

## CONCEPT: 2) AVOIDING PHAGOCYTOSIS

● *Recall:* Phagocytic cells destroy microbes in a series of steps (*chemotaxis, recognition, attachment, engulfment, fusion*).

### Pathogens Avoid an Encounter with Phagocytes

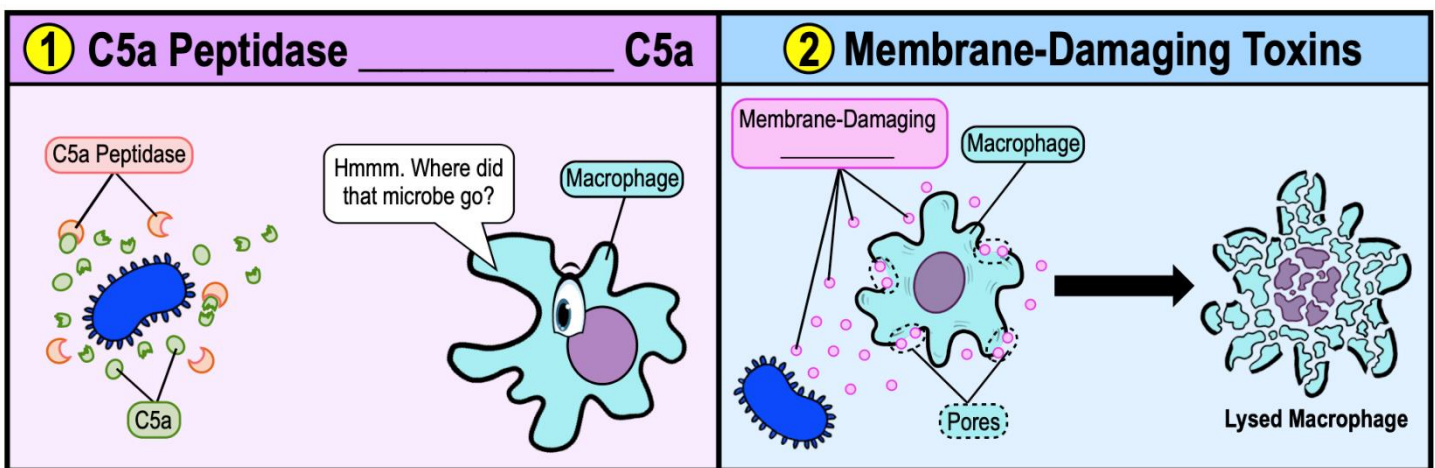
● Certain pathogens can avoid encountering phagocytic cells using one of \_\_\_\_ mechanisms:

1) Production of the enzyme \_\_\_\_\_ **Peptidase** which degrades the complement system protein C5a.

□ *Recall:* C5a is the chemoattractant that recruits phagocytic cells to sites of infection.

2) Production of \_\_\_\_\_-Damaging Toxins that kill phagocytes & other cells.

□ Form \_\_\_\_\_ in membranes of phagocytic cells.



● Some pathogens can survive phagocytosis by phagocytic cells & induce apoptosis once inside the cell.

**PRACTICE:** Why do some pathogens destroy the host organism's C5a proteins?

- The phagocytic immune cells are not attracted to the pathogen without C5a.
- The phagocytic immune cells cannot bind to the pathogen without C5a.
- The membrane of the phagocytic immune cells degrades without C5a.

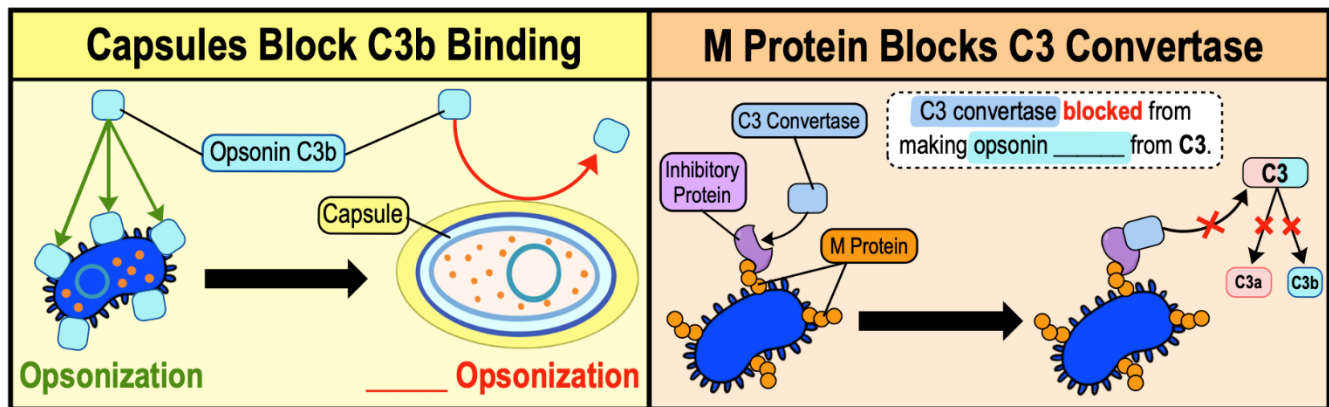
**PRACTICE:** What benefit do pathogenic bacteria obtain from secreting membrane-damaging toxins?

- The toxins will damage any surrounding immune cells.
- The toxins will stop the immune cells from phagocytosing the pathogen.
- The toxins will stop the immune cells from sending messages to other immune cells.
- All of the above.
- None of the above.

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### Pathogens Use Capsules & M Proteins to Avoid Opsonization

- **Recall: Opsonization:** process that makes microbes *easier* to bind/engulf during *phagocytosis*.
- Some pathogens can prevent opsonization by the production of a \_\_\_\_\_.
  - Capsules can prevent immune cells & opsonins (like C3b) from adhering to pathogen.
- Similar to capsules, production of the cell wall protein (\_\_\_\_\_ **Protein**) allows a pathogen to avoid opsonization.
  - M protein binds a regulatory protein that degrades C3 Convertase, thereby \_\_\_\_\_ C3b production.

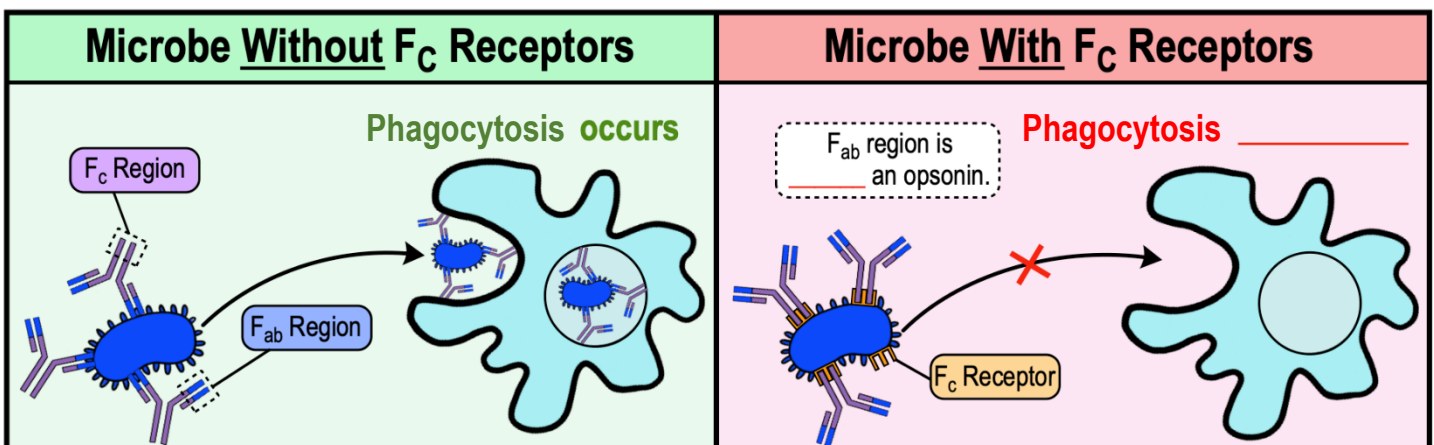


**PRACTICE:** The C3b complement protein is incredibly important for opsonization of pathogenic material. How do some pathogens avoid opsonization by interfering with C3b?

- a) Pathogen has outer layer that C3b cannot bind to.
- b) Pathogen inhibits the functions of C3 convertase.
- c) Pathogen stops the formation of C3b.
- d) All of the above.

### Pathogens use F<sub>c</sub> Receptors to Prevent Opsonization

- **Recall:** When antibodies bind a pathogen, the F<sub>c</sub> region points *outward* & can serve as an *opsonin* for phagocytes.
- **F<sub>c</sub> Receptors:** surface proteins that bind the F<sub>c</sub> region of antibodies & *interfere* with their function as \_\_\_\_\_.
  - Pathogens with F<sub>c</sub> receptors bind antibodies & orient them so the F<sub>c</sub> region is points *outward* (instead of F<sub>c</sub>).
  - Since the F<sub>ab</sub> region is NOT an opsonin, pathogens with F<sub>c</sub> receptors \_\_\_\_\_ opsonization & phagocytosis.



**CONCEPT: 2) AVOIDING PHAGOCYTOSIS**

**PRACTICE:** Certain pathogens have Fc receptors on their surface to avoid which immune responses to infection?

- a) Opsonin proteins binding to the pathogen and triggering opsonization.
- b) C5a attracting phagocytic immune cells to the pathogen's location.
- c) Triggering the classical pathway of the complement system.
- d) Release of histamines trigger an inflammatory response.

**PRACTICE:** Why would some pathogens inhibit the functions of C3 convertase as a defense mechanism against the host's immune cells?

- a) To inhibit the formation of opsonin proteins that bind the pathogen.
- b) To inhibit the production of antibodies that recognize the pathogen.
- c) To inhibit the activation of T cells that trigger a larger immune response.
- d) To inhibit the differentiation of naive B cells into plasma cells.

**PRACTICE:** *Streptococcus pyogenes* creates a C5a peptidase enzyme which breaks down the host's C5a proteins.

Breaking down the C5a proteins has what result?

- a) Stopping the opsonization of *Streptococcus pyogenes*.
- b) Killing the phagocytic cells.
- c) Cell lysis of *Streptococcus pyogenes*.
- d) Inhibition of host's membrane attack complexes.
- e) Decreased accumulation of phagocytic immune cells.