$\mathbf{R05}$ 



# IV B.Tech I Semester Examinations, November 2010 METABOLIC ENGINEERING Bio-Technology

Time: 3 hours

Code No: R05412310

Max Marks: 80

## Answer any FIVE Questions All Questions carry equal marks $\star \star \star \star \star$

1.	Explain briefly how radiolabel materials are utilized in experimental determination of metabolic flux. [16]
2.	What is Jacob Monod model? Explain its regulation with reference to Lac operon. [16]
3.	Explain the advantages of writing an algorithm for a metabolic pathway synthesis, explain with an example. [16]
4.	What is substrate range extension? Explain pathway manipulations to substrate range extension in the production of ethanol production [16]
5.	Write about the alteration of feed back regulation.[16]
6.	Discuss in detail the conversion of insoluble substances by mixed or sequential bioconversions. [8+8]
7.	Write about various producers of secondary metabolites. [16]
8.	Explain the goals of biotechnological improvements in crops? Explain various strategies for metabolic engineering in plants. [16]

**R05** 

Set No. 4

## **IV B.Tech I Semester Examinations, November 2010** METABOLIC ENGINEERING **Bio-Technology**

Time: 3 hours

Code No: R05412310

Max Marks: 80

[16]

[16]

### Answer any FIVE Questions All Questions carry equal marks \*\*\*\*

- 1. Write about various producers of secondary metabolites.
- 2. Explain briefly how radiolabel materials are utilized in experimental determination of metabolic flux. 16
- 3. What is Jacob Monod model? Explain its regulation with reference to Lac operon.
- 4. Explain the advantages of writing an algorithm for a metabolic pathway synthesis, explain with an example. [16]
- 5. Explain the goals of biotechnological improvements in crops? Explain various strategies for metabolic engineering in plants. [16]
- 6. What is substrate range extension? Explain pathway manipulations to substrate range extension in the production of ethanol production. [16]
- 7. Discuss in detail the conversion of insoluble substances by mixed or sequential bioconversions. |8+8|
- 8. Write about the alteration of feed back regulation. [16]

 $\mathbf{R05}$ 

# Set No. 1

# IV B.Tech I Semester Examinations, November 2010 METABOLIC ENGINEERING Bio-Technology

Time: 3 hours

Code No: R05412310

Max Marks: 80

## Answer any FIVE Questions All Questions carry equal marks \* \* \* \* \*

1.	What is substrate range extension? Explain pathway manipulations to substrate range extension in the production of ethanol production. [16]
2.	What is Jacob Monod model? Explain its regulation with reference to Lac operon. [16]
3.	Write about various producers of secondary metabolites. [16]
4.	Discuss in detail the conversion of insoluble substances by mixed or sequential bioconversions. [8+8]
5.	Write about the alteration of feed back regulation. [16]
6.	Explain briefly how radiolabel materials are utilized in experimental determination of metabolic flux. [16]
7.	Explain the goals of biotechnological improvements in crops? Explain various strategies for metabolic engineering in plants. [16]
8.	Explain the advantages of writing an algorithm for a metabolic pathway synthesis, explain with an example. [16]

 $\mathbf{R05}$ 

Set No. 3

# IV B.Tech I Semester Examinations,November 2010 METABOLIC ENGINEERING Bio-Technology

Time: 3 hours

Code No: R05412310

Max Marks: 80

[16]

[16]

## Answer any FIVE Questions All Questions carry equal marks \* \* \* \* \*

1. Explain the goals of biotechnological improvements in crops? Explain various strategies for metabolic engineering in plants. [16]

2. Write about various producers of secondary metabolites.

- 3. Write about the alteration of feed back regulation.
- 4. Explain the advantages of writing an algorithm for a metabolic pathway synthesis, explain with an example. [16]
- 5. Discuss in detail the conversion of insoluble substances by mixed or sequential bioconversions. [8+8]
- 6. What is Jacob Monod model? Explain its regulation with reference to Lac operon.
  [16]
- 7. What is substrate range extension? Explain pathway manipulations to substrate range extension in the production of ethanol production. [16]
- 8. Explain briefly how radiolabel materials are utilized in experimental determination of metabolic flux. [16]