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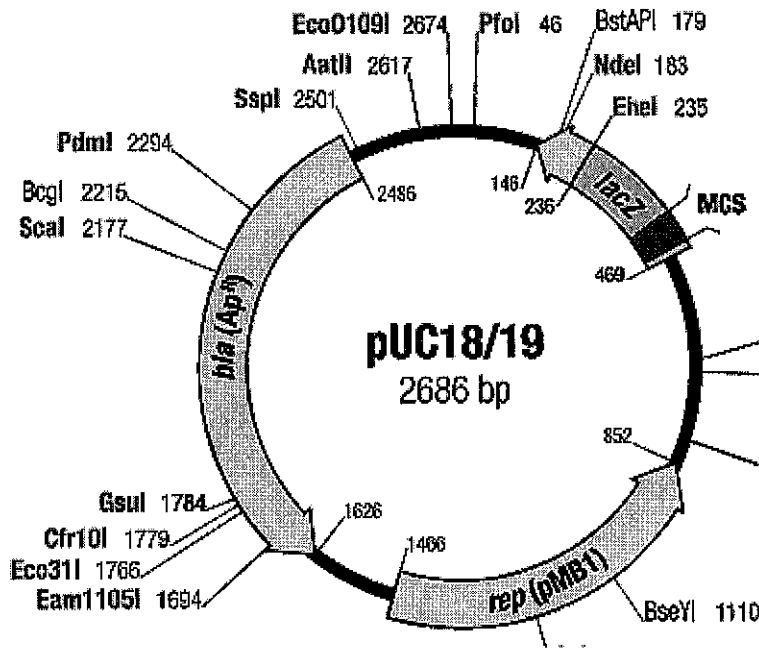
COURSE CODE: 10B11BT513  
 COURSE NAME: GENETIC ENGINEERING

MAX. MARKS: 50  
 MAX. TIME: 2 HR

Q1. You are provided with diagram of a pUC18/19 vector. The gene for ampicillin resistance (*bla*) was amplified using primers designed from the start and end codon of the gene. [15]

**Provide the following:**

- Size of the PCR product obtained. [2]
- Number and size of products obtained when the amplified product is digested with **ScaI**. [2]
- Number and size of products obtained when the amplified product is digested with **ScaI** and **SspI**. [2]
- Number and size of products obtained when the amplified product is digested with **ScaI** and **GsuI**. [2]
- Draw a well labeled agarose gel showing different bands obtained when the PCR product/digested product(s) would be run from a, b, c, and d. [4]
- Utility of '*bla*' gene for cloning of insert using the vector. [3]



Q2. How are the following useful in selection of 'insert containing vector' in genetic engineering? [5 X 3 = 15]

- Blue and white screening
- Selection marker
- Reporter genes

Q3. What are restriction enzymes? Differentiate between different types of restriction enzymes with examples and their application in Genetic Engineering. [5]

Q4. Write Short Notes on: [3 X 3 = 9]

- T4 DNA ligase
- T4 Polynucleotide Kinase
- Alkaline phosphatase

Q5. Illustrate the life cycle of M13 filamentous phage and discuss its applications in Genetic Engineering. [6]