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Working with Protein: Proteomics and Purification

What is the common range for polypeptide lengths?

1 message

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Thanks for filling out Working with Protein: Proteomics and Purification

Here's what we got from you:

Working with Protein: Proteomics and Purification

0 10 - 1000 amino acids
100 - 1000 amino acids
50 - 500 amino acids
Other:
Multisubunit proteins are made up of two or more polypeptide chains that are identical.
O True
False
What is the minimum size of a polypeptide chain that commonly carries out a particular function?
40 amino acids

Why are proteins greater than 1000 amino acids not very common?

Because it takes a lot of effort and resources from a cell to ensure that the polypeptide chain gets synthesized properly and folds into the correct structure.

William Diliversity Wall - Working Will Frotein. Frotesines and Furnication	
What four amino acids are the rarest in proteins?	
Histidine, Tryptophan, Cysteine, Methionine	
What five variable need to be considered and controlled when a bioch protein stable?	emist wants to keep a
pH, Temperature, presence of enzymes that can degrade or damage the protein, surface adsorption potential, long term storage	
What does ELISA stand for and what is it used for?	
enzyme linked immunosorbent assay	
Trp and Tyr. NOT Phe! Describe the relationship between salting out and salting in. How are they different?	they similar and how are
Really low salt concentrations will cause proteins to be insoluble. Really high salt concentrations will cause proteins to be insoluble. So as you increase the concentration of salt, you encourage solubility - this is salting in. As you keep increasing the salt, the protein precipitates - this is salting out.	
What does HPLC stand for?	
High pressure liquid chromatography	
DAEA is a common functional group used in exchange ch	romatography.

Why is the isoelectric point of a protein important to consider when doing ion exchange chromatography?

It dictates the charge on a protein at a certain pH value.

cation

anion

polyvalent

size	
What does PAGE stand for?	
polyacrylamide gel electrophoresis	
Why is SDS commonly added to PAGE experiments?	
it denatures the protein and coats it with negative charge giving each protein has a uniform charge and has "snake-like" properties. This ensures that the only variable that influences protein migration is a PAGE experiment is the size.	
What can 2D electrophoresis do that 1D cannot?	
separate proteins with similar masses or pl values. 1D can only separate based on one variable, but it is very possible (in fact, very likely) that there are multiple proteins in a cell that have the same MW or pl.	
	1
Which protease cleaves adjacent to large hydrophobic residues?	
trypsin	
chymotrypsin	
elastase	
thermolysin	
pepsin	
Mass spectrometry has emerged as an important tool to sequence pro on a collision cell. What does the collision cell do?	teins. This technique relies
fragment one peptide bond in a peptide chain.	
	1
When comparing two protein sequences, what is meant by an invariar	nt residue?
This amino acid exists in all similar proteins at the same position. It strongly suggests that this amino acid is required for structure or	

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