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preparatory stage and will
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of the ring you just drew to

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represent the origin of replication. Now, using your circular paper plasmid, mark clockwise from "O" and write on your drawing the following information as it appears in your paper plasmid: (a) Show where the

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inserted gene is located on
your loop.

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demo on the process of cloning a gene into a plasmid. It reviews the concepts of restriction enzymes, "sticky ends", cloning, and transformation.

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...

LAB ____: CLONING PAPER
PLASMID In this exercise you
will use paper to simulate
the cloning of a gene from
one organism into a
bacterial plasmid using a
restriction enzyme digest.

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The plasmid (puc18 plasmid) can then be used to transform bacteria so that it now expresses a new gene and produces a new protein.

1. From the white paper, cut out the puc18 plasmid DNA in a long strip.
- 2.

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I need help with a few
questions: 1: Why did we

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make sure to include the start and stop DNA sequences for the Jellyfish Glo gene in our cut segment 2:What would have happened if we had cut both the Jellyfish Glo gene and puc18 plasmid with the EcoR1 restriction

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enzyme? Be sure to look on the paper DNA sequences to find the EcoR1 restriction enzyme cut sites.

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Recombinant Paper Plasmid
Background

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of 4 Adapted by Kim B.

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_____: CLONING PAPER PLASMID

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In this exercise you will use paper to simulate the cloning of a gene from one organism into a bacterial plasmid using a restriction enzyme digest. The plasmid (puc18 plasmid) can then be used to transform bacteria

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so that it now expresses a
new gene and ...

16 biology lab plasmid

Flashcards and Study Sets |

Quizlet

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Answers ...

LAB ____: CLONING PAPER
PLASMID In this exercise you
will use paper to simulate
the cloning of a gene from
one organism into a
bacterial plasmid using a
restriction enzyme digest.

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Answers Key

The plasmid (puc18 plasmid) can then be used to transform bacteria so that it now expresses a new gene and produces a new protein.

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Biology ...

The E. coli Insulin Factory

BACKGROUND Bacteria have not only their normal DNA, they also have pieces of circular DNA called plasmids.

Plasmids are a wonderfully ally for biologists who

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desire to get bacteria to produce very specific proteins. The plasmids conveniently can be cut, fused with other DNA and then reabsorbed by bacteria.

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Isolation of DNA Fragments
from Complex Mixtures A DNA
fragment of a few base pairs
up to 20 kb can be inserted
into a plasmid vector . When

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such a recombinant plasmid transforms an E. coli cell, all the antibiotic-resistant progeny cells that arise from the initial transformed cell will contain plasmids with the same inserted sequence of DNA (Figure 7-3

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