### **Objectives**

1.White book: Read Chap 3 & p 77-98 & 1082.Black book: Read Chap 3 & p75-96 & 106

### Objectives:

- 1. List metric measurement units for microorganisms and convert to other metric units (m, mm, um, nm).
- 2. Identify parts & functions of the compound light microscope.
- 3. Define/calculate total magnification & resolution.
- 4. Compare, contrast, and identify uses (diseases/organisms) for brightfield, darkfield, fluorescent, electron-transmission, and electron-scanning microscopy.
- 5. Differentiate, compare, and explain the appearance and uses of each of the following: acidic & basic dyes, simple, differential & special stains, capsule, endospore, acid-fast and flagella stains.

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## Objectives, Cont'd

- Identify the functions of the cell/plasma membrane, chromatophores/thylakoids, nucleoid, ribosomes, endospores (including location), inclusions.
- Transport: passive (simple diffusion, osmosis, facilitated diffusion), active transport, hypertonic, hypotonic, isotonic, osmotic lysis, plasmolysis
- 13. Discuss several pieces of evidence that support the endosymbiotic theory of eukaryotic evolution.
- Describe the overall structure and defining characteristics of prokaryotes, as compared to eukaryotes.
- On given slides identify shape, gram reaction, arrangement, type of stain.

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# Objectives, cont'd

- List specific chemicals that are used for each type of stain in the objective above, primary stain, mordant, decolorizer, counterstain.
- 7. Gram stain: list the steps, purpose, and the appearance of GP & GN cells after each step.
- 8. Identify the 3 basic <u>shapes</u> of bacteria <u>and</u> secondary arrangements.
- 9. Describe the structure & function of the glycocalyx, flagella (including arrangement), axial filaments, fimbriae, pili. Identify flagellar arrangements.
- 10. Compare & contrast the cell walls of GP bacteria, GN bacteria, archaea, mycoplasmas, and mycobacteria. (Including composition, antibiotic & chemical resistance, presence of toxins, staining reactions, effect of penicillin, lysozyme, etc.)

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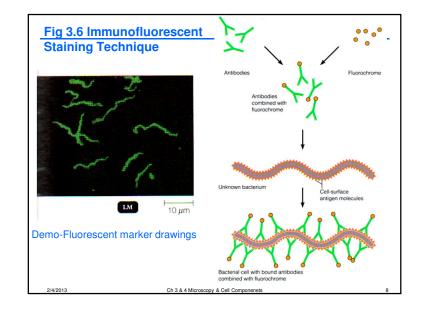
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	Measurement Units & Terms	
1.	<u>Units</u>	
	A. Micrometer (μm) =	
	B. Nanometer (nm) =	
	i. Example: Convert 21.5 nm to m	
	•	
2.	Total Magnification	
3.	Resolution: Distance apart needed to see(Ability to see)	
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Resol	ution & Refractive Index					
A. Res	olving power =					
N.A.	depends on:					
i	of material between lens &					
S	lide.					
ii. T	heof most divergent light ray					
B. To ir	B. To improve resolution:					
i						
ii						
C. Impr	rove conditions but NOT resolution:					
i						
ii						
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Using oil does improve resolution, as it increases the	Unrefracted Oil immersion objective lens
numerical aperture, which will cause a better (smaller) resolving power number	Without immersion most light is refract and lost  Air  Glass sl
	Condenser lenses  Light source
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Scope	Enhanced by	<u>Advantages</u>	<u>Uses</u>
Light, Brightfield: Background Visible light Res: Mag:	& light	Inexpensive Easy to use	Live specimens (unstained) Stained specimens Bacteria, protozoa
Light, Darkfield:  Background  & microbes  Same	N/A	Easier to see microbes	Live microbes:
Light, Fluorescent: Background & microbes Same	Fluorescent dyes: Fluorescent dye on to microbe nicrobe fluoresces	directly from specimen, w/o culture  Detection of microbes compared to other light microscopy	When immediate diagnosis neede When cultures aren't avail, or take long



<u>Scope</u>	Enhanced by	<u>Advantages</u>	<u>Uses</u>
Electron, Scanning Res; Mag;		3-D Book from U of I	Surfaces structures - eukaryote to virus
Electron, Transmission Res Mag	Stain w/+ salt of heavy metal	res & mag  DISADVANTAGE:  Need slice as e- can't All e- scopes due to killing, & fixing under vacuum	Virus particles, bacterial flagella, _ cell structures, protein molecules
Scanned-Probe Res 1/100 of atom	Ch 3 & 4 Microscopy &	Res No special prep	Map atomic & molecular shapes & processes, ie. DNA, fibrin (clot) formation

Electronbeam  Electronagnetic condenser fins Specimen  Viewn open condenser fins Specimen  Viewn open condenser fins Specimen  Viewn open condenser fins Specimen  Specimen  Specimen  Specimen  Specimen  Specimen  Specimen	Pinary electron beam  Electromagnetic lenses  Viewing screen Electron collector  Bectron	Fig 3.8 Transmission vs. Scanning
	The state of the s	
(a) Transmission. (Top) In a transmission electron microscope, electrons pass through the specimen and se scattered Magnetic lenses focus the image orto a fluorescent scene or photogramic plate (Bottom) This collected transmission exclude micrograph collected transmission exclude micrograph that the properties of the properties of the pro- training that the properties of the properties of the transmission of the properties of the pro- training transmission of the properties of the transmission of the properties of the pro- perties of the properties of the properties of the pro- perties of the properties of the properties of the pro- training transmission of the pro- trai	(b) Scanning. (Top) ha scanning electron microscope, primary electrons sweep across the specimen and whose electrons from its surface. These secondary electrons are picked up by a collectric amount of the property of the property electrons are picked property of the property electrons are picked property of the property electrons in this colorized scanning electron micrographs (SEAT), the surface structures of a Paramacum can be seen. Note the three-dimensional appearance of this cell, in contrast	renets

_							
	Stains-Slide Prep & Basic Stains						
<u>Sli</u>	Slide Prep:						
1.	<u>Smear</u>						
2.	<u>Fix</u> –	_ to slide (won't off)					
	A						
	В						
	C						
	D. HOPEFULLY-prese	erves w/					
Sta	<u>aining</u>						
1.	Basic dye/	stain: Colored () ion of a salt					
	A. Attracted to (	_) bacterial cell; stains					
	B. Crystal violet, meth	hylene blue, safranin, malachite green					
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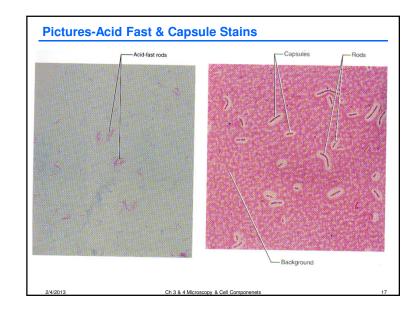
Acidic Dye / Negative S	tain
2. Acidic dye /	stain: Colored () ion
A& s	stains
B. For cell	, to detect
C. Advantage:	(no & stain
	so accurate size & shape)
D. Examples: Acid fuc	chsin, nigrosin
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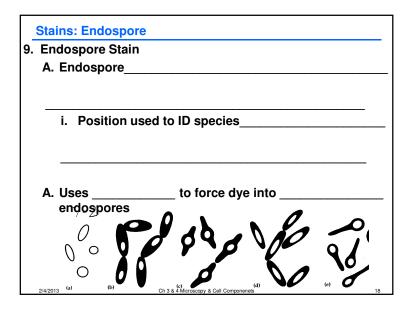
Mordant, Simple Stain, Diffe	erential Stain
3. Mordant: Substance used t NOTE: This is not the stain that only helps the stain be more in	t gives color,
4. Simple stain:	_basic dye
A. All microbes	
B. Only for	
5. Differential Stain: Use of	to
	groups of bacteria
A. Examples: gram stain, a	cid fast stain
A Common of the	
2/4/2013 (b) Spin 'osco	рру & Ce 👊 🖂 13

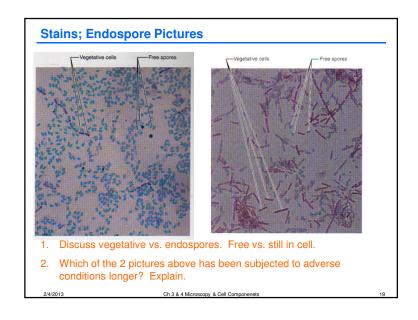
2/4/2013 (b)	oscopy & Ce	M i—i	13
Gram Stain Di	agram		
			Crystal viole lodine Alcohol Safranin
(1:1/3·) →		* - (1.37.*)	
Application of crystal violet (purple dye)	Application of iodine (mordant)     Application of iodine (mordant)		
• Shapes above?			
• GN or GP?			
• Combine?			

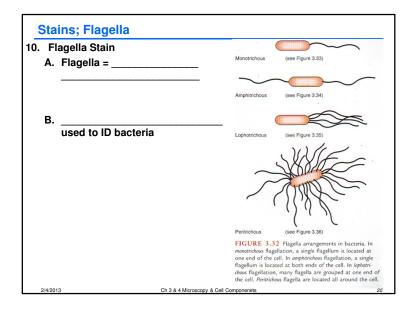
6.	Gram Stain: differences		- due	to	
	A.		= gram positive, Us	, retain to penicillin	stain
	В.		= gram negative, red,		stain &
		i.		to penici	llin
	C.	i.	ining problems Need Some bacteria stain	cultures	
				timing is	
		iv.	Potential	-structures/c ng procedures NOTE: th	listortions that appear
		ow p	ch step, how cell wall o	• • • • • • • • • • • • • • • • • • • •	appearance of cells after ing (Chap 4)

Stains	Stains: Acid Fast & Capsule						
7. Acid F	ast Stain						
A. Ac	id-fast positive =	(due to	in cell)				
B. Ac	cid-fast neg =						
C. ID		species,					
8. Capsu	ıle Stain (w/	stain)					
A. Ca	psule =	covering on o	utside of bacteria				
B. Va	riation w/2 stains:						
i.	•						
ii.							
iii	•	of capsule left be	tween the stains				
C. Pr	oblems: capsule may						
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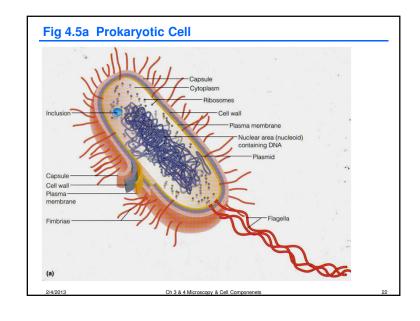


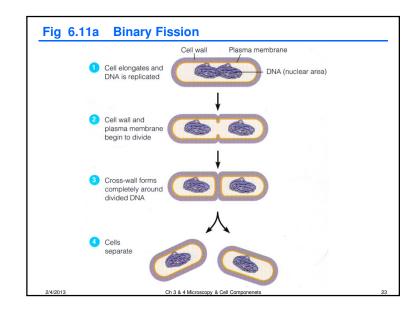


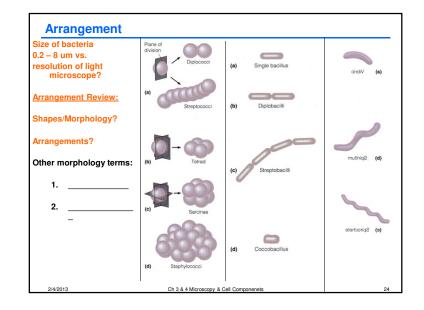




Chapter 4: Prok	caryotic Cells	
<b>Prokaryote</b>		
1		
2		
3		
4		
5. Bacteria – d	cell wall	
6. Archaea – _		
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Cell Wall - Bacteria	
Bacterial Cell Wall	
1	
2. Clinical importance	
, ,	
A	
В	
3	
<b>5</b>	
4. Penicillin interferes	
	_
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Teichoic acid  Peptidoglycan layer  Cell membrane	(+) Gram (-) Bacterial
Inside Cell	Inside Cell

Table – GP vs. GN Cell Wall Characteristics	
GP Wall	GN Wall
1	1
2. Contains	2. None
3. None	OUTER Wall Membrane     A. Evades      B. Contains  C
4. None	4. Periplasm(where peptidoglycan is) A. Contains
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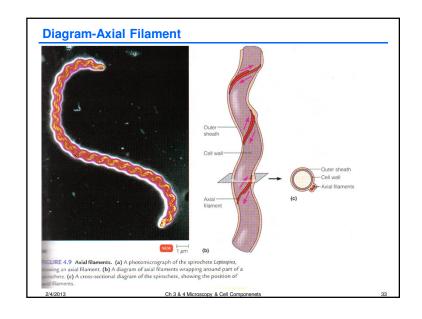
Gra	m Stain & the Cell Wall
Cell W	all & gram stain
1.	lodine =
2.	Alcohol
	A. GP:
	B. GN:
	C. GP falsely stain GN when cell wall damaged due to
3.	GPR/GPB only:
	A: Bacillus & Clostridium
	B: Mycobacterium (TB)
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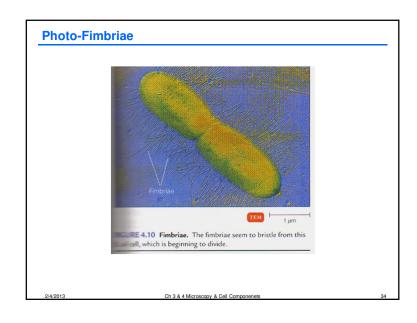
# Chemicals & the Cell Wall Chemical Effects on Cell Wall 1. Lysozyme: A. Most effective on 2. Penicillin

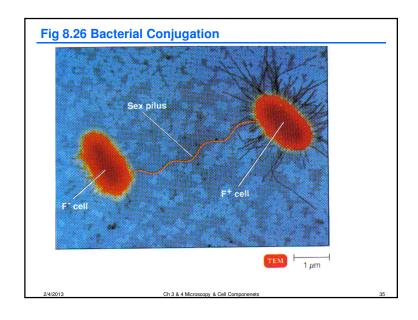
xternal Str	res External to Cell Wall uctures	
1. Glyc	ocalyx/Capsule:	
A. E	EPS (Extracellular polysaccharid	le) & polypeptide polymer
В		
		Capsules
C. 1	Negative Stain, but uses 2 dyes	
i	. Basic stains	
i	ii. Acidic stains	
i	ii	
		Background

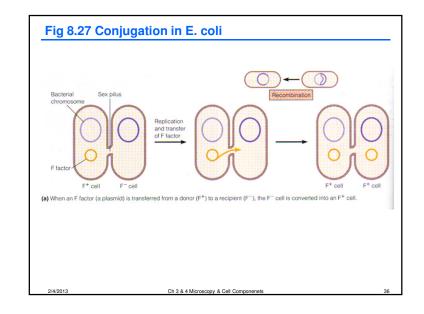
Atypical Cell W Atypical Cell Walls	/alls		
1. <u>Mycoplasma</u>	species:		-
	ount from lysis	in plasma membrane, s	
2. <u>Mycobacteria</u> A	<u>a</u> - High	in wall	
В			
3. Archea;			-
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External Fi 2. Table:	lamentous Struc	tures	
<u>Flagella</u>	Axial Filaments	<u>Fimbrae</u>	<u>Pili</u>
Monotrichous -			
	Spiralled around cell within		
Amphitrichous-	(AKA endoflagella)		
Lophotrichous-	endonagena)		
Peritrichous-	Ch 3 & 4 Microso	copy & Cell Componenets	32



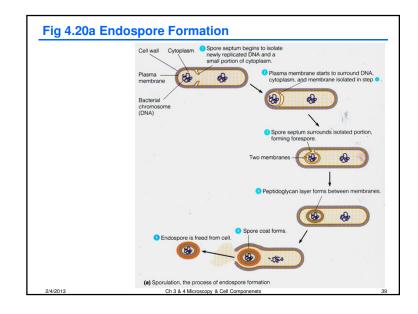


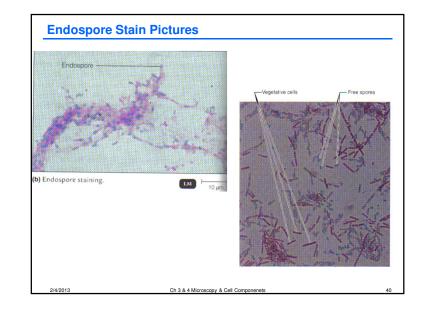


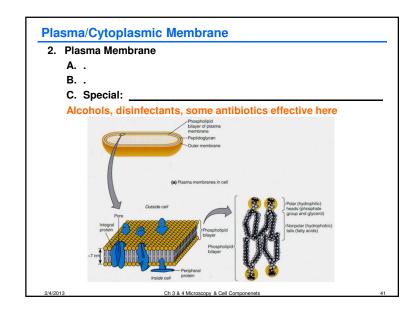


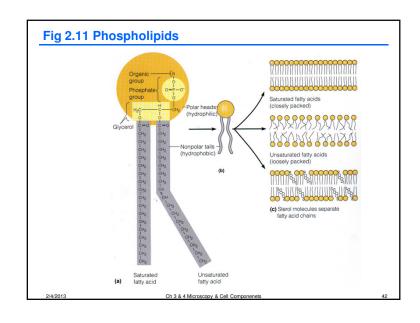
External Filamentou	s Structures, Cont'd	
3. NO		
4. Taxis:		
A. Chemotaxis		
B. Phototaxis		
Discuss serovars		
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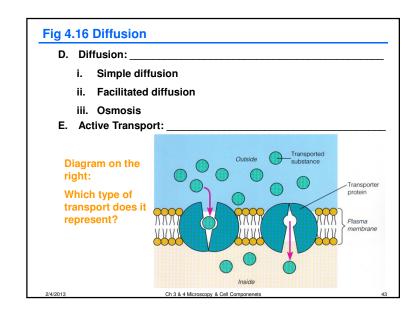
Endospores	
Structure Internal to Cell Wall	
1. Endospores:structures	to
adverse conditions	
A. <u>.</u>	
B. Sporulation / Sporogenesis	
C. Germination – return tostate	
D	
E. Location:	
F. Survive	
G. Stains:	
i. Gram	
ii. Endospore Stain:	
Primary: basic stain	
Rinse: removes stain from	
Counterstain: basic stain colors	
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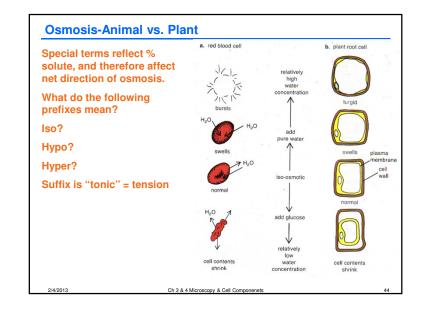




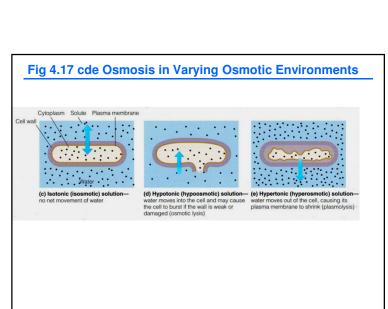








Osmosis & Solution Types	
F. Osmotic Environments	
i. Isotonic/isoosmotic solution:	
» .	
» Water movement	
» .	
ii. Hypotonic solution:	_
» Net H <sub>2</sub> O moves	
» <u>.</u>	
iii. Hypertonic <u>solution</u> :	_
» Net H <sub>2</sub> O movement	
» .	
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	Isotonic solution	Hypotonic solution	Hypertonic solution
Animal cell	HO HO	HO HO	140
	(1) Normal	(2) Lysed	(3) Shriveled
Plant cell	H,O	H <sub>0</sub> 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	Plasma membrane H,O
	(4) Flaccid	(5) Turgid	(plasmolyzed)

Inte	ernal Cell Structures continued	
	Chromatophores/thylakoids:	
4.	Nucleoid/nuclear area: No nuclear membrane	
	A. Contains	
5.	Plasmids:	
	A	
	B. Conjugation: transfer through	
	i. GN	
	ii. GP	
	C. Biotech:	
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