

focal adhesion kinase (FAK)

Übung 10

Vorlesung Bio-Engineering

Frühlingssemester 2019

Foundations o	f Cel	I-Matrix	Mec	hano	bio	logy
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I. TRUE/FALSE Decide whether each of these statements is true or false, and then explain why.

Q1: The extracellular matrix is a relatively inert scaffolding that stabilizes the structure of tissues.		
Q2: One of the main chemical differences between proteoglycans and other glycoproteins lies in the structure of their carbohydrate side chains: proteoglycans mostly contain long, unbranched polysaccharide side chains, whereas other glycoproteins contain much shorter, highly branched oligosaccharides.		
II. CALCULATIONS		
Q3: Platelets are flat, disc-like cells about 2 μ m in diameter. Estimates of the number of integrin molecules on their surface vary around a mean of about 80,000. If the integrins themselves are about 10 nm in diameter, how tightly packed are they? (Assume that the total membrane area is $2\pi r^2$.)		
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III. DEFINITIONS Define or briefly explain	the following terms:	
Q4: Terms to be defined	A4: Definitions	
anchorage dependent cell behaviors		
An integrin and its ligand		



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IV. MIX AND MATCH Select the best matched term from the list to the provided description in the table.

TERMS TO LEARN: collagen; glycosaminoglycan (GAG); collagen fibril; hyaluronan; elastin; matrix metalloprotease; proteoglycan; fibril-associated collagen; RGD sequence; fibrillar collagen; fibronectin; extracellular matrix; chondroblast; fibroblast; osteoblast; osteoclast; osteocyte;

A5: Best suited	Q5: Description:
term from the list:	
	Fibrous protein rich in glycine and proline that, in its many forms, is a major
	component of the extracellular matrix and connective tissues.
	Complex network of polysaccharides (such as glycosaminoglycans or cellulose) and
	proteins (such as collagens) secreted by cells that serves as a structural element in
	tissues and also influences tissue development and physiology.
	General name for long, linear, highly charged polysaccharides composed of a
	repeating pair of sugars, one of which is always an amino sugar, that is found
	covalently linked to a protein core in the extracellular matrix.
	Type of collagen molecule that assembles into ropelike structures and larger,
	cablelike bundles.
	Extracellular matrix protein that binds to cell-surface integrins to promote adhesion
	of cells to the matrix and to provide guidance to migrating cells during
	embryogenesis.
	Hydrophobic protein that forms extracellular extensible fibers that give tissues their
	stretchability and resilience.
	Common cell type in connective tissue that secretes an extracellular matrix rich in
	collagen and other extracellular matrix macromolecules.

V. Experimental Design and Analysis:

Binding of fragments and competition for binding can be used to identify the portion of a larger ligand that is critical for binding. Fibronectin, which is a large glycoprotein component of the extracellular matrix, binds to fibronectin receptors on cell surfaces. Fibronectin can stick cells to the surface of a plastic dish, to which they would otherwise not bind, forming the basis of a simple binding assay. By attaching small fragments of fibronectin to dishes, researchers identified the cell-binding domain as a 108-amino acid segment about three-quarters of the way from the N-terminus.

Synthetic peptides corresponding to different portions of the 108-amino acid segment were tested in the cell-binding assay to precisely localize the active binding region. Two experiments were conducted.



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Experiment 1: first, peptides were linked covalently to plastic dishes via a disulfide bond to an attached carrier protein, and then tested for their ability to promote cell sticking (Table E1).

Experiment 2: Plastic dishes were coated with native fibronectin, and cells that stuck to the dishes in the presence of the synthetic peptides were counted (Table E2).

Table E1. Fibronectin-related peptides tested for their ability to promote cell sticking

PEPTIDE	SEQUENCE	CONCENTRATION REQUIRED FOR 50% CELL ATTACHMENT (nM)
Fibronectin		0.10
Peptide 1	YAVTGRGDSPASSKPISINYRTEIDKPSQM(C)*	0.25
Peptide 2	VTGRGDSPASSKPI(C)	1.6
Peptide 3	SINYRTEIDKPSQM(C)	>100
Peptide 4	VTGRGDSPA(C)	2.5
Peptide 5	SPASSKPIS(C)	>100
Peptide 6	VTGRGD(C)	10
Peptide 7	GRGDS(C)	3.0
Peptide 8	RGDSPA(C)	6.0
Peptide 9	RVDSPA(C)	>100

Table E2. Fibronectin-related peptides tested for their ability to block cell sticking.

PEPTIDE	PERCENT OF INPUT CELLS STICKING
GRGDSPC	2.0
GRGDAPC	1.9
GKGDSPC	48
GRADSPC	49
GRGESPC	44
None	47

Q6. The two experiments used different assays to detect the cell-binding segment of fibronectin. Does the sticking of cells to the dishes mean the same thing in both assays? Explain the difference between the assays.

Q7. From the results in Tables E1 and E2, deduce the amino acid sequence in fibronectin that is recognized by the fibronectin receptor.

^{*}The (C) at the C-terminus indicates the cysteine linkage to the carrier protein.



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Q8. Cell matrix linking via integrin - TRUE/FALSE (Decide whether each of these statements is true or false, and then explain why.)

- a) RGD is a small amino acid responsible for integrin binding of fibronectin.
- b) Inside the cell integrin binds to vinculin, which binds to actin filament.
- c) Inside out and outside in activation have the same result of strong binding.
- d) ECM protein network consists of nidogen, type III collagen, laminin and perlecane.

Q9. Nuclear deformation and related signaling – TRUE/FALSE (Decide whether each of these statements is true or false, and then explain why.)

- a) Integrin senses mechanical stimuli from neighboring cells.
- b) Chromatin is the main cytoskeletal filament in the nucleus.
- c) Nuclear deformation and molecular mechanisms are both mechanisms of nuclear mechanosensing.