

# Biology - DNA Cheat Sheet

by emilyaltmann via cheatography.com/81523/cs/19518/

#### **Translation**

## **Gene Expression**

transfer of genetic info from DNA to RNA to protein

#### Codons

mRNA is read in groups of 3 nucleotides. Codes for amino acid

#### Transfer RNA (tRNA)

single stranded RNA of 80 nucleotides. Bonds to amino acids and mRNA codon

#### Ribosomes

catalyzes the peptide bonds between amino acids

#### 1) INITIATION

eukaryotes, ribosome small subunit recognizes and binds to the mRNA at the 5' cap. Initiator tRNA attaches at AUG codon

#### Ribsomal binding sites

1 site where mRNA binds, 3 sites where tRNA binds (**A site** - aminoacyl-tRNA site...... **P site** - peptide-tRNA site...... **E site** exit site, leaves the ribosome)

# 2) ELONGATION

initiator tRNA binds to ribosome, incoming tRNA binds to A site. H bonds form between mRNA codon and tRNA anticodon. Requires GTP (E)..... Peptide bond formed between A site and P site by ribosomes = longer peptide chain

# 3) TERMINATION

At stop codon, a protein release factor binds to A site. (Adds H2O instead of amino acid, polypeptide chain is released)

# **Polysome**

single strand of mRNA can be used to make multiple copies of a polypeptide simultaneously

## **Translation (cont)**

## **Polysome**

single strand of mRNA can be used to make multiple copies of a polypeptide simultaneously

#### **History**

#### Friedrich Miescher

discovered DNA. White blood cells from pus- isolated nuclei (high in P)

#### Frederick Griffith

studied bacteria that caused pneumonia (used Rough and Smooth Strains)defined it as transformation in cell's function

#### Avery, LcLeod, and McCarty

purified S strain bacteria, added it to R strain bacteria. No S cells appeared in the tube w/ no DNA, but they did appear in that w/ no proteins and no RNA

# Hershey & Chase

used bacteriophage to infect bacteria. Light up DNA and Protein case, only DNA was passed to bacteria.

#### **DNA Replication**

# **Replication Origin**

Specific sites where replication begins, then bidirectional. (can be more than one)

#### Helicase

enzyme that disrupts H bonds, creating replication fork

## **Single Stranded Binding Proteins**

relieve pressure. bind to unwound single stranded DNA to keep strands apart

# **Topoisomerases**

relieve pressure. break bonds in DNA then reform them.

## **DNA Replication (cont)**

## **RNA Polymerase**

adds primer (RNA nucleotides)

## **Priming**

Required as DNAP (DNA Polymerase) can only add nucleotides, but RNAP can start a new chain.

#### **DNA Polymerase III**

adds nucleotides to the 3' end of pre-existing nucleotides. (hydrolyzes last two phosphate groups)

# **Leading Strand**

synthesized continuously, moving along replication fork

#### **Lagging Strand**

synthesized in short, discontinuous segments of 1000-2000 nucleotides (**Okazaki fragments**)

#### DNAP I

replaces RNA primer with DNA

#### **DNA Ligase**

joins broken pieces of DNA by catalyzing formation of phosphodiester bonds

## DNAP III

can correct errors as it moves down the strand

# **DNAP II**

checks for errors and corrects them

# **Transcription**

# **Archibald Garrod**

studied patients w/Alkaptonuria (pee turns black with O2), faulty genes meant they couldn't break down alkapton (no enzyme)

# George Beadle & Edward Tatum

bread mould, using x-rays to create mutations- and then the moulds couldn't grow



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## **Transcription (cont)**

## **Central Dogma**

**TRANSCRIPTION** (DNA- mRNA) nucleus **TRANSLATION** (mRNA- protein) ribsomes The Flow Of Info

#### 1) INITIATION

RNAP binds to promoter. Composed of TATA box (less energy to break H bonds) RNAP recognizes the promoter and begins unwinding DNA

# 2) ELONGATION

RNA polymerase unwinds, exposing 10-20 base pairs. Uses template strand to add complementary RNA nucleotides, from 5' to 3'

#### 3) TERMINATION

**Prokaryotes:** protein, mRNA binds to itself (hairpin) **Eukaryotes:** many A's = many U's added = weak=proteins bind

## **Multiple Transcription Machinery**

multiple RNAP can transcribe simultaneously on the same gene

# **Post-Transcriptional Modifications**

in prokaryotes mRNA can be used directly, in eukaryotes it needs to be modified (pre-mRNA to mature mRNA) in order to leave the nucleus

# mRNA Modifications- capping

**poly(A)** tail - 50-250 adenine added to 3' end, preventing degredation ----- 5' cap 7 G's added to prevent degradation, signals for ribosomes to attach

#### mRNA Modifications- splicing

removal of introns (non-coding regions) & mature mRNA will only contain exons. Occurs in spliceosome (snRNPs bind to splice sites, excises introns, rejoins exons)



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