

01

RULES SHEET

02

DIFFICULT TOPICS

03

COMMON QUESTIONS

04

TIPS FROM A VETERAN

05

OTHER FREE RESOURCES



The Rules Sheet

- May bring 18.5" by 11" sheet of paper, both sides
 - No additions on top of the paper
 - Can be in a a sheet protector sealed with tape or laminated
- 2 non-programmable, non-graphing calculators allowed
- Make sure that you and your teammate know every topic at least a surface level.
- Either as lab-practical stations or an exam



ATEAM OF UPTO: 2



EYE PROTECTION: C

- THE COMPETITION: This Event may be administered as a written test or as a series of lab-practical THE COMPETITION. This Frest may be administred as a writen test or as a series of abspacetical statements which can include but are not inflamed to experiments, including appratum, models, illustrations, as the contract of the contract of
- Microscopy;

 (1) Describe the parts, functions, images, and sample preparation of bright-field, phase contrast, fluorescence, and electron (TEM & SEM) microscopes.
- fluorescence, and electron (TEM, & SEM) microscopes.

 (2) Identify and explain which microscopy method is most appropriate to address a given hypothesis or experimental goal.

 (2) Estimate the sixt or microbes tising scale bars. Calculate magnification and resolution using power and numerical aperture data. Determine direct cell counts (in cellibrill) using a Neulus counting chamber (exact chamber dimensions to be provided by the Exam writer).
- and eukaryotic (i.e., microalgal and fungal) cells (i.e., membrane, cell wall, flagella, pilus, fimbria, mclevid, cytoplasma, and organicles) and of specialized situetures in buceria and eukaryotic microbes (i.e., gas vesicles, endospores, contractile vacuoles, cycopots,
- carroxysomes).

 Contract Gram (+), Gram (•), and acid-fast cells and explain the Gram stain procedure.

 Describe basic structural components of vinuses and their functions.

 State and Nationals only. Describe different forms of cell locomotion (swimming and glidling motility) and discuss chemotaxis and phototaxis.
- The Describe applications of different methods to culture bacteria (i.e., liquid vs. agar) and different (1) Discribe applications of different methods to culture bacteria (i.e., liqual vs. agar) and different (i.e., liqual vs. agar) and different liqual vs. agar) and vs. agar and vs. agar and vs. agar as a second vs. aga

- filtration, and chemical) are able to compromise/eliminate microbes. Understand the limitations of culture-based approaches to study microbes.
- (1) Outline the steps of bacterial cell division (i.e., binary fission) and genome replic
- including the function and properties of the origin of replication, DNA unwinding element DuaA, and DNA polymerase. <u>State and Nationals only</u>: Outline the steps of relling circle replication and identify microbes or agents that use this strategy. Outline the steps of bacterial transcription and translation, including major enzymes invol-

MICROBE MISSION C (CONT.)



- (4) Sing and Nationals only: Describe the properties and function of plasmids in bacteria. Discuss how recombinant DNA technology is used to produce useful products such as human insulin. Metabolism and Applications:
 (1) Describe microbial metabolis strategies based on carbon and energy sources.

- (i) Describe microdula metabolic instiguis based on orders and emerge source.

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 strictural production, policyophiese in biolical production, and strapper factors in the

 strictural recipitation and carbon fraction, using the Wingmahay column as a model system.

 (ii) Discoult for conversability theory of agentic revolution,

 (iii) Discoult for conversability theory of agentic revolution,

 (iii) Discoult for conversability theory of agentic revolution,

 (iii) Discoult for conversability theory of agentic revolution,

 - Describe lytic and lysogenic viral life cycles with examples from the Microbes and Agents
- (4) Describe how ornomic analysis can be used to determine the functional potential and
- evolutionary history of a microbe.

 (5) Outline the mechanisms of horizontal gene transfer (i.e., transduction, conjugation, antransformation). Explain the role of horizontal gene transfer and viral infection in evolutio (6) Describe applications and limitations of IcS amplicon sequencing, interpret data from
- amplicon sequencing experiments (i.e., bacterial community composition, alpha diversity beta diversity), outline how PCR is used to target specific genes in amplicon sequencing
- ensalism, predation, parasitism). Explain how these interactions can be mediated b
- metabolic portways.

 (8) State and Nationals only: Describe applications and limitations of metagenomic and metamatabolic particular and metagenomic and met
- (6) State and Microsoft such: "Distribute applications and allimitations of metaposomers and meta-posomers are present to a large and present to a large and present to a separation and particular and prospensition and part pure hypothesis or experimental gual." Continued and present and particular and present and present and particular and parti
- aformation will be provided to answer questions.

 Bacteria: Escherichia coli, Richettia richettiii, Mycobacterium leprae, Mycobacterium tuberculos Microcytis aeruginosa, Saphylococcus ameus, Helicobacter pejori
- ii. Archsen: Pyrococcus, suppyrococcus sp. iii. Euktryotes: Plasmodium faiciparum, Saccharonyces cerevisiae, Nannochloropsis sp., Paramecius

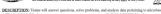
Recommended Resources: The Science Olympiad Store (store.soinc.org) carries a variety of resources to

Fundamental Topics

- Microscopy
- Structures of Bacteria, Archaea, Eukarya, and Viruses.
- Gram Staining.
- Molecular Biology of Bacteria*
- Metabolic Pathways
- **Evolution & Ecology**



ATEAM OF UPTO: 2



EYE PROTECTION: C

- ACLICATOR Cass 11
 2. ENEXT PRABETERS: For events with a lab practical portion, each shadent must wear goggles. Each tram may bring one 8.5" XII" sheet of puper, which may be in a sheet protector realed by uper of unimated, but may cotain information on book sides in any form and from any source without any amoutations or labels affixed along with two stand-since non-programmable, non-graphing calculators (Class II), Any measurements must be made to the precision of the device.
- THE COMPETITION: This Event may be administered as a written test or as a series of lab-practical . III. COMPETITION: This levest may be administered as a writte set or as a series of alb-practical incincion which can unitable but are not intermited but are present incincion spatrate, manchine, limentations, manchine and the second of t

- below without any overemphasis in any one particular topic. The list of topics is exhaustive.

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 A for each of the defining upone, particular, will be expected to see quantiture transmitting and the list of th
- Describe the basic sometime, composition, and anticensor or composition to basic retaining and cultury robis (i.e., microsaga) and fungas) cells (i.e., membrane, cell wall, flagella, polus, fimbria, meckoid, cytoplasm, and organelles) and of specialized structures in succeria and cultury for microsaga (i.e., gas vesicles, endospores, contractile vacuo les, eyespots,
- Carronysomes).

 (Contrast Gram (+), Gram (•), and acid-fast cells and explain the Gram stain procedure.

 (Describe basic structural components of viruses and their functions.

 State and Nationals only. Describe different forms of cell locomotion (swimming and glidling modifier) and discuss chemotaxis and phototaxis.
- gating mortinty) and discuss enemotaxis and phototaxis.

 Culture and Growth:

 (1) Describe applications of different methods to culture bacteria (i.e., liquid vs. agar) and different
- (1) Describe applications of different methods in culture bacteria (i.e., Inquid vs. agar) and different (i.e., Inquid vs. agar) and different methods in culture bacteria (i.e., Inquid vs. agar) and different (2) Integret bacterial growth curves and discuss white is happening at each stage. (3) Describe how plate count data (i.e., CFUs) and optical dentity measurements are used to calculate the manner of cribe in a culture and populating growth rate. (i.e., See a consistent of the contract of the contract of the contract of the contract of the calculation of the contract of the co

- filtration, and chemical) are able to compromise/eliminate microbes. Understand the limitations of culture-based approaches to study microbes.
- Molecular Biology:

 (1) Outline the steps of bacterial cell division (i.e., binary fission) and genome replicatio including the function and properties of the origin of replication, DNA unwinding element DnaA, and DNA polymerase. State and Nationals only: Outline the steps of rolling circle replication and identify microbes or agents that use this strategy.

 (2) Outline the steps of bacterial transcription and translation, including major enzymes involved.

MICROBE MISSION C (CONT.)



- (1) Describe microbilat attachioù interigeire based on carbon and emergy source; mentation, arrayen; phosphorpulents, misogra franciazion al devent free your in the cell, describe a repetit production; and service procure in the cell, describe a repetit production; photocytichen in holding production; and interpet fusation in the attachment production; photocytichen in holding production; and interpet fusation in the attachment production; photocytichen in holding production; and interpet fusation in the attachment production; photocytichen in microbilat progression data in Clark and interpetation and carbon fusation, using the Williagendoky column as a model system.
 (1) Describe the consupposition describe of specialize revolution.
- (2) Describe common adaptations to environmental extremes (i.e. temperature, salinity, pH).
 (3) Describe lytic and lysogenic viral life cycles with examples from the Microbes and Agents
- (4) Describe how genomic analysis can be used to determine the functional potential and
- evolutionary history of a microbe.

 (5) Outline the mechanisms of horizontal gene transfer (i.e., transduction, conjugation, and transformation). Explain the role of horizontal gene transfer and viral infection in evolution.

 (6) Describe applications and limitations of 16S amplicon sequencing, interpret data from 16S
- amplicon sequencing experiments (i.e., bacterial community composition, alpha diversity, beta diversity), outline how PCR is used to target specific genes in amplicon sequencing experiments.

 (7) Identify and describe community interactions between microbes (i.e., coor
- ommensalism, predation, parasitism). Explain how these interactions can be mediated by metabolic pathways.

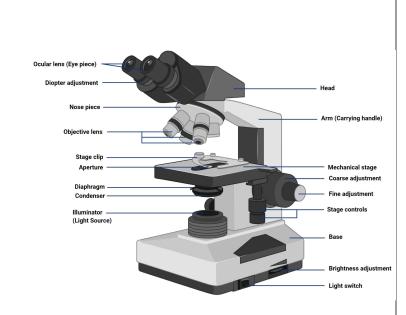
 (8) State and Nationals only: Describe applications and limitations of metagenomic and meta-
- (6) Size and Miritarshi, and P. Discible explorations and intendions of metaponenic and metaponenic and metaponenic properties of the p
- information will be provided to answer questions.

 Batteria: Excherichia coli, Rickettria rickettsii, Mycobacterium leprae, Mycobacterium tuberculosis, Microcystis aerugiosas, Saphylococcus aureus, Helicobacter pylori
- Archives: Pyrococcus furiosus, Methanococcus sp.
 Eukaryotes: Plasmodium falciparum, Saccharonyces cerevisiae, Nannochloropsis sp., Paramecium
- High score wins. Selected questions may be used as tiebreakers
 Points will be awarded for quality and accuracy of answers, quality of supporting reasoning, and the use of proper scientific methods.

Recommended Resources: The Science Olympiad Store (store.soinc.org) carries a variety of resources to

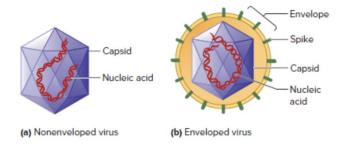
Topic 1: Microscopy

- The anatomy of bright-field microscopy
 - Know the functions and its limits as well.
- Know the appropriate formulas for resolution
 - O Abbe's Equation: $d = \frac{0.5\lambda}{n \times \sin(\theta)}$
- The purpose of dark-field microscopy.
 - Living & unstained microbes
- Fluorescence Microscopy
- SEM & TEM
 - Differences and use



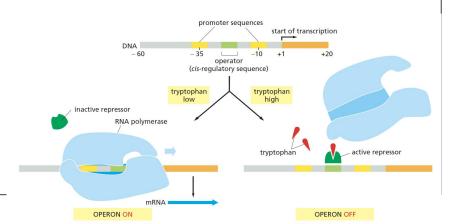
Topic 2: Structures

- Common bacterial and archaeal internal structures and their functions.
 - Carboxysomes, (CO2 Fixation); Storage Inclusions (PHB)
- Compare and contrast the differences of cell membranes & wall's of archaea, eukarya, and bacteria.
 - Ether-linkage vs. ester-linkages
- Compare and contrast Gram (+) and Gram (-).
 - What about the atypical bacteria? Why?
- Noneveloped vs. enveloped viruses.



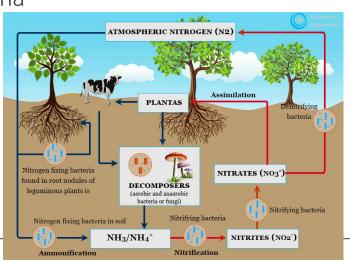
Topic 3: Molecular Bio

- Rule sheet limits the molecular biology aspect to bacteria.
- Model bacteria is E.coli.
- Know the functions of each commonly asked proteins:
 - DnaA (initiator of replication)
 - O DNA Pol I & III
 - DNA Ligase
- Lac and Trp Operons
- Bacterial Translation
 - Shine-Dalgarno Sequence?



Topic 4: Metabolic Pathways

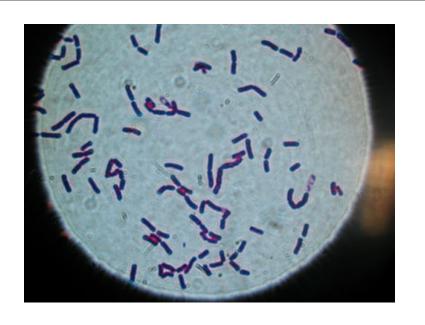
- Cheat Sheet!
- Understand difference between catabolism and anabolism.
- Know the general concepts of common pathways: Glycolysis, TCA, and ETC.
- Then, known the more specialized pathways:
 - ED-Pathways: Gram-negative, mostly soil-bacteria
 - o ETCs of E.coli (bo and bd?)
- Different types of fermentation.
 - Lactic Acid
 - Ethanol
- Nitrogen Fixation



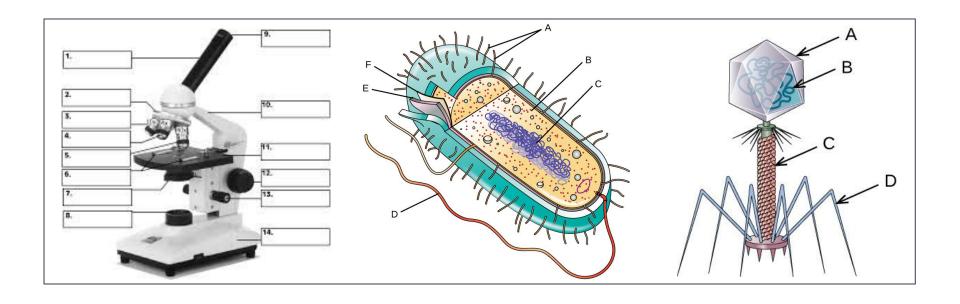


Topic 1: Identification

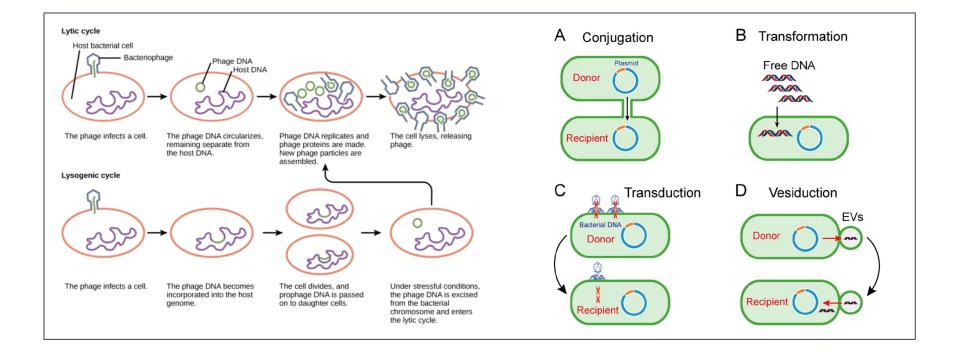




Topic 2: Specific Naming



Topic 3: Processes

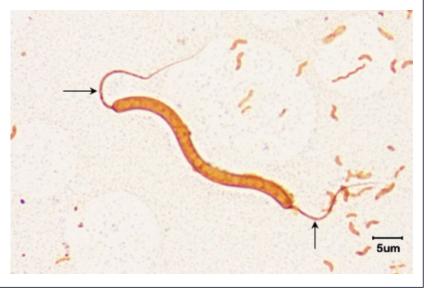




All of the following questions have been pulled from past YJI exams (which can be found on our website) or the Text Exchange on SciOly Wiki

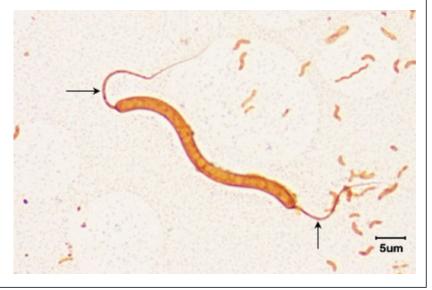
• Name this type of flagellar arrangement.

- Monotrichous
- Amphitrichous
- Lophotrichous
- Peritrichous



• Name this type of flagellar arrangement.

- Monotrichous
- Amphitrichous
- Lophotrichous
- Peritrichous
- Amphitrichous bacteria have flagella on both ends of their body.



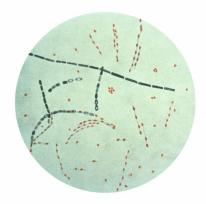
• Which of the following is NOT a viral disease?

- AIDS
- Measles
- Anthrax
- Rubella



Which of the following is NOT a viral disease?

- AIDS caused by the human immunodeficiency virus (HIV)
- Measles caused by Morbillivirus hominis
- Anthrax caused by Bacillus anthracis
- Rubella caused by *Rubivirus rubellae*



- Which of the following types of microscopy would most likely be used to view the insides of organelles?
 - Darkfield
 - Scanning Electron
 - Transmission Electron
 - X-Ray
 - Phase Contrast
 - Differential Interference Contrast

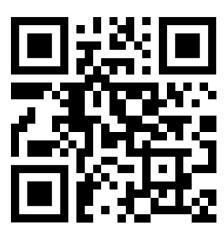
- Which of the following types of microscopy would most likely be used to view the insides of organelles?
 - Answer: Transmission Electron
 - Electron microscopy High Resolution at High Magnification
 - Scanning Electron Sees structure surface by reflecting electrons
 - Transmission Electron Electrons pass through structures, visualizing internal structure

• Given a stock protein solution with a concentration of 15 mg/ml, determine the protein concentration (in mg/ml) of a solution made by mixing 2 μL of the stock with 8 μL of a buffer.

- Given a stock protein solution with a concentration of 15 mg/ml, determine the protein concentration (in mg/ml) of a solution made by mixing 2 μL of the stock with 8 μL of a buffer.
 - Answer: 3 mg/mL
 - \circ V1 * M1 = V2 * M2
 - \circ (2 μ L) * (15 mg/mL) = (10 μ L) * (M2)
 - \circ 30 = 10 * M2
 - \circ M2 = 3 mg/mL

More Example Q's

• If you want to find more example questions, head to the Scioly Wiki Test Exchange located at https://scioly.org/tests/.

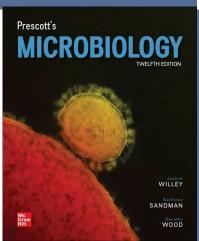


Tips from a Veteran

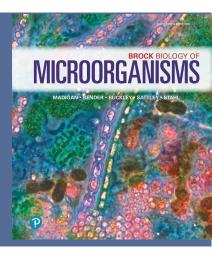
- Prepare well beforehand. Don't cram the information.
- Only include information on your notes sheet that you are sure you can't remember. The limited space is valuable!
- Include images over words.
- Be sure to practice several tests to learn how questions are asked.
 - Tests v. Stations
- Study the classic microbiology textbooks many questions are usually based on the contents found in these books.

Additional Resources

Prescott's Microbiology



Brock's Biology of Microorganisms



OpenStax Microbiology



Your Teachers!

THANKS!

