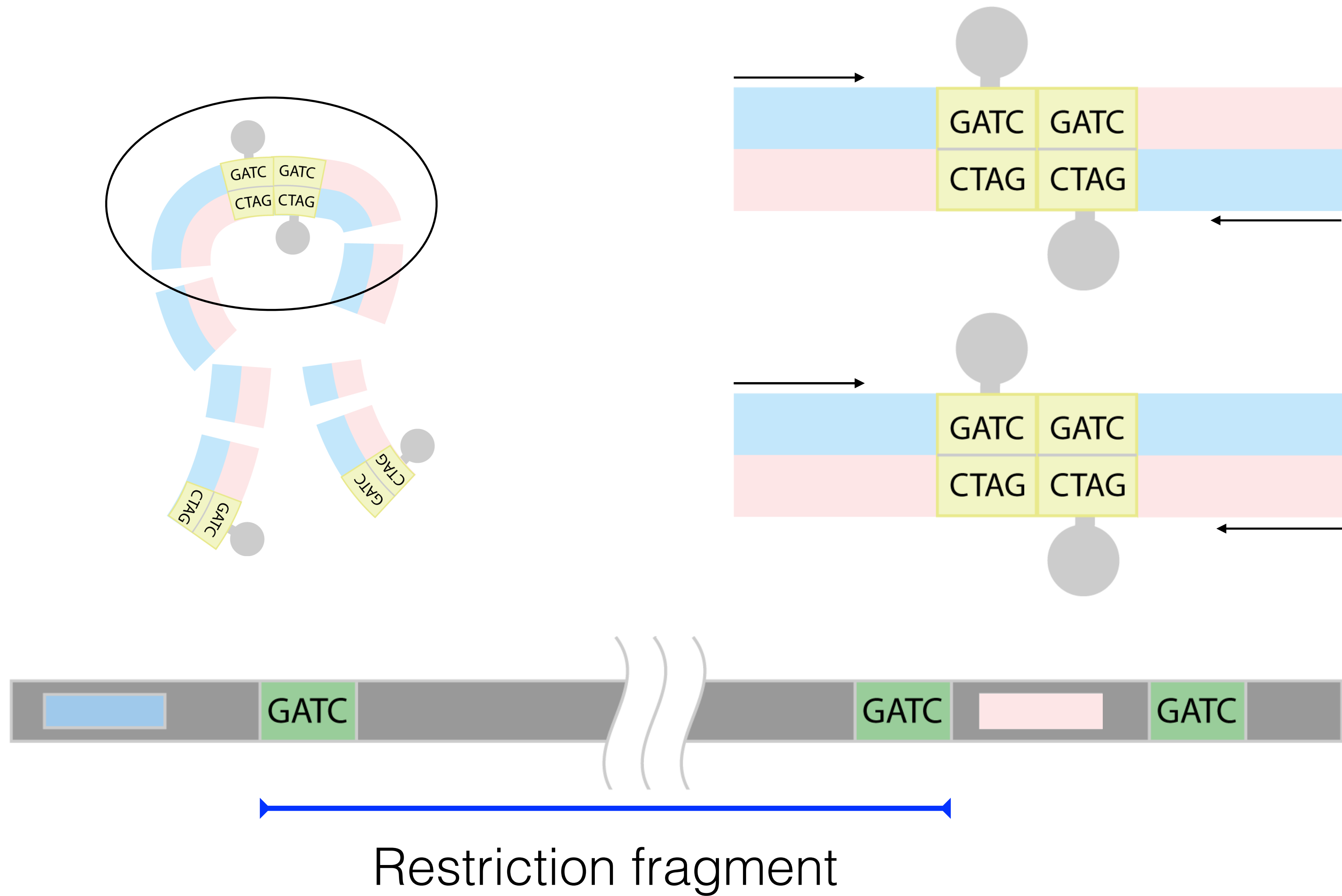
Valid pair (~36% of the mapped read-pairs)



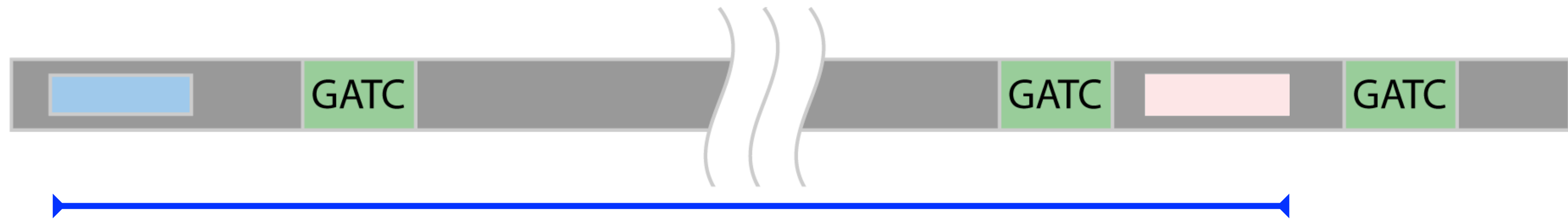
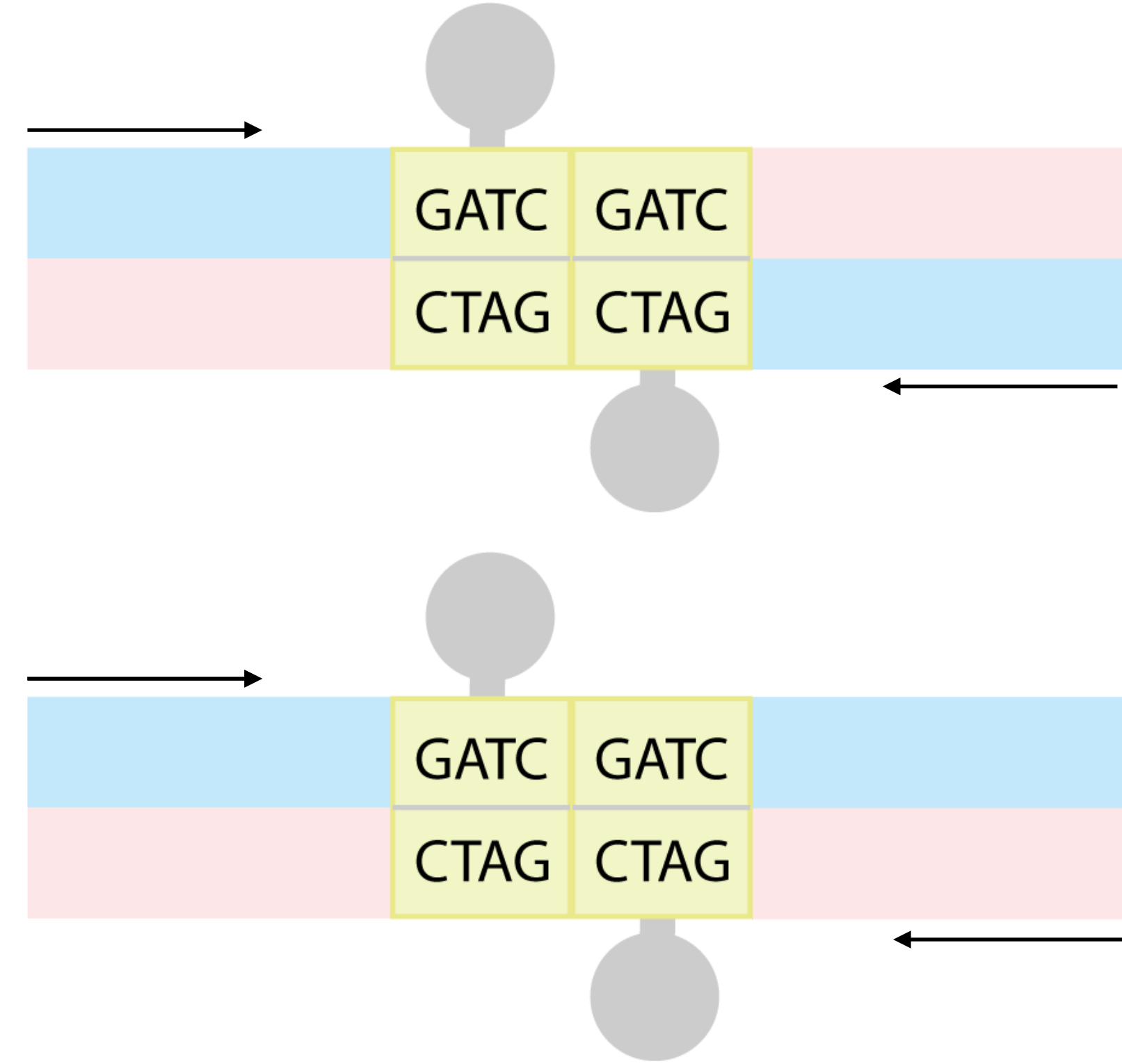
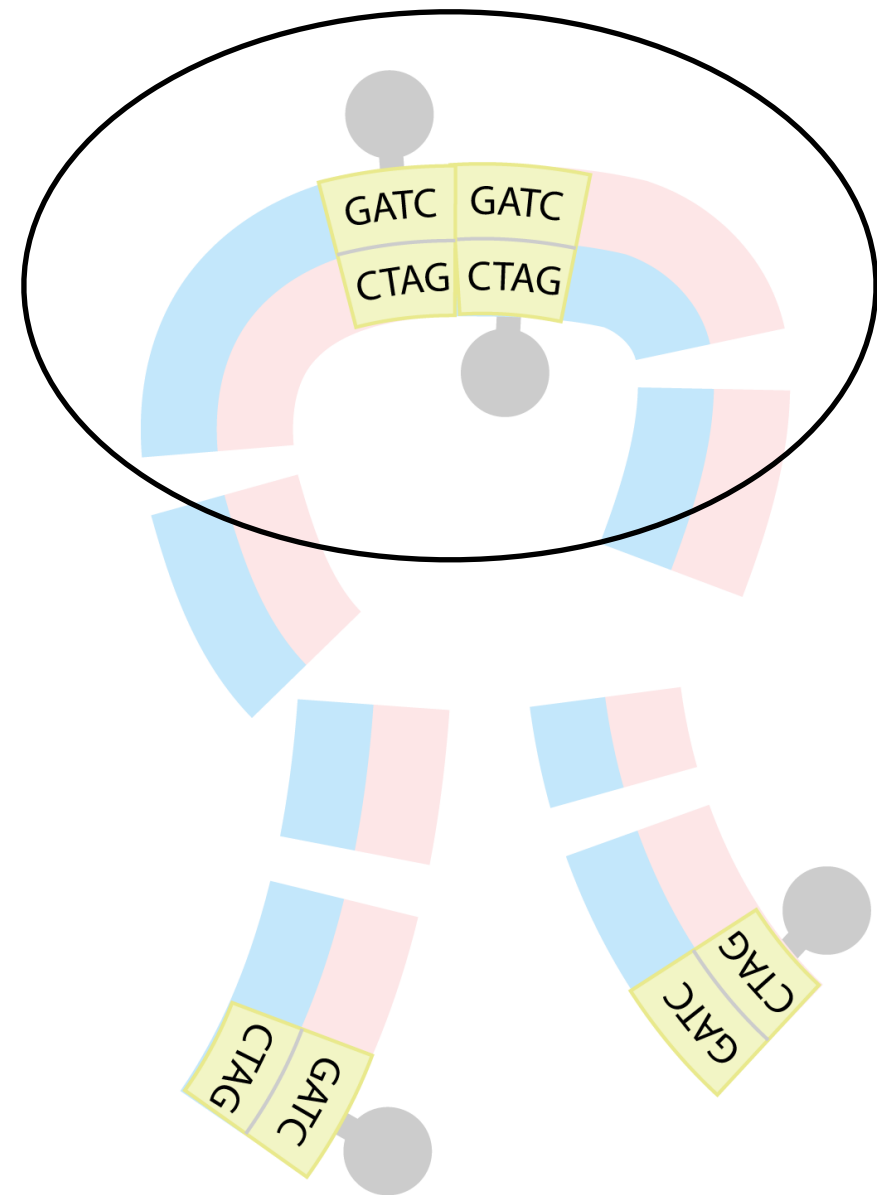
Valid pair (~36% of the mapped read-pairs)



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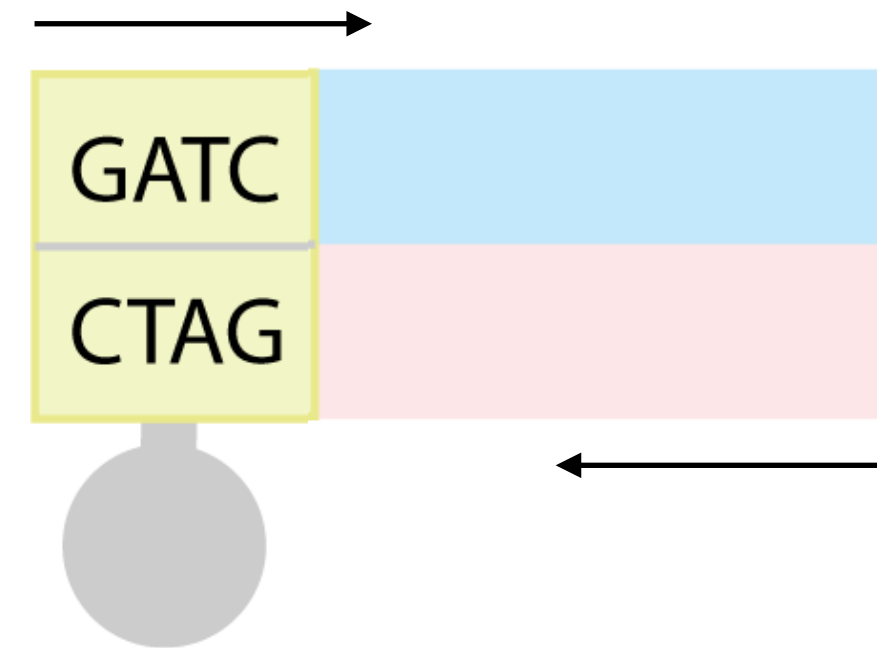
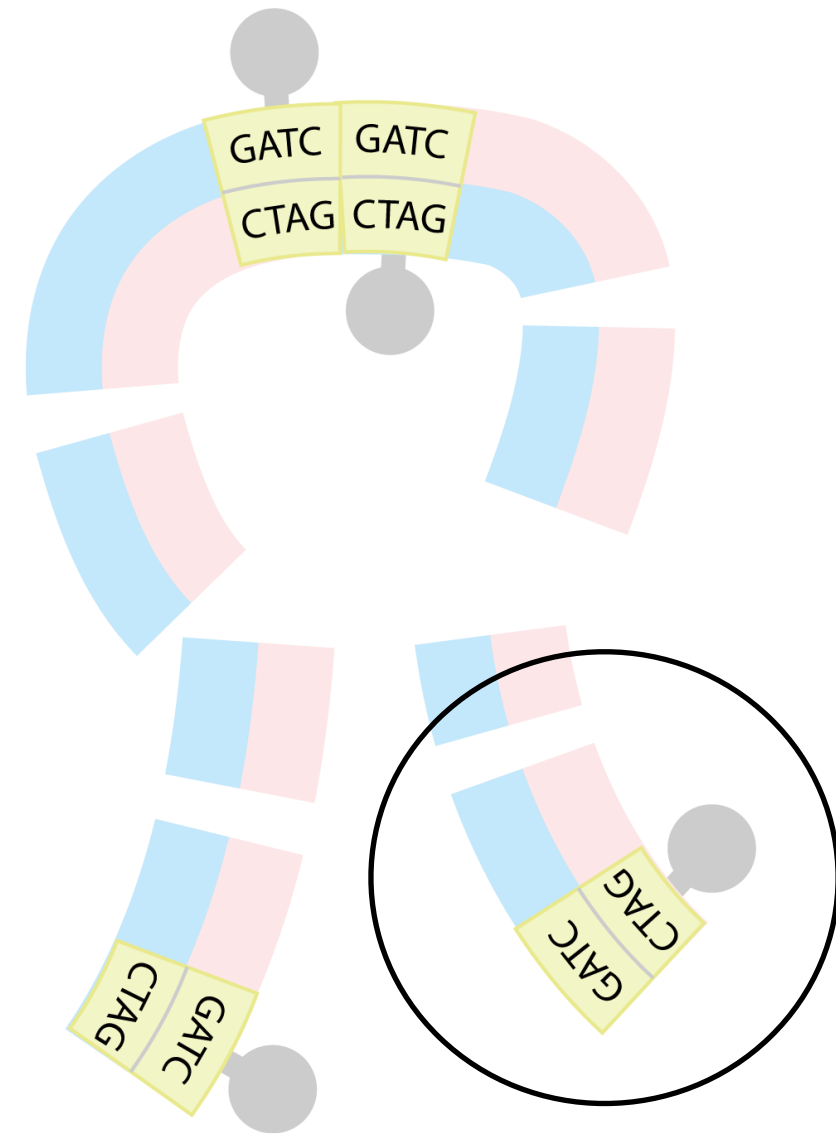


Valid pair (~36% of the mapped read-pairs)

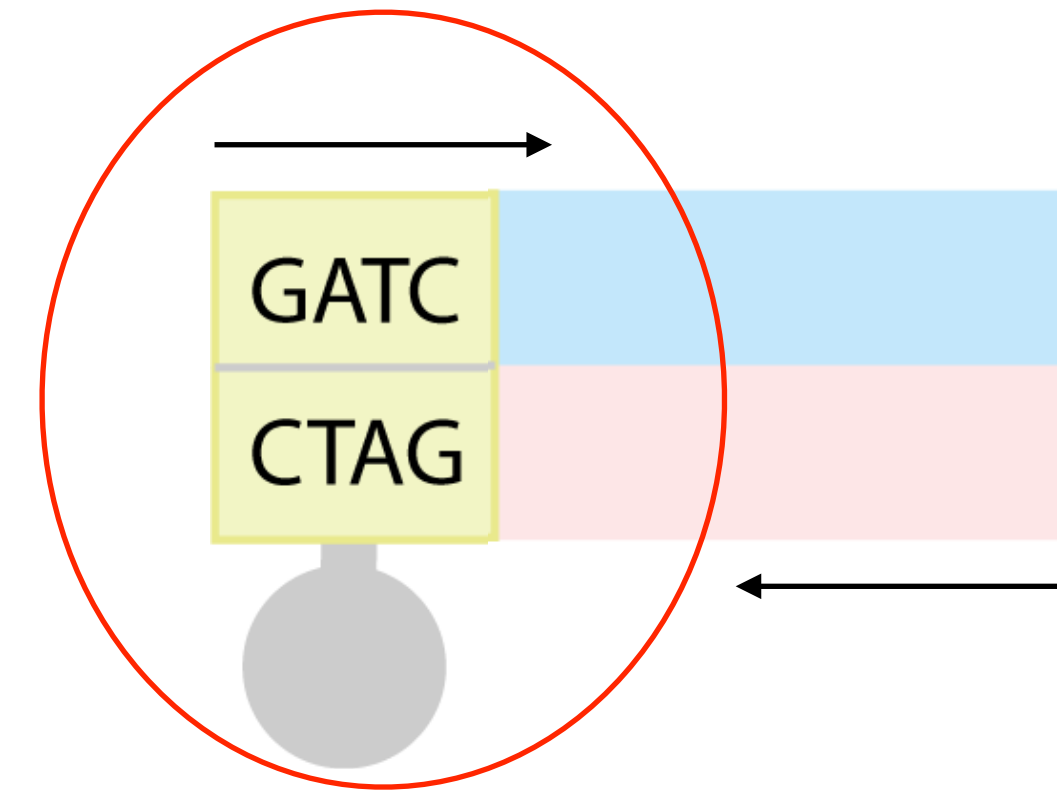
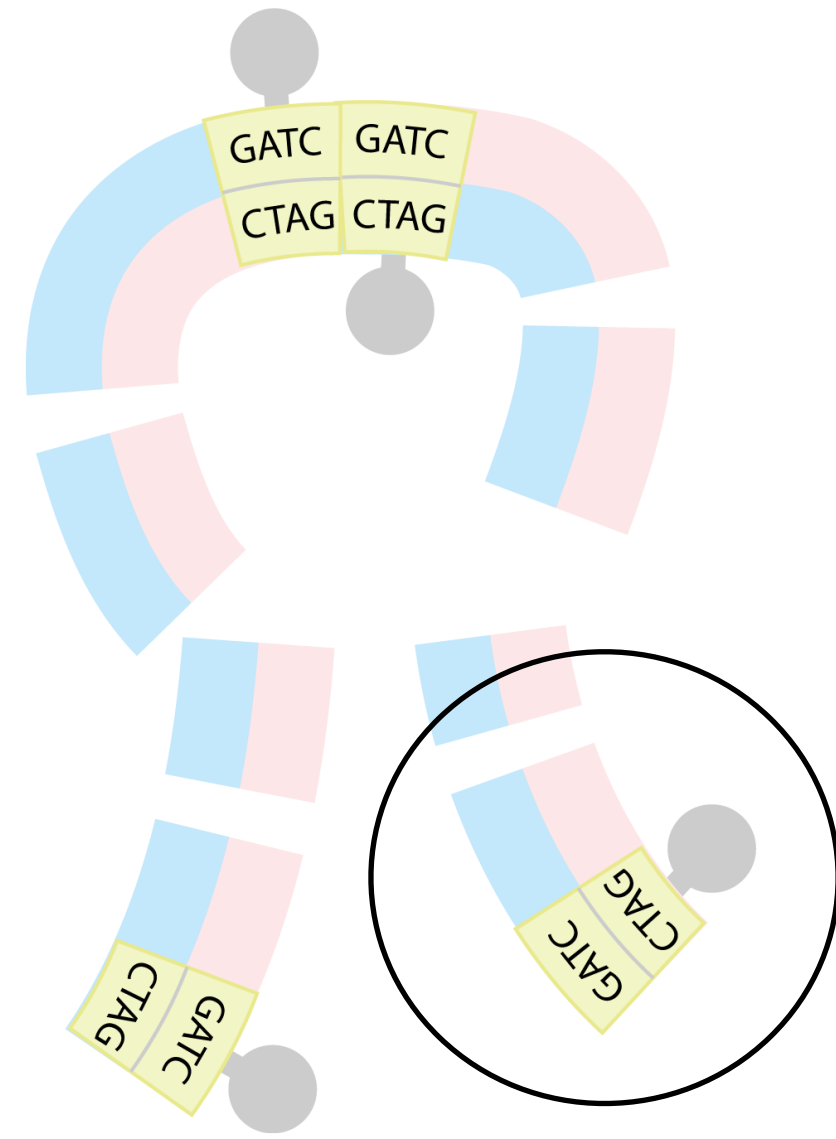


Fragment

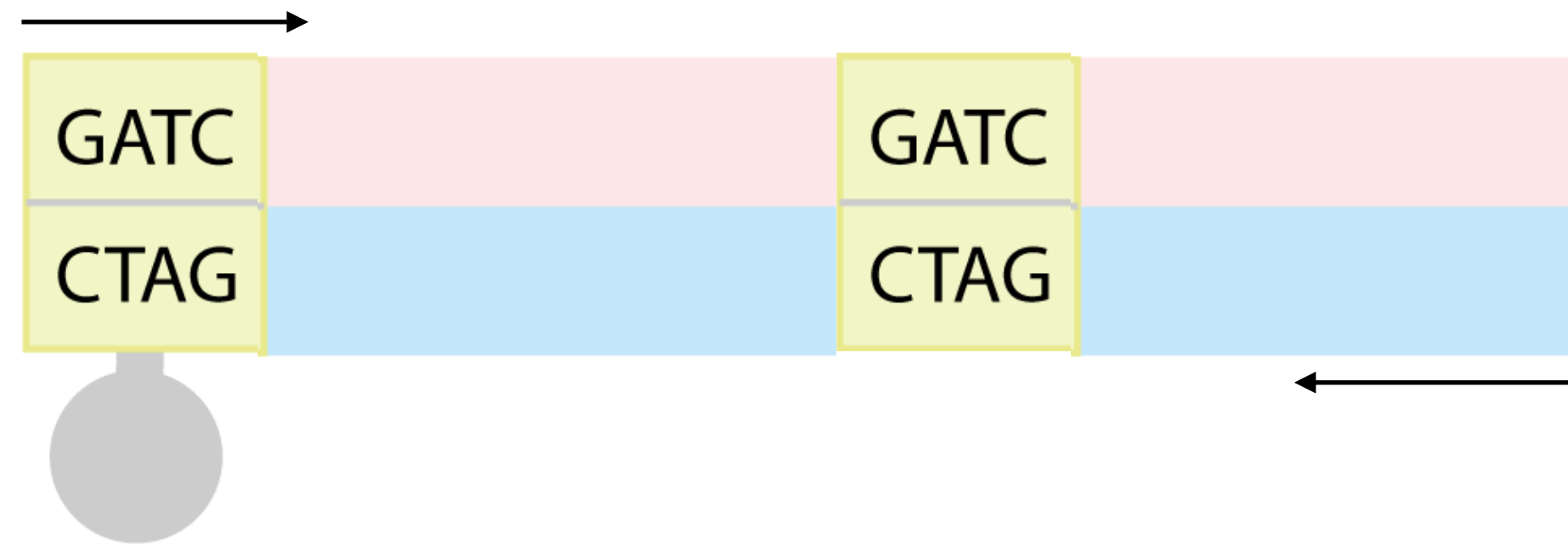
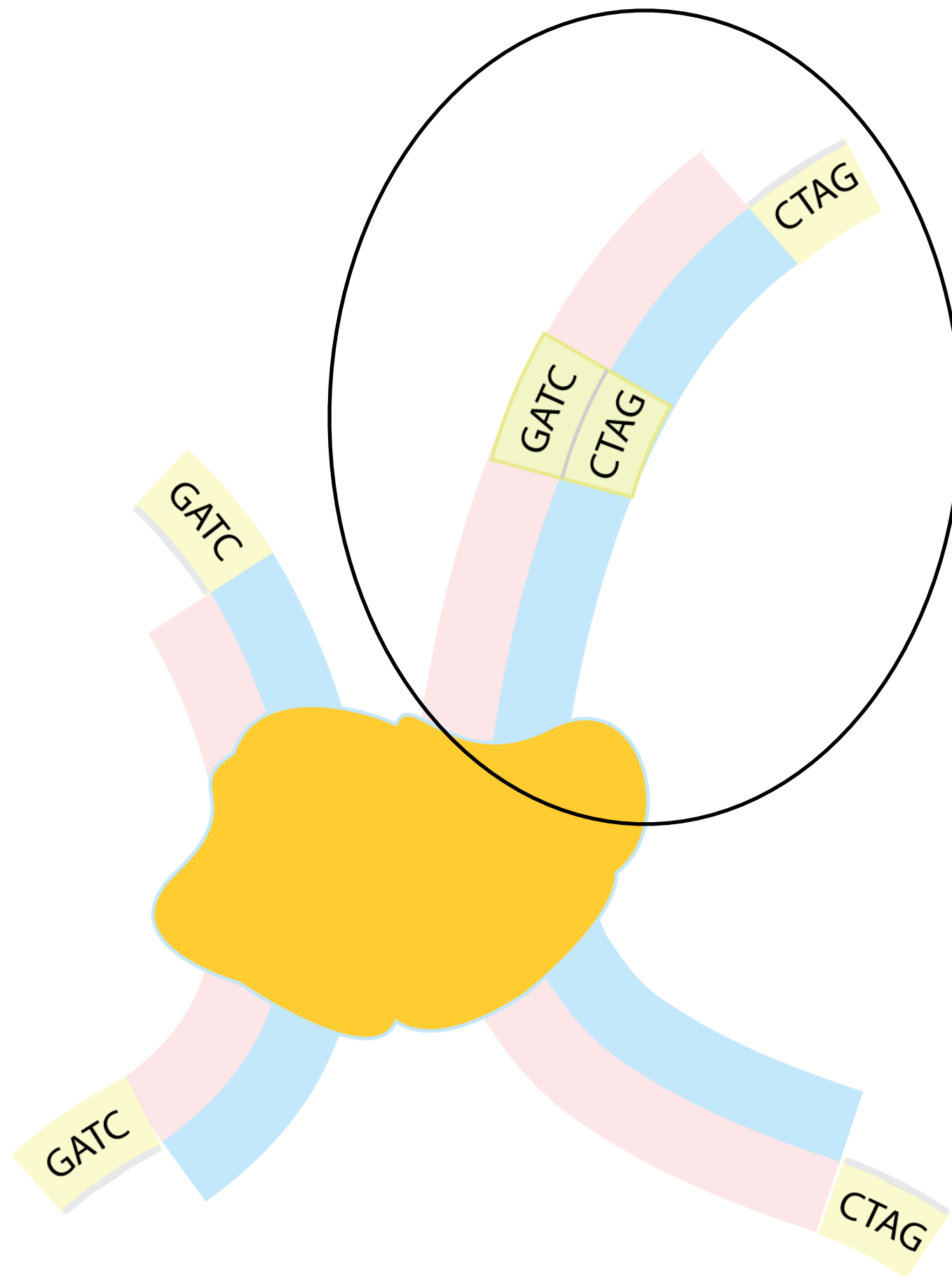
Dangling-end (~15%)



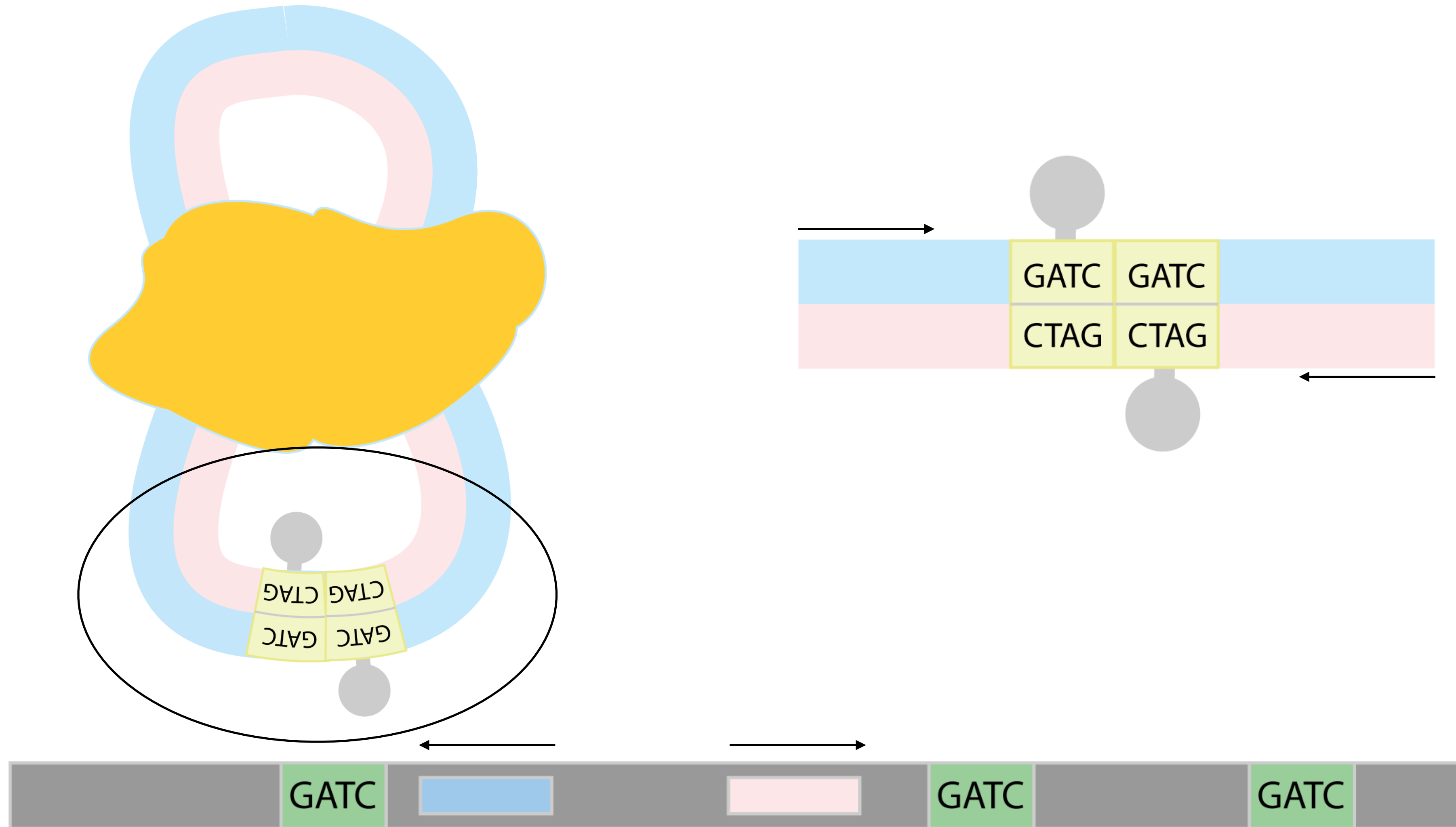
Dangling-end (~15%)



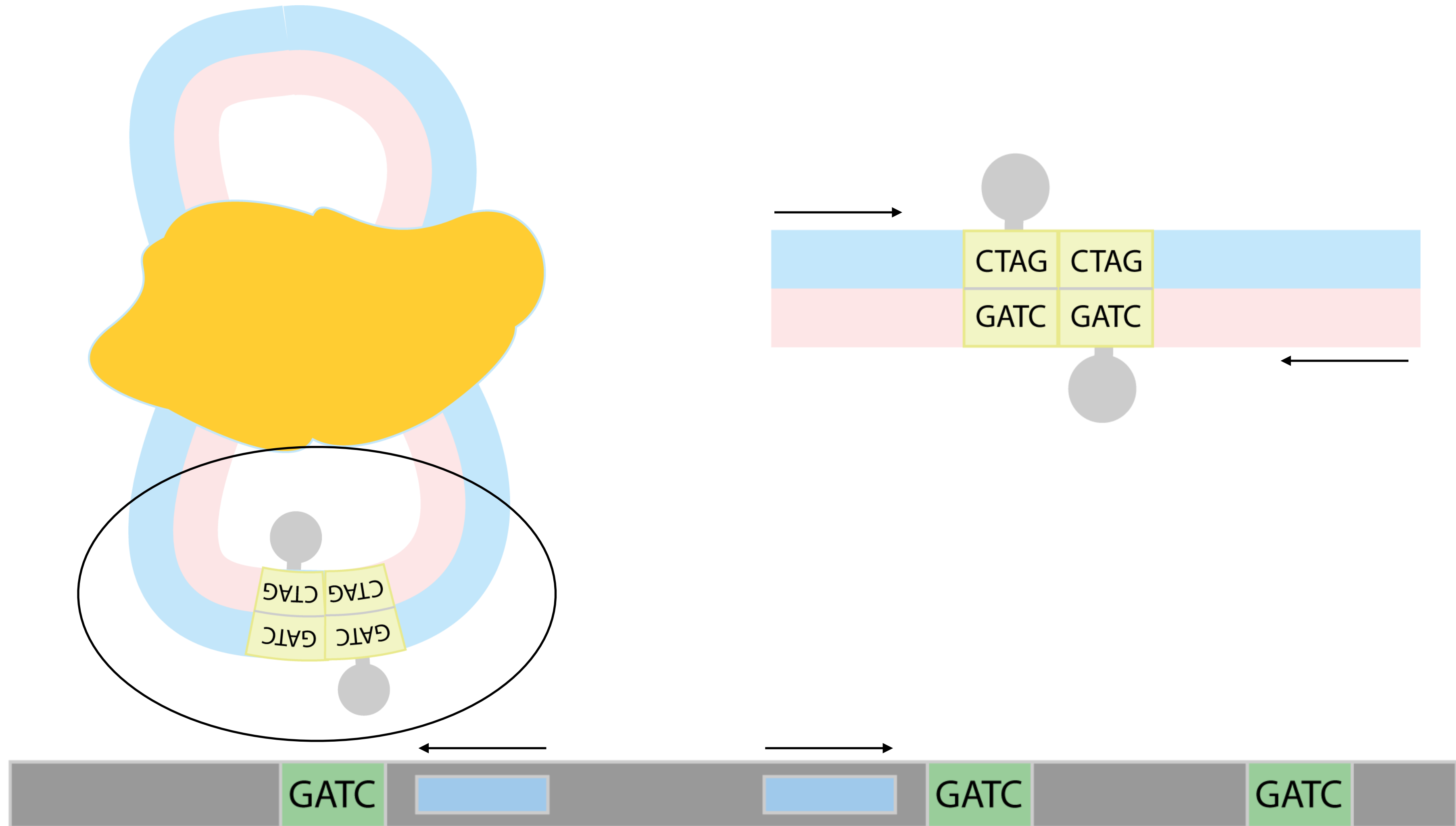
Extra dangling-end (~5%)



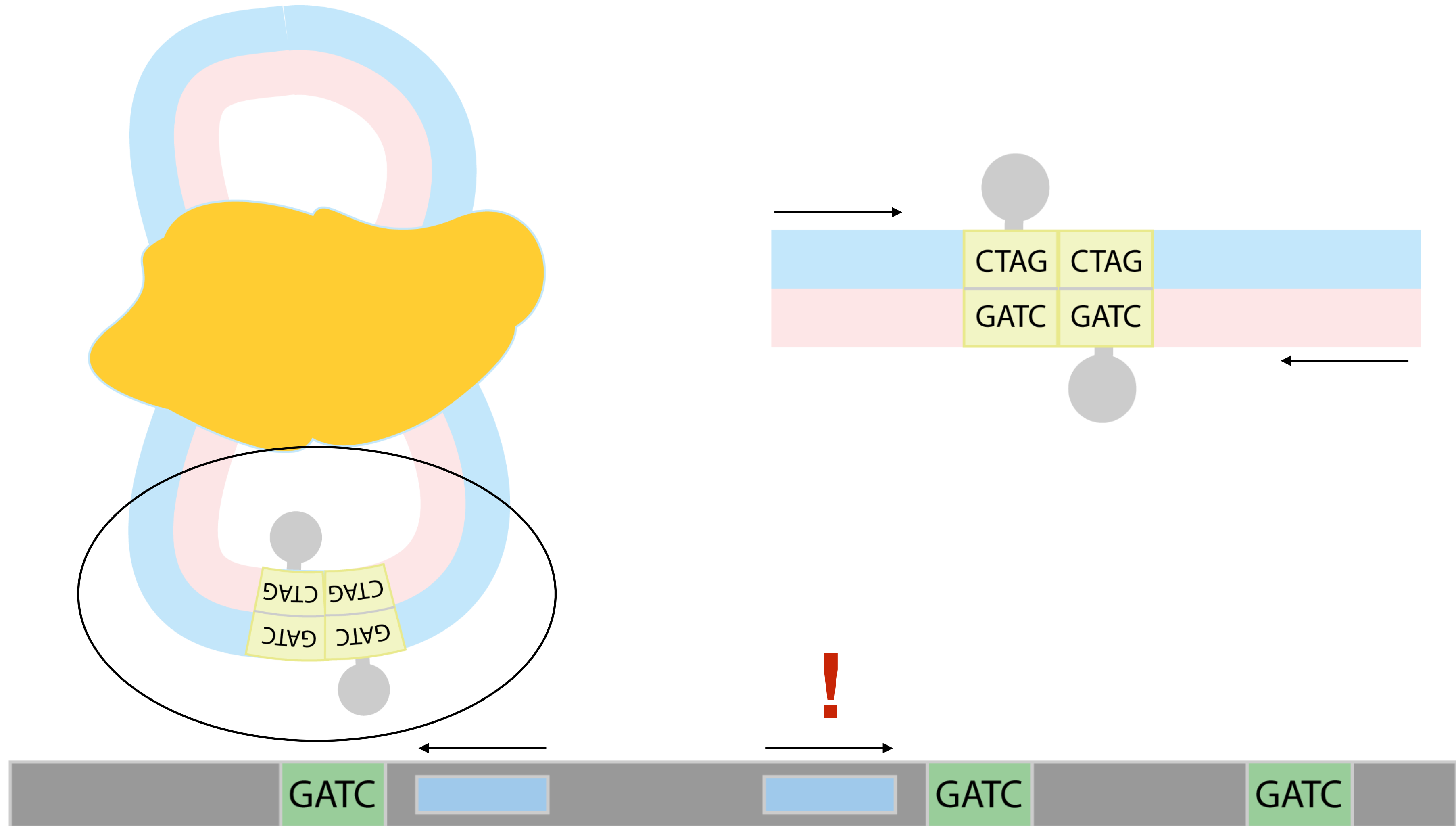
Self-circle (~10%)



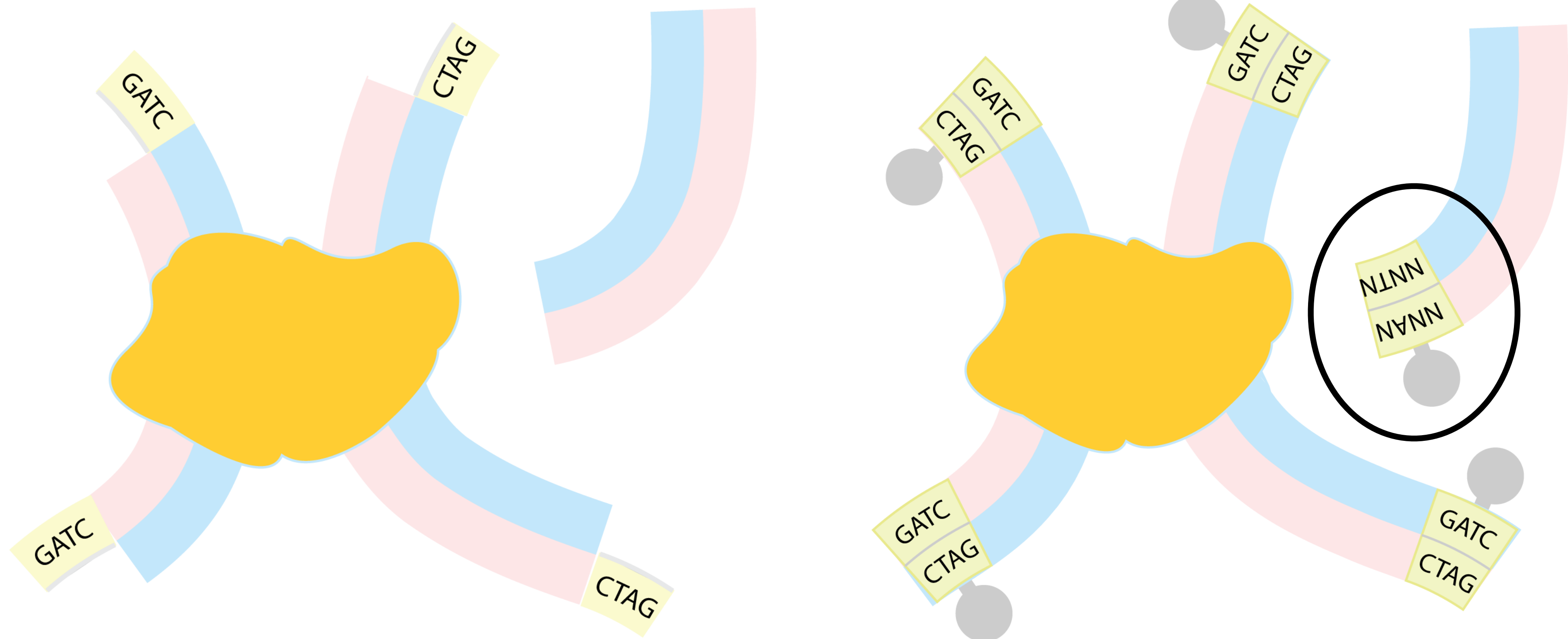
Error (<2%)



Error (<2%)



Random break (~20%)



> min_distance_to_RE

> min_distance_to_RE

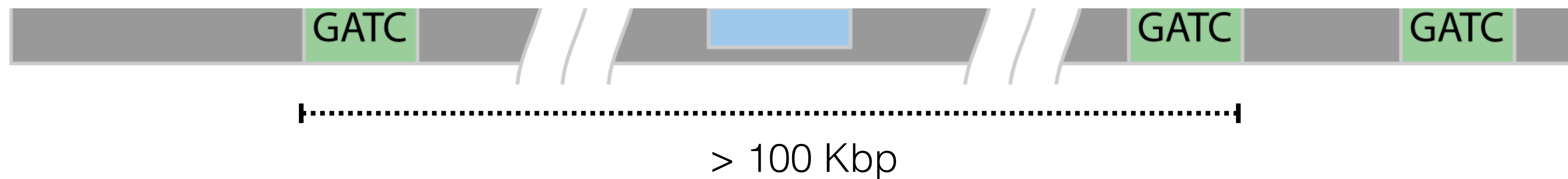
Too close from RE sites (~2%)



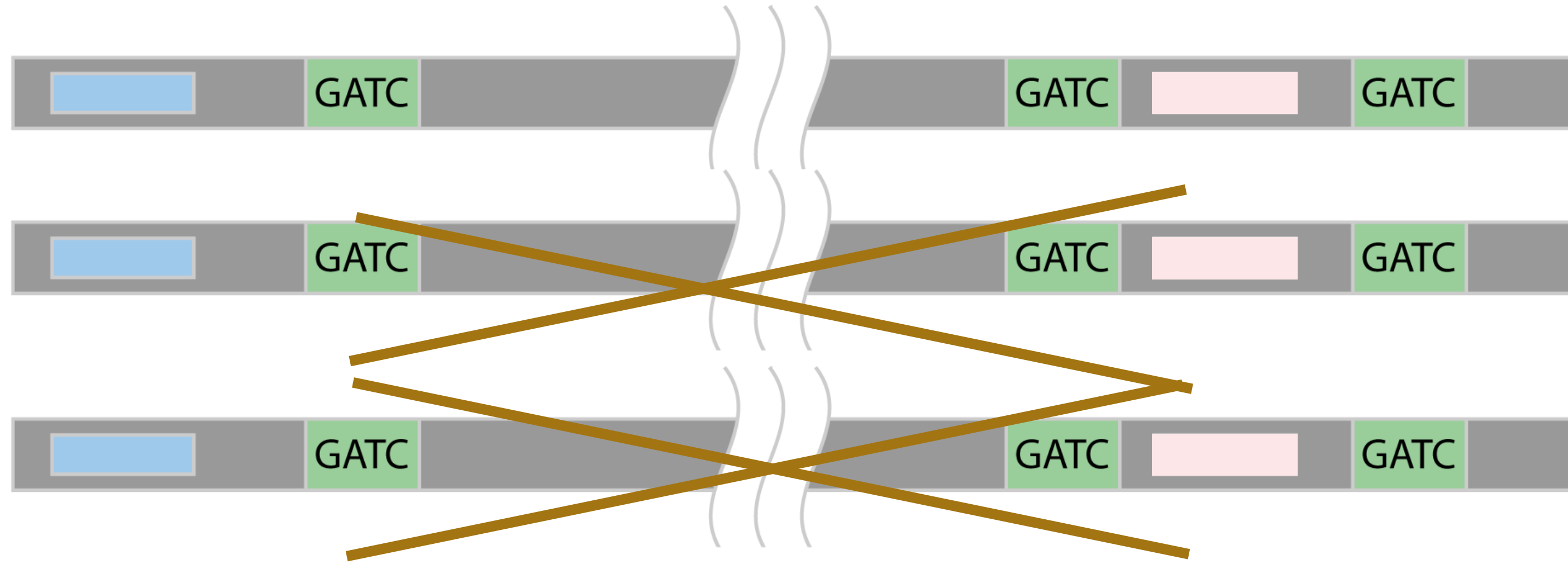
Too short (<1%)



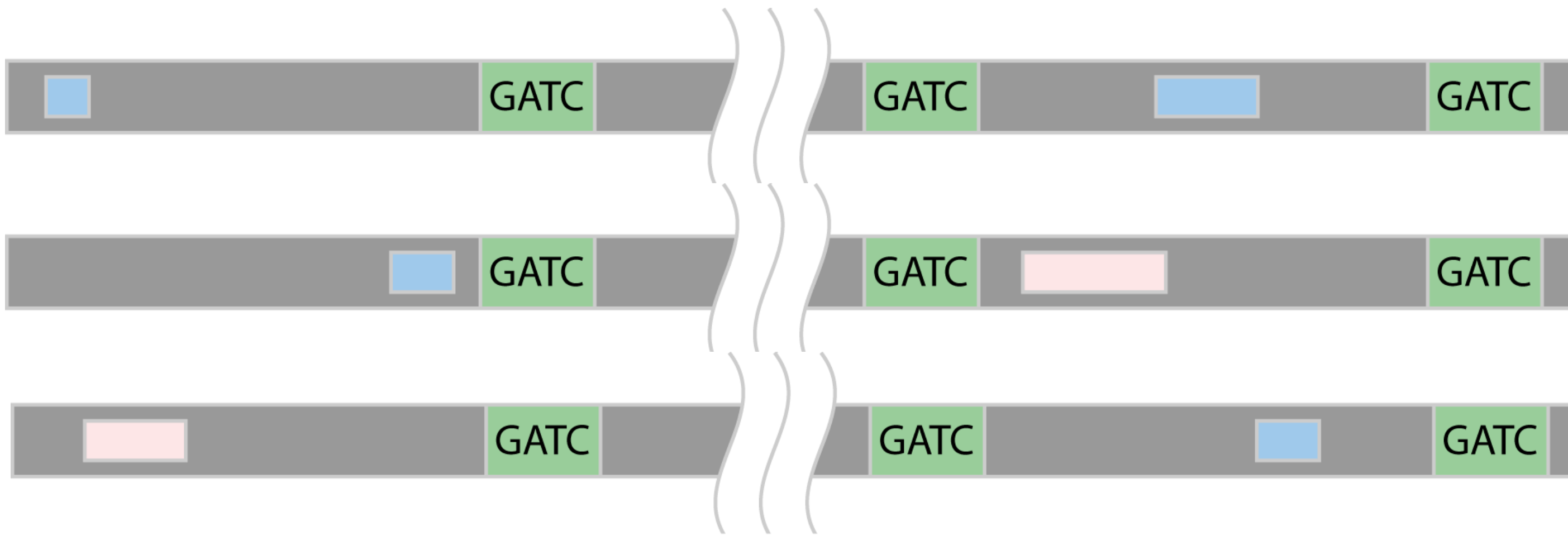
Too large (<1%)



Duplicated (~20%)



Over represented (<1%)



How much do we map?

- Valid reads: 80-90% each end => 64-81%
intersection => 60% of the intersection, 36% of the total
- 1% multiple contacts
- many of the reads will be lost in the filtering...