

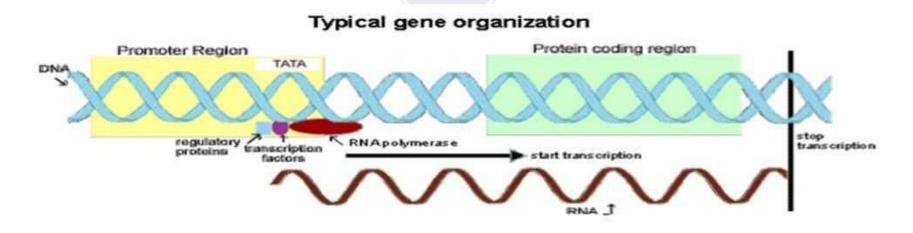
# FACULTY OF ENGINEERING AND TECHNOLOGY

**Department of Biotechnology** 

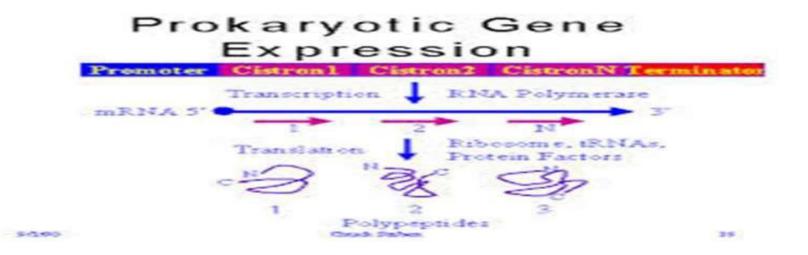
Genes are subunits of DNA, the information database of a cell that is contained inside the cell nucleus.

This DNA carries the genetic blueprint that is used to make all the proteins the cell needs.

Every gene contains a particular set of instructions that code for a specific protein



- In prokaryotes the primary control point is the process of transcription initiation.
- Different ways for regulation of gene expression in bacteria:
  - 1- Promoter recognition.
  - 2-Transcription elongation (Attenuation).
- Regulation of gene expression can be done by some operon pathways such as
  1.lac operon.
  2.tryptophan operon.

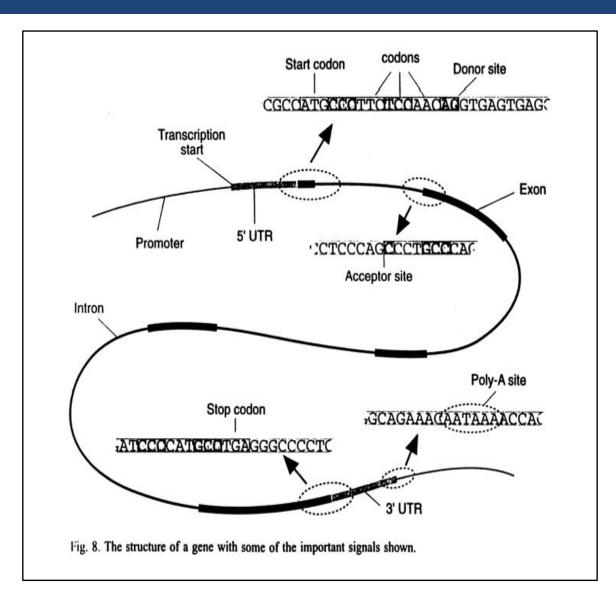


✓ Eukaryotic DNA wrapped around histones that might result in repeated patterns (periodicity of 10) for histone binding. The promotor regions might be near these sites so that they remain hidden.

✓ Prokaryotes have no introns.

✓ Promotor regions and start sites more highly conserved in Prokaryotes

✓ Different codon use frequencies



- Codon usage patterns vary by species
- Functional regions (promoters, splice sites, translation initiation sites, termination signals) vary by species
- Common repeat sequences are species- specific
- Gene finding programs rely on this information to identify coding regions

#### Pattern-based gene finding

- ORF finding based on start and stop codon frequency is a pattern-based procedure
- Other pattern-based procedures recognize characteristic sequences associated with known features and genes, such as ribosome binding sites, promoter sites, histone binding sites, etc.
- Statistically based.

#### Content-based gene finding

- Content-based gene finding methods rely on statistical information derived from known sequences to predict unknown genes
- Some evaluative measures include: "coding potential" (based on codon bias), periodicity in the sequence, sequence homogeneity, etc.

## A standard content-based alignment procedure

- Select a window of DNA sequence from the unknown. The window is usually around 100 base pairs long
- Evaluate the window's potential as a gene, based on a variety of factors
- Move the window over by one base
- Repeat procedure until end of sequence is reached; report continuous high-scoring regions as putative genes









### MCQs

- 1. A
- 2. A
- 3. A
- 4. A
- 5. A
- 6. A
- 7. A
- 8. A
- 9. A
- 10. A

