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## Unit 11 - Week 9

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### Course outline

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Week 9

Lecture 38 : Isolation and Characterization of DNA Part - I

Lecture 39 : Isolation and Characterization of DNA Part - II

Lecture 40 : Bacterial Culture for

## Assignment 9

The due date for submitting this assignment has passed.

As per our records you have not submitted this assignment. **Due on 2019-04-03, 23:59 IST.**

1) To lyse the bacterial cell wall for extraction of DNA, which of the following reagents is commonly used? **1 point**

- a. Lysozyme
- b. Phenol
- c. Penicillin
- d. Cetrimonium bromide

No, the answer is incorrect.

Score: 0

Accepted Answers:

a. Lysozyme

2) Isolation of DNA from plant sources possess challenges due to the presence of **1 point**

- a. high amount of RNA
- b. high amount of protein
- c. presence of polysaccharide and secondary metabolites
- d. low amount of DNA

No, the answer is incorrect.

Score: 0

Accepted Answers:

c. presence of polysaccharide and secondary metabolites

3) The function of EDTA during cell lysis is to **1 point**

- a. act as metal chelating agent
- b. inhibit cellular enzymes that degrade DNA
- c. destabilize the cell wall

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Characterization  
of DNA  
Summary

Quiz :  
Assignment 9

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Solution

- a. Denature genomic and plasmid DNA.
- b. Break down the cell wall.
- c. Degrades the RNA.
- d. Remove proteins from the cell lysate.

**No, the answer is incorrect.**

**Score: 0**

**Accepted Answers:**

- a. Denature genomic and plasmid DNA.*
- b. Break down the cell wall.*

5) RNase A degrades RNA that are present in cell lysate. What will happen if RNase A treatment is not done during DNA isolation? **1 point**

- a. DNA will get degraded.
- b. RNA will not be removed and as both DNA and RNA absorbs at 260nm, quantification of isolated pure DNA will be problematic.
- c. Both DNA and RNA will form complex and precipitate out of the solution.
- d. All of the above.

**No, the answer is incorrect.**

**Score: 0**

**Accepted Answers:**

- b. RNA will not be removed and as both DNA and RNA absorbs at 260nm, quantification of isolated pure DNA will be problematic.*

6) What is the role of SDS in lysis buffer for plasmid DNA isolation? **1 point**

- a. Solubilizes the membrane.
- b. Denatures the proteins.
- c. Disrupts DNA-Protein complexes.
- d. All of the above.

**No, the answer is incorrect.**

**Score: 0**

**Accepted Answers:**

- d. All of the above.*

7) What is the purpose of plasmid isolation? **1 point**

- a. Cloning
- b. Gene expression
- c. Vehicle for delivering target DNA in another organism
- d. All of the above

**No, the answer is incorrect.**

**Score: 0**

**Accepted Answers:**

- d. All of the above*

8) Isolating plasmid DNA from genomic DNA is based upon **1 point**

- a. conformation
- b. charge difference
- c. size difference
- d. cellular location

No, the answer is incorrect.

Score: 0

Accepted Answers:

a. conformation

c. size difference

9) The neutralizing buffer used in plasmid isolation contains

1 point

- a. triton-X
- b. potassium/sodium acetate
- c. SDS
- d. phenol



No, the answer is incorrect.

Score: 0

Accepted Answers:

b. potassium/sodium acetate

10) When neutralizing buffer is added after lysis to the cell lysate

1 point

- a. Chromosomal DNA gets tangled and precipitates out.
- b. Plasmid DNA anneals and remains in the solution.
- c. Proteins and SDS forms insoluble precipitate.
- d. All of the above.

No, the answer is incorrect.

Score: 0

Accepted Answers:

d. All of the above.

11) DNA bound to silica membrane can be eluted out by addition of

1 point

- a. Water
- b. Tris pH 6.0
- c. Isopropanol
- d. 0.5% SDS solution

No, the answer is incorrect.

Score: 0

Accepted Answers:

a. Water

12) To wash the DNA bound to silica membrane which of the following reagents are used?

1 point

- a. Tris pH 8.0
- b. Deionized water
- c. Ethanol/Isopropanol
- d. Detergents

No, the answer is incorrect.

Score: 0

Accepted Answers:

c. Ethanol/Isopropanol

13) Quality of the purified plasmid DNA has to be checked by

1 point

- a. Size exclusion chromatography

- b. Agarose gel electrophoresis
- c. Polyacrylamide gel electrophoresis
- d. UV spectroscopy

**No, the answer is incorrect.**

**Score: 0**

**Accepted Answers:**

*b. Agarose gel electrophoresis*

*d. UV spectroscopy*



14) The quantity of the purified DNA can be estimated by measuring absorbance at

**1 point**

- a. 260nm
- b. 280nm
- c. 560nm
- d. 190nm



**No, the answer is incorrect.**

**Score: 0**

**Accepted Answers:**

*a. 260nm*

15) In agarose gel electrophoresis, the plasmid DNA runs ahead of linear DNA as

**1 point**

- a. Linear DNA is more compact
- b. Plasmids have higher mass/charge ratio
- c. Plasmids are supercoiled and migrates faster
- d. All the above

**No, the answer is incorrect.**

**Score: 0**

**Accepted Answers:**

*c. Plasmids are supercoiled and migrates faster*

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