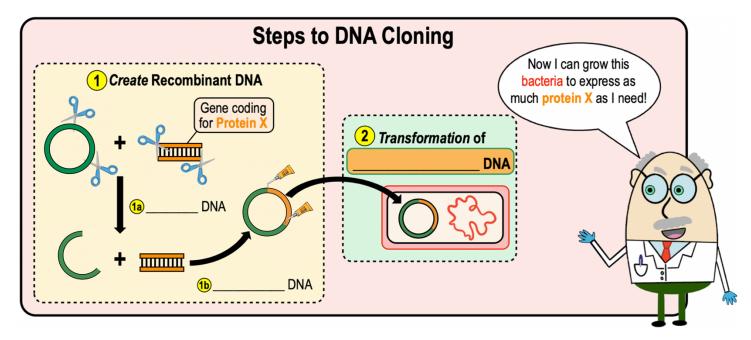
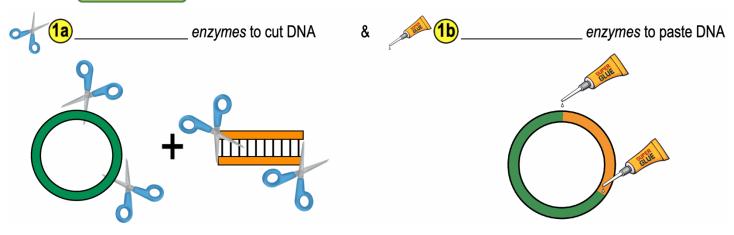
• Recall: There are _____ general steps in DNA Cloning:





Step 1) Create Recombinant DNA

• Creating a recombinant DNA molecule is a 2-step process that requires:



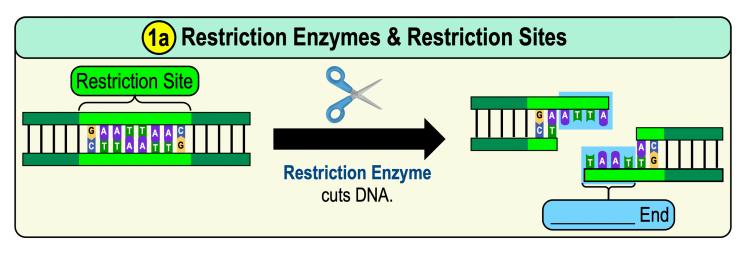
PRACTICE: Human DNA cut with restriction enzyme A can be joined to:

- a) A bacterial plasmid cut with restriction enzyme A.
- b) Human DNA cut with restriction enzyme B.
- c) Human DNA that is uncut.
- d) A bacterial plasmid that is uncut.

1a) Use Restriction Enzymes

 Enzymes: cleave DNA at restriction sites producing "sticky ends". 	
Restriction	specific sequence of DNA where restriction enzymes bind & cut the DNA.
□ Sticky ends: a	-stranded DNA "overhang" produced from a restriction digestion reaction.

EXAMPLE: Restriction enzymes cut DNA at the restriction site leaving a sticky end overhang.



PRACTICE: What is the enzymatic function of restriction enzymes?

- a) To add new nucleotides to the growing strand of DNA.
- b) To join nucleotides during replication.
- c) To join sticky ends of DNA.
- d) To cleave DNA at specific sequences.

PRACTICE: Each restriction enzyme has a specific sequence of nucleotides where it cuts the DNA. These sequences of DNA are unique to each restriction enzyme and are known as:

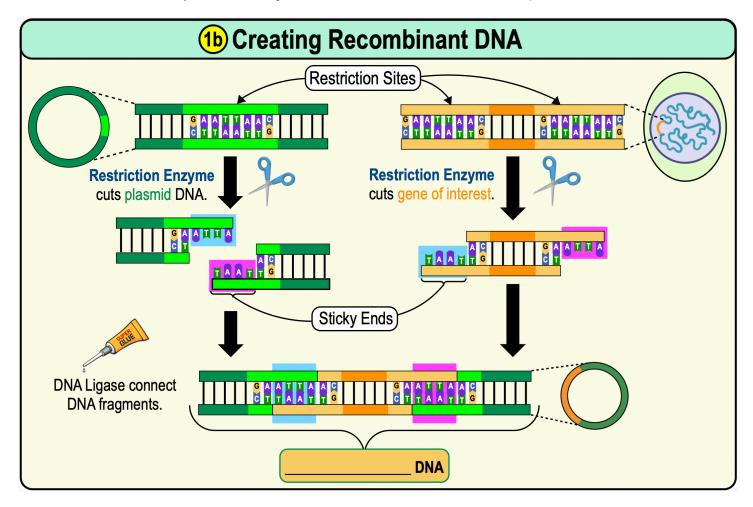
- a) Restriction Sites.
- b) Restriction Signals.
- c) Restriction Sequences.
- d) Restriction Sticky Ends.

1b) Use Ligation Enzymes

•DNA _____: enzyme that *ligates* (or joins) the two *sticky ends* together creating the recombinant DNA.

□ Note: only DNA fragments cut by the ______ restriction enzyme can be ligated back together.

EXAMPLE: A restriction enzyme and DNA ligase are used to clone a recombinant DNA plasmid.



PRACTICE: The single-stranded ends of DNA molecules can be joined together by:

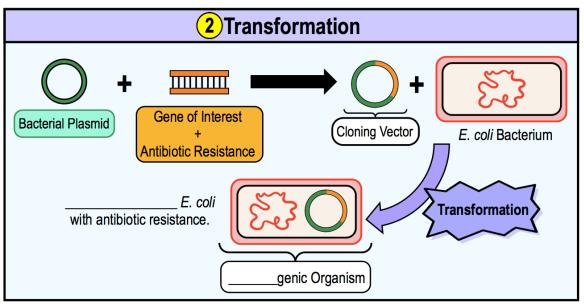
- a) Restriction Endonucleases.
- b) DNA ligase.
- c) DNA polymerase.
- d) Helicase.
- e) Primase.

2) Transform Recombinant DNA into Bacteria

- ●The second and final step of DNA cloning is to ______ the recombinant DNA.

 □ Transformation: process allowing cells to directly uptake foreign DNA (ex. cloning vector).
- _____ Organism: organism that receives & expresses recombinant DNA.
- Phenotypic markers (ex. _____ resistance) are used to confirm a positive transformation.

EXAMPLE: Creating a transgenic organism with antibiotic resistance by transformation of recombinant plasmid DNA.



• Transformed bacterial cells *replicate* & *express* the gene of interest, which the researcher can then purify & study.

PRACTICE: The process of using DNA from one organism to alter the characteristics of another is called:

- a) Translation.
- b) Transduction.
- c) Transcription.
- d) Transformation.

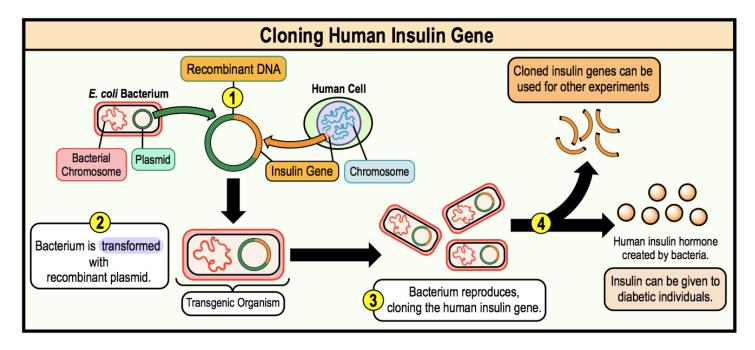
PRACTICE: An organism which has foreign genes incorporated into its genomes is known as a:

- a) Recombinant organism.
- b) Transgenic organism.
- c) Hybrid organism.
- d) Polyploid organism.

Review & Application of DNA Cloning in Medicine

- •Now that we have discussed the techniques used in DNA cloning, let's see how they are used together.
- Diabetics do not produce enough ______ protein to metabolize blood glucose & they require daily injections.
 - □ Researchers have found a way to use ______ organisms to mass produce insulin for patients.

EXAMPLE: Human Insulin Protein is expressed & purified in large amounts using transgenic *E. coli*.



PRACTICE: What is the most logical sequence of steps for splicing foreign DNA into a plasmid and inserting the plasmid into a bacterium?

- I. Transform bacteria with a recombinant DNA molecule.
- II. Cut the plasmid DNA using restriction enzymes (endonucleases).
- III. Extract plasmid DNA from bacterial cells.
- IV. Hydrogen-bond the plasmid DNA to non-plasmid DNA fragments.
- V. Use ligase to seal plasmid DNA to non-plasmid DNA.
 - a) II, III, V, IV, I
 - b) III, II, IV, V, I
 - c) III, IV, V, I, II
 - d) IV, V, I, II, III