The background features a series of concentric circles in light gray and dashed lines, creating a ripple effect. A large, solid blue speech bubble shape is centered on the page, containing the title and author information.

# Bacterial Structure (Lab 2)

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# 1. Why study Bacterial Cell Structure?

- **Mechanisms of virulence.**
- **Drug development.**
- **Identification**

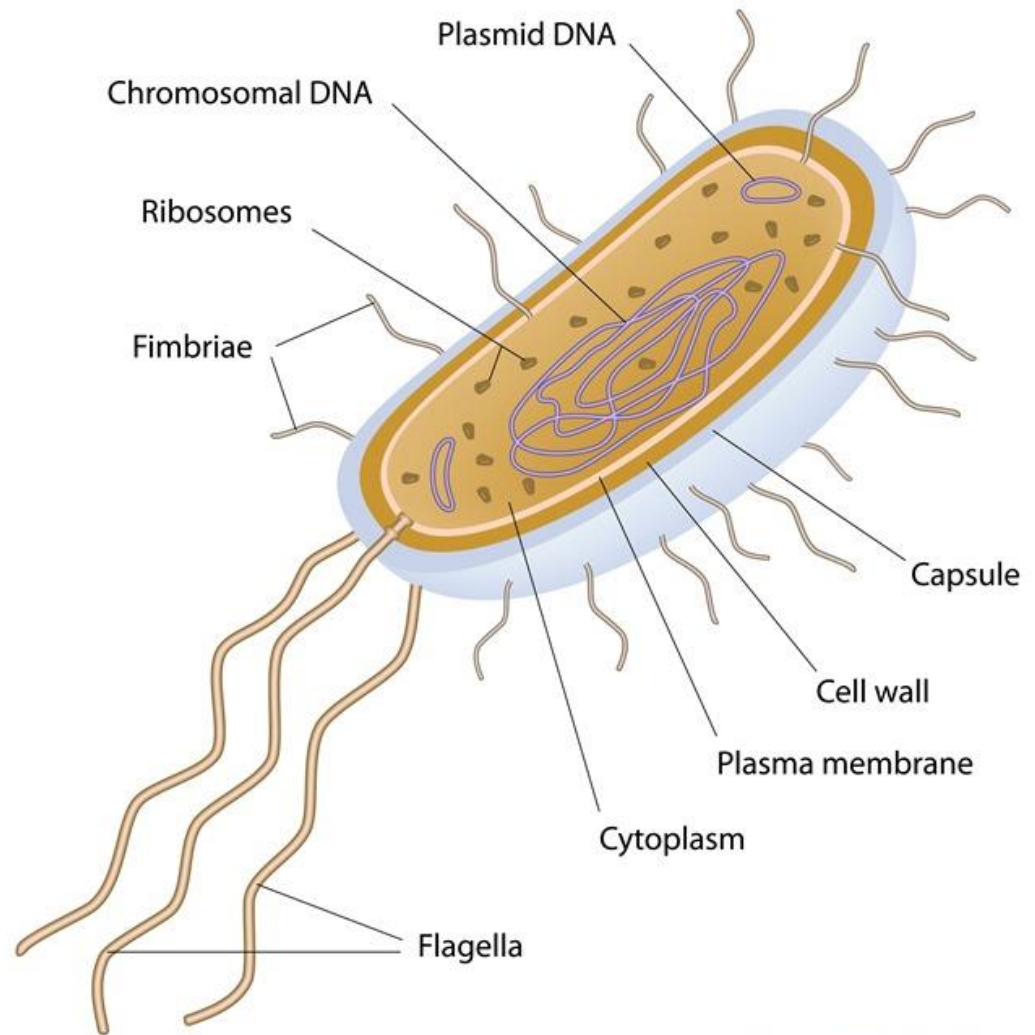
## 2. Essential structures

- **Cell wall.**
- **Cell membrane.**
- **Cytoplasm.**
- **Nuclear material.**

### 3. Particular structures

- **Capsule.**
- **Flagella.**
- **Pili.**
- **Fimbriae.**
- **Spore.**

## Bacterial Structure (Cell wall)



## Characteristics and Functions of Cell wall

- **Outer most portion /barrier.**
- **Protection from turgor pressure.**
- **Gives shape.**
- **Surrounds plasma membrane.**

## Common Cell wall Components

- The bacterial cell wall consists of peptidoglycan, an essential protective barrier for bacterial cells that encapsulates the cytoplasmic membrane of both Gram-positive and Gram-negative bacterial cells.
- Peptidoglycan is a rigid, highly conserved, complex structure of polymeric carbohydrates and amino acids

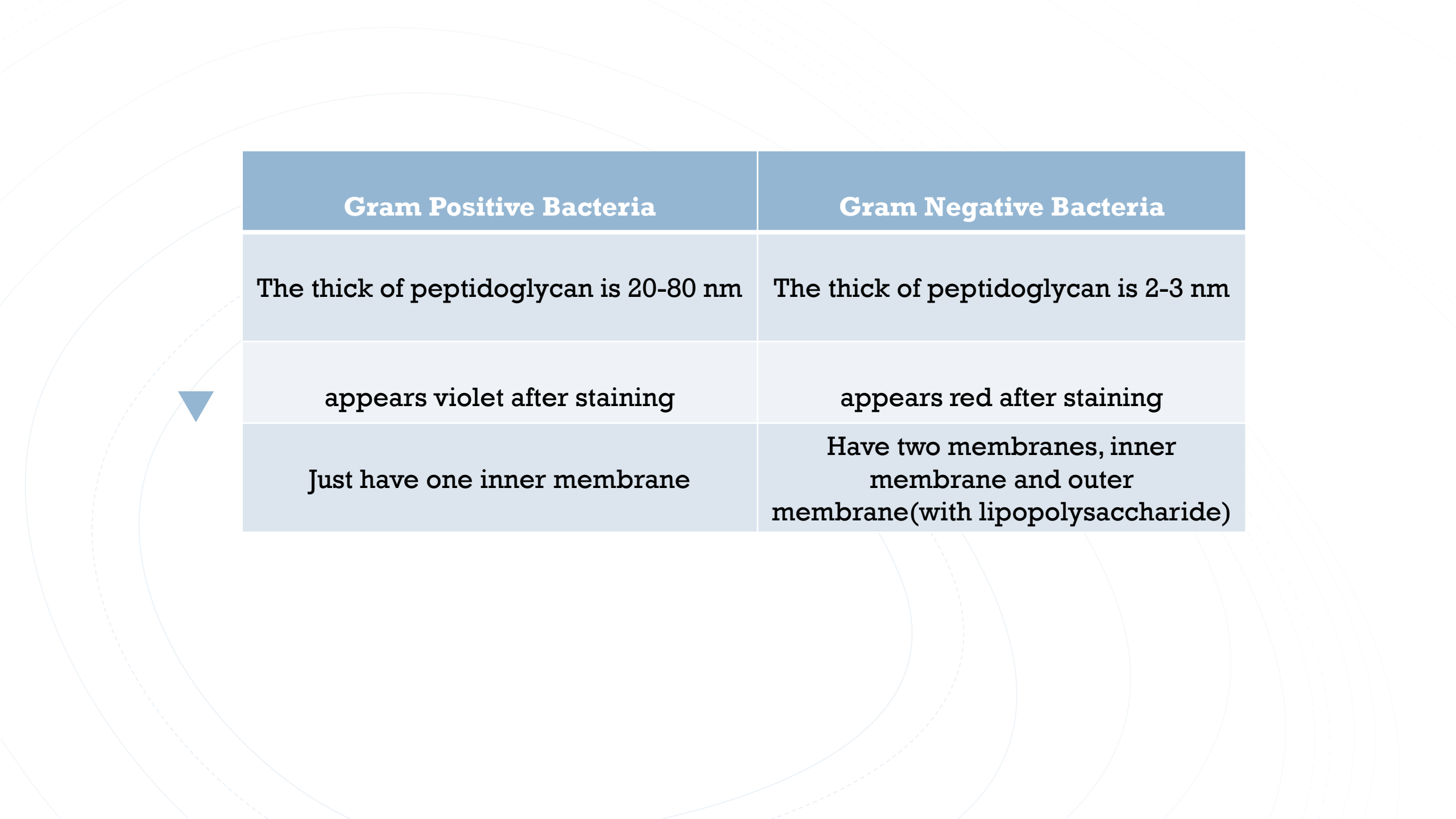
# Cell wall Identification

- **1884 Christian Gram (First publication for the Gram stain method)**



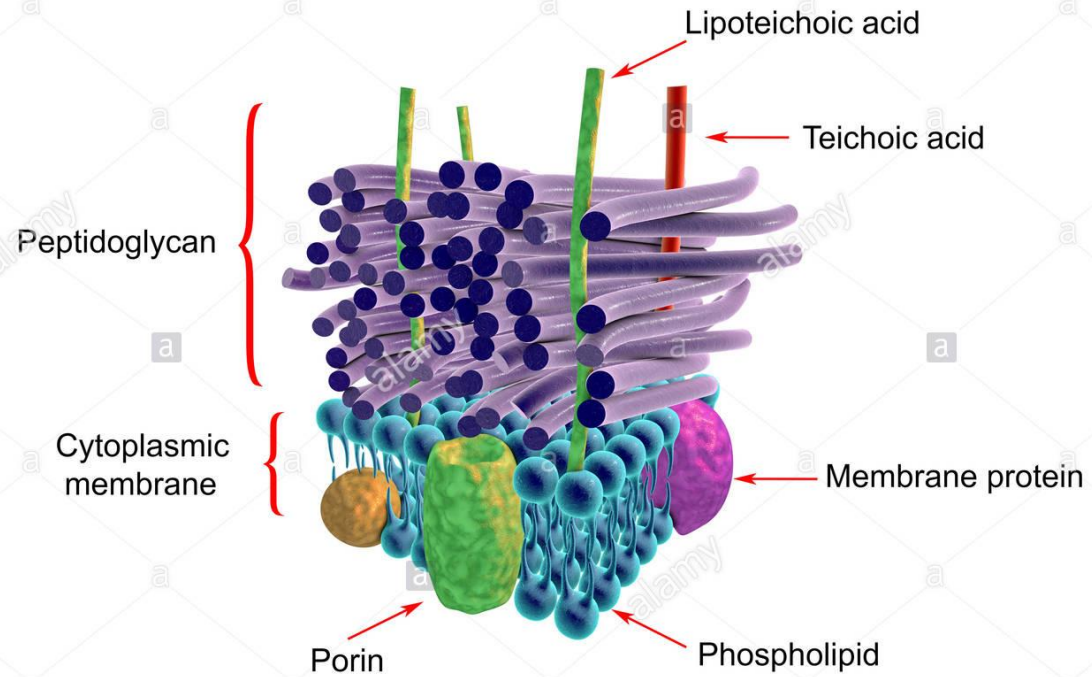
Classification  
based upon  
staining.

- **Gram Positive vs Gram Negative:**
  1. **Hans Christian Gram (1884) differentiate between Positive and Negative.**
  2. **It is the most important differential stain used in bacteriology because (it classified bacteria into two major groups)**
  3. **Gram-positive organisms have a thicker peptidoglycan cell wall compared with gram-negative bacteria**

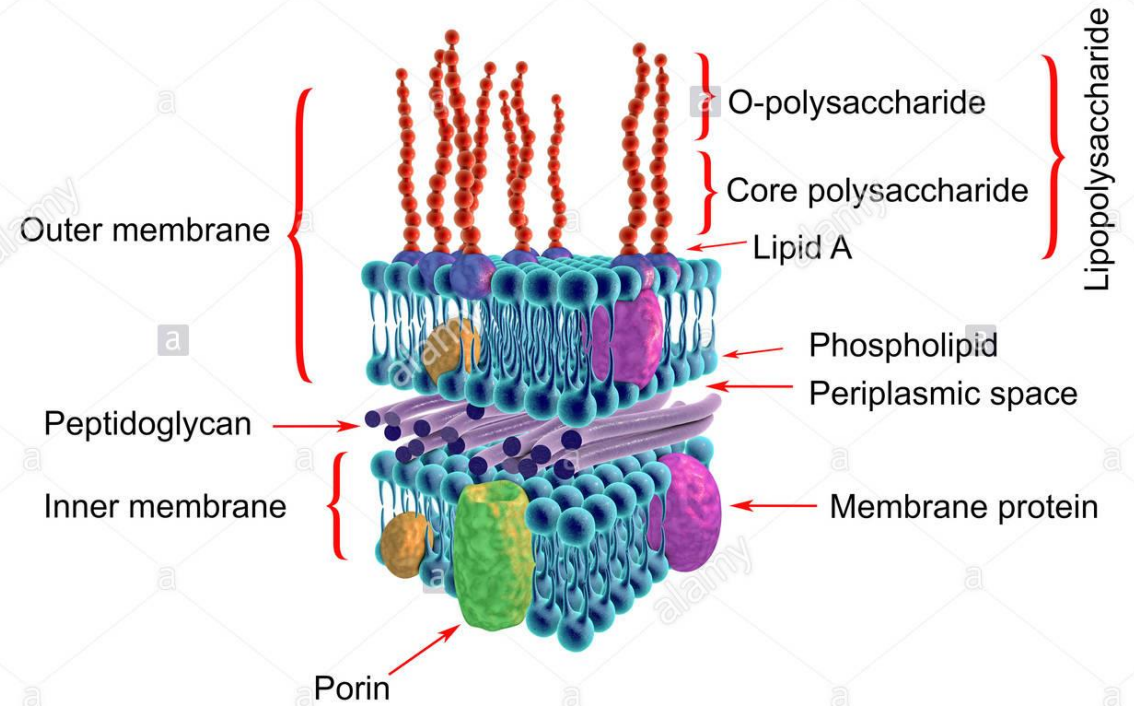


<b>Gram Positive Bacteria</b>	<b>Gram Negative Bacteria</b>
The thick of peptidoglycan is 20-80 nm	The thick of peptidoglycan is 2-3 nm
appears violet after staining	appears red after staining
Just have one inner membrane	Have two membranes, inner membrane and outer membrane(with lipopolysaccharide)

## Gram Positive



## Gram Negative



# Gram Stain

- This stain for differentiate between **G +ve** bacteria (which accept staining), and **G -ve** bacteria (which not accept staining).

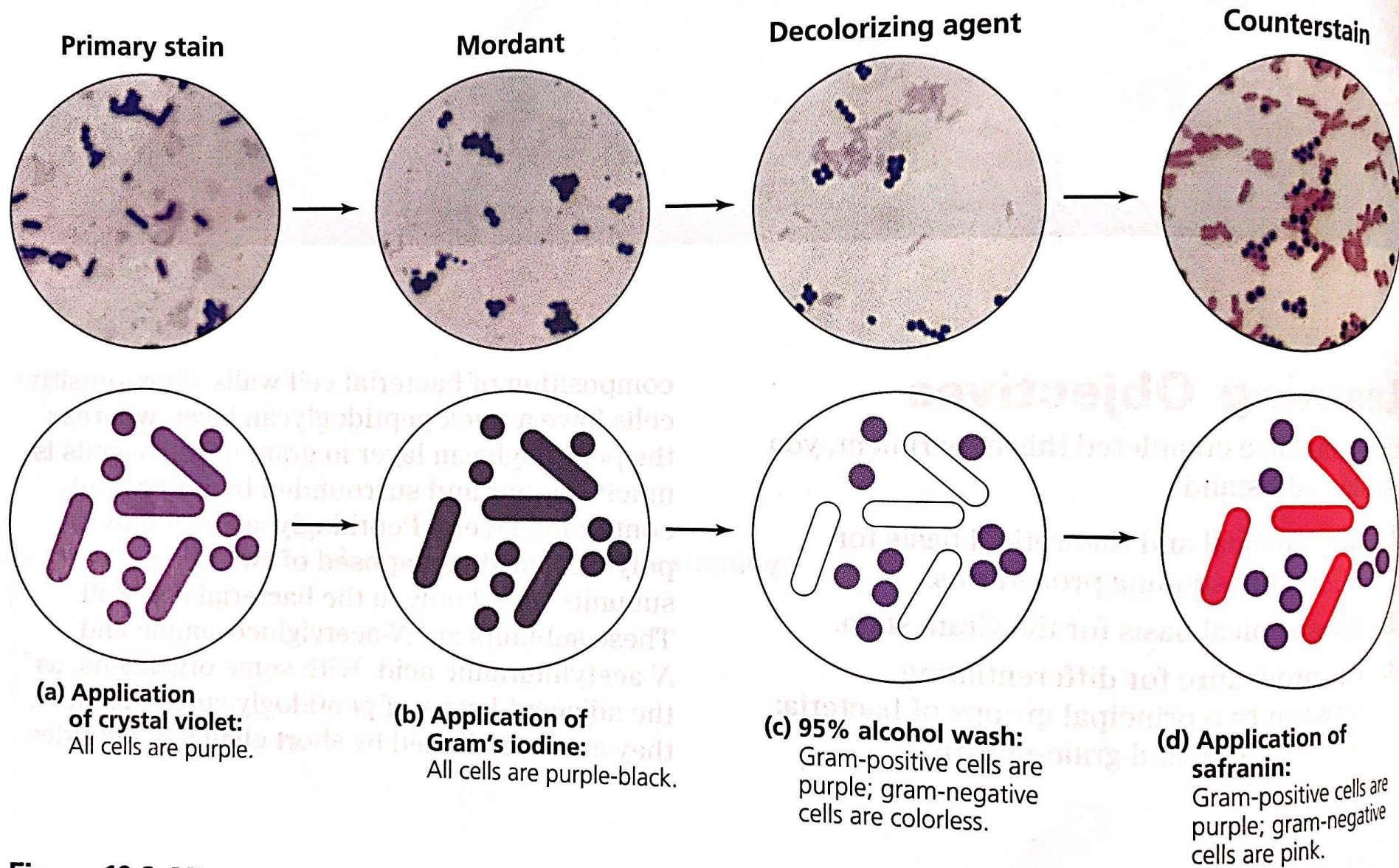
# Procedure

- Prepare your clear slide and put one drop of H<sub>2</sub>O .
- Take specimen from the colony and mix it well with water or normal saline.
- Leave for dryness on air then fix it on burner.
- Then pour Crystal Violet stain on the slide and leave it for 1 minute.
- Wash gently with water..

# Procedure

- Then pour Iodine on the slide and leave it for 1 minute.
- Wash gently with water.
- Wash with Alcohol to decolorize the slide.
- Then pour Safranin stain on the slide and leave it for 1 minute.
- Examine the finished slide under a microscope





**Figure 10.2** Microscopic observation of cells following steps in the Gram staining procedure





## Reference

- <https://www.sciencedirect.com/topics/medicine-and-dentistry/bacterial-cell-wall>
- <https://www.ncbi.nlm.nih.gov/books/NBK470553/>
- **James G. Cappuccino, Natalie Sherman. 2014. Microbiology a laboratory manual.**