

### NPTEIN

reviewer2@nptel.iitm.ac.in ▼

#### Courses » Mass spectrometry based proteomics



**Announcements** 

Course

Ask a Question

**Progress** 

## Unit 4 - Week 3: Quantitative proteomics



# Course

How to access the portal

Week 1: Proteomics introduction and sample preparation

Week 2: Basics of mass spectrometry

#### Week 3: Quantitative proteomics

- Lecture 11: Introduction to quantitative proteomics
- Lecture 12:
   Hybrid mass spectrometry configurations
- Lecture 13:SILAC: In Vivo labeling
- Lecture 14: iTRAQ: In Vitro labeling
- Lecture 15:TMT: In vitro labeling
- Lab session3.1: Dataanalysis
- Lab session
   3.2: Targeted proteomics-experimental design
- Assignment-III Answer key

## **Assignment-III**

The due date for submitting this assignment has passed. Due on 2016-04-12, 05:20 IS As per our records you have not submitted this assignment.

- 1) You want to treat a human cell line with drug Imatinib and analyze its proteomic **0.5 points** alterations using SILAC. Which of the following radio-labeled amino acid could be used for SILAC experiment?
  - Leucine
  - Glycine
  - Proline
  - Lysine

No, the answer is incorrect.

Score: 0

**Accepted Answers:** 

Lysine

- 2) You want to perform an in vitro labeling method for quantitative proteomics experiment. *0.5 points* Which of the following method could NOT be preferred for in vitro labeling?
  - Quest
  - CDIT
  - GIST
  - ICAT

No, the answer is incorrect.

Score: 0

**Accepted Answers:** 

**CDIT** 

- 3) In an iTRAQ experiment, the quantification of peptides could be performed by measuring **0.5 points** intensity of reporter ion in MS/MS spectra. Which of the following chemical group is reporter ion in iTRAQ?
  - N, N-dimethyl peperazine
  - N, N-peperazine
  - N, N- methyl peperazine
  - N-methyl peperazine

No, the answer is incorrect.

Score: 0

**Accepted Answers:** 

N, N-dimethyl peperazine

Quiz : Assignment-III

Week-4: Proteomics and systems biology

	teolysis using trypsin, peptides are labeled with iTRAQ for quantitative prote- ich of the following amino acids could be involved in iTRAQ labeling?	omic <i>0.5 point</i> s
Onl	y lysine	
	ine and N-terminal amino acid	
Bot	h lysine and Arginine	
Onl	y Arginine	
No, the a	nswer is incorrect.	-
•	Answers: d N-terminal amino acid	<u> </u>
	Q based quantitative analysis, we always rely on MS/MS spectral data rather nat is the information that we could obtain from MS/MS spectrum during iTRA	-
Pro	vide amino acid sequence information	
	vide the quantitative information	I
Bot	n A & B	
O Pro	vide the sample injection to ensure ionization	[2
	nswer is incorrect.	
Score: 0		
Accepted Both A &	l <b>Answers:</b> B	
proteome an	t to grow bacterial sample in a media having N15 heavy labels for quantitativalysis. Though metabolic labeling of the proteins in bacteria is considered to be but still it has some disadvantages. What is the drawback of N15 labeling?	be highly
Qua	antification using MS	
O In v	ivo labeling for quantitation	
○ Use	d for microorganism	
O Une	equal incorporation of isotopes in protein	
	nswer is incorrect.	
Score: 0		
Accepted	Answers: ncorporation of isotopes in protein	
Accepted Unequal in 7) Match th		0.5 points
Accepted Unequal in 7) Match the	ncorporation of isotopes in protein e reporter ion to the respective balancer ion in iTRAQ labeling method i. 184	0.5 points
Accepted Unequal in 7) Match the a. 113 b. 114	ncorporation of isotopes in protein e reporter ion to the respective balancer ion in iTRAQ labeling method i. 184 ii. 192	0.5 points
Accepted Unequal in 7) Match th a. 113 b. 114 c. 115	ncorporation of isotopes in protein re reporter ion to the respective balancer ion in iTRAQ labeling method i. 184 ii. 192 iii. 31	0.5 points
Accepted Unequal in 7) Match that 113 or 114 or 115 dr.116	ncorporation of isotopes in protein e reporter ion to the respective balancer ion in iTRAQ labeling method i. 184 ii. 192	0.5 point
Accepted Unequal in 7) Match the a. 113 b. 114 c. 115 d.116 e. 121	ncorporation of isotopes in protein e reporter ion to the respective balancer ion in iTRAQ labeling method i. 184 ii. 192 iii. 31 iv. 30	0.5 point
Accepted Unequal in 7) Match the a. 113 b. 114 c. 115 d.116 e. 121	ncorporation of isotopes in protein e reporter ion to the respective balancer ion in iTRAQ labeling method i. 184 ii. 192 iii. 31 iv. 30 v. 29	0.5 point
Accepted Unequal in 7) Match the a. 113 b. 114 c. 115 d.116 e. 121	ncorporation of isotopes in protein  e reporter ion to the respective balancer ion in iTRAQ labeling method  i. 184  ii. 192  iii. 31  iv. 30  v. 29  , b-l, c-iv, d-v, e-iii	0.5 point
Accepted Unequal in 7) Match the a. 113 b. 114 c. 115 d.116 e. 121 a-ii, a-ii, a-ii,	ncorporation of isotopes in protein  e reporter ion to the respective balancer ion in iTRAQ labeling method  i. 184  ii. 192  iii. 31  iv. 30  v. 29  , b-l, c-iv, d-v, e-iii  b-iii, c-iv, d-v, e-i	0.5 point
Accepted Unequal in 7) Match the a. 113 b. 114 c. 115 d.116 e. 121 a-ii, a-ii, a-iii a-iii	ncorporation of isotopes in protein  e reporter ion to the respective balancer ion in iTRAQ labeling method  i. 184  ii. 192  iii. 31  iv. 30  v. 29  , b-I, c-iv, d-v, e-iii  b-iii, c-iv, d-v, e-i  b-iii, c-I, d-v, e-iv	0.5 point
Accepted Unequal in 7) Match the a. 113 b. 114 c. 115 d.116 e. 121 a-ii, a-ii, a-iii a-iii	ncorporation of isotopes in protein  e reporter ion to the respective balancer ion in iTRAQ labeling method  i. 184  ii. 192  iii. 31  iv. 30  v. 29  , b-l, c-iv, d-v, e-iii  b-iii, c-lv, d-v, e-i  b-iii, c-l, d-v, e-iv  , b-ii, c-iv, d-v, e-i	0.5 point
Accepted Unequal is 7) Match the a. 113 b. 114 c. 115 d.116 e. 121  a-ii, a-iii  No, the as Score: 0 Accepted	ncorporation of isotopes in protein  e reporter ion to the respective balancer ion in iTRAQ labeling method  i. 184  ii. 192  iii. 31  iv. 30  v. 29  , b-l, c-iv, d-v, e-iii  b-iii, c-lv, d-v, e-i  b-iii, c-l, d-v, e-iv  , b-ii, c-iv, d-v, e-i	0.5 point
Accepted Unequal is 7) Match the a. 113 b. 114 c. 115 d.116 e. 121  a-ii, a-iii No, the as Score: 0 Accepted a-ii, b-iii, co	ncorporation of isotopes in protein  e reporter ion to the respective balancer ion in iTRAQ labeling method  i. 184  ii. 192  iii. 31  iv. 30  v. 29  b-I, c-iv, d-v, e-iii b-iii, c-I, d-v, e-iv b-iii, c-I, d-v, e-iv h-iii, c-iv, d-v, e-i  mswer is incorrect.	0.5 points
Accepted Unequal in 7) Match the a. 113 b. 114 c. 115 d.116 e. 121 a-ii, a-iii, a-iiii No, the as Score: 0  Accepted a-ii, b-iii, c. 8) Which on	ncorporation of isotopes in protein  e reporter ion to the respective balancer ion in iTRAQ labeling method  i. 184  ii. 192  iii. 31  iv. 30  v. 29  , b-I, c-iv, d-v, e-iii  b-iii, c-iv, d-v, e-i  b-iii, c-l, d-v, e-iv  , b-ii, c-iv, d-v, e-i  nswer is incorrect.  Answers:  e-iv, d-v, e-i  ne of the following statements is not correct with regards to SILAC?	
Accepted Unequal is 7) Match the a. 113 b. 114 c. 115 d.116 e. 121  a-ii, a-iii No, the as Score: 0 Accepted a-ii, b-iii, co 8) Which o	ncorporation of isotopes in protein  e reporter ion to the respective balancer ion in iTRAQ labeling method  i. 184  ii. 192  iii. 31  iv. 30  v. 29  , b-I, c-iv, d-v, e-iii  b-iii, c-iv, d-v, e-i  b-iii, c-l, d-v, e-iv  , b-ii, c-iv, d-v, e-i  nswer is incorrect.  I Answers:  e-iv, d-v, e-i  ne of the following statements is not correct with regards to SILAC?	
Accepted Unequal in 7) Match the a. 113 b. 114 c. 115 d.116 e. 121 a-ii, a-iii, a-iiii No, the as Score: 0  Accepted a-ii, b-iii, c. 8) Which o	ncorporation of isotopes in protein  e reporter ion to the respective balancer ion in iTRAQ labeling method  i. 184  ii. 192  iii. 31  iv. 30  v. 29  , b-I, c-iv, d-v, e-iii  b-iii, c-iv, d-v, e-i  b-iii, c-l, d-v, e-iv  , b-ii, c-iv, d-v, e-i  nswer is incorrect.  Answers:  e-iv, d-v, e-i  ne of the following statements is not correct with regards to SILAC?	0.5 points

Mass spectrometry based proteomics Unit 4 - Week 3: Quantitative proteomics	
<ul> <li>SILAC can be used for tissue and body fluid samples</li> </ul>	
No, the answer is incorrect. Score: 0	
Accepted Answers: SILAC can be used for tissue and body fluid samples	
9) In Tandem Mass spectrometry, the peptides are resolved using two mass analyzers, <b>0.5 pc</b> separated by collision cell. What is the function of mass analyzer-I, collision cell and mass analyzer-I	12
<ul> <li>Select fragment, fragmentation, select parent</li> <li>Select parent, fragmentation, scans fragment ions</li> <li>Select parent ion, select fragment ion, fragmentation</li> <li>Fragmentation, parent ion selection, fragment ion selection</li> </ul>	f
No, the answer is incorrect. Score: 0	in
Accepted Answers: Select parent, fragmentation, scans fragment ions	in σ+
10 Different types of mass analyzers are available to resolve the ions based on its respective <b>0.5 pc</b> principle. Which of the following mass analyzer has highest resolution?	)ints
Triple Quadrupole Fourier Transformer Fourier Transformer Ion Cyclotron Resonance Electron-positron cyclotron collider	
No, the answer is incorrect. Score: 0	
Accepted Answers: Fourier Transformer Ion Cyclotron Resonance	
11)Many protein labeling methods are available for the mass spectrometry based protein quantitation. Which of the following labeling method, which is also first gel-free quantitative labeling method, depends on cystine residue for labeling of protein?	oints
<ul><li>iTRAQ</li><li>TMT</li><li>iCAT</li><li>SILAC</li></ul>	
No, the answer is incorrect. Score: 0	
Accepted Answers: iCAT	
12 During the in vitro labeling, tagging of the protein is either by N-terminal, C-terminal or amino acid based. Which of the following method is an amino acid based labeling method?	oints
<ul><li>iTRAQ</li><li>Esterification</li><li>MCAT</li><li>GIST</li></ul>	
No, the answer is incorrect. Score: 0	
Accepted Answers: MCAT	
13)Match the following labeling method with its respective statement  a. VICAT  i. In vivo labeling  b. TMT  ii. Amino acid based in vitro labeling  c. Proteolysis O16/O18  iii. N-terminal peptide in vitro labeling	oints

d. CDIT	iv. C-terminal peptide in vitro labeling	
a-iv, b-		
a-i, b-ii		
a-iii, b-		
○ a-ii, b-i	iii, c-iv, d-i	
No, the ansv	wer is incorrect.	
Accepted And a-ii, b-iii, c-iv,		
14)Which of th	e following is the proteomic data management database?	0.5 p
PRIDE		
Ensem	ldl	
○ HapMa	ар	
O None o	of the above	
No, the ansv Score: 0	wer is incorrect.	
Accepted A	nswers:	
	study effects of Doxorubicin on cancer patients post chemotherapy using  . Which one of the following labeling methods could be used for quantification	of ser
O N15 La	abeling	
C13 La	abeling	
itraq	Labeling	
All of the	he above	
No, the ansv Score: 0	wer is incorrect.	
Accepted A		
*	een given a peptide digest to run on MALDI-TOF. How will you be confident mass fingerprinting data is accurate?	0.5 p
By con	sidering threshold score	
O By con	sidering the peptide number	
Seque	nce coverage	
O All of the	he above	
No, the answ Score: 0	wer is incorrect.	
Accepted A		
17Key feature following?	of Selective Reaction Monitoring (SRM) is best described by which of the	0.5 p
Full fra	igment ion spectrum of each precursor in target list is recorded continuously	
	ively monitors desired transitions	
	re quantitation of samples	
	ed sample monitoring to obtain transitions	

### Selectively monitors desired transitions

Score: 0

No, the answer is incorrect.

**Accepted Answers:** 

 $https://online courses-archive.nptel.ac.in/noc16\_bt03/unit?unit=37\&assessment=74$ 

Mass spectrometry based proteomics - - Unit 4 - Week 3: Quantitative proteomics 18)You have run two peptides (X and Y) on LC MS/MS instrument and obtained same MS 0.5 points spectra for both peptides. How will you distinguish these two peptides? By considering score By considering the quantity of the protein By acquiring MS/MS data All of the above No, the answer is incorrect. Score: 0 **Accepted Answers:** By acquiring MS/MS data 19)You have a complex protein mixture from serum and you want to quantify the protein of interest. Which of the following techniques will fit best? Shotgun proteomics MALDI -TOF/TOF analysis Quantitative analysis using TMT Multiple reaction monitoring No, the answer is incorrect. Score: 0 **Accepted Answers:** Multiple reaction monitoring 20) There are many vendor specific software to build and run the MRM assays. Which of the 0.5 points following is an open source software widely used for MRM assays to generate peptide transition list? Multiquant Skyline Lab Solutions Proteome Discoverer No, the answer is incorrect. Score: 0 **Accepted Answers:** Skyline

Previous Page

End

© 2014 NPTEL - Privacy & Terms - Honor Code - FAQs -





Funded by

Government of India Ministry of Human Resource Development

Powered by











