



## Lecture-14: *Streptococcus pneumoniae*

### Morphology:

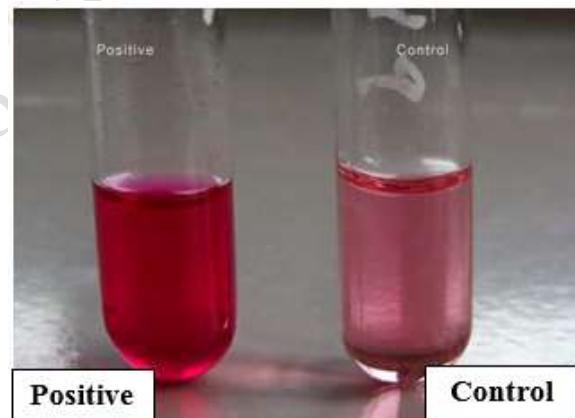
1-gram-positive cocci in pairs (diplococci), 2- slightly elongated cocci, with one end rounded, 3- non-motile and non-sporing, 4- all freshly isolated strains are capsulated and the capsule encloses each pair.

### Cultural Characteristics

- 1-They are aerobes and facultative anaerobes.
- 2- It grows best in air or hydrogen with 5-10 percent CO<sub>2</sub>.

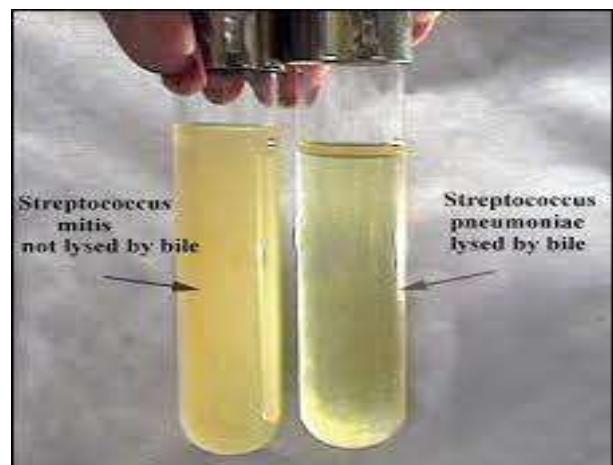
### Biochemical Reactions

1. **Inulin Fermentation:** Pneumococci ferment inulin with the production of acid without gas. Fermentation of inulin by pneumococci is a useful test for differentiating them from streptococci as the latter do not ferment it.

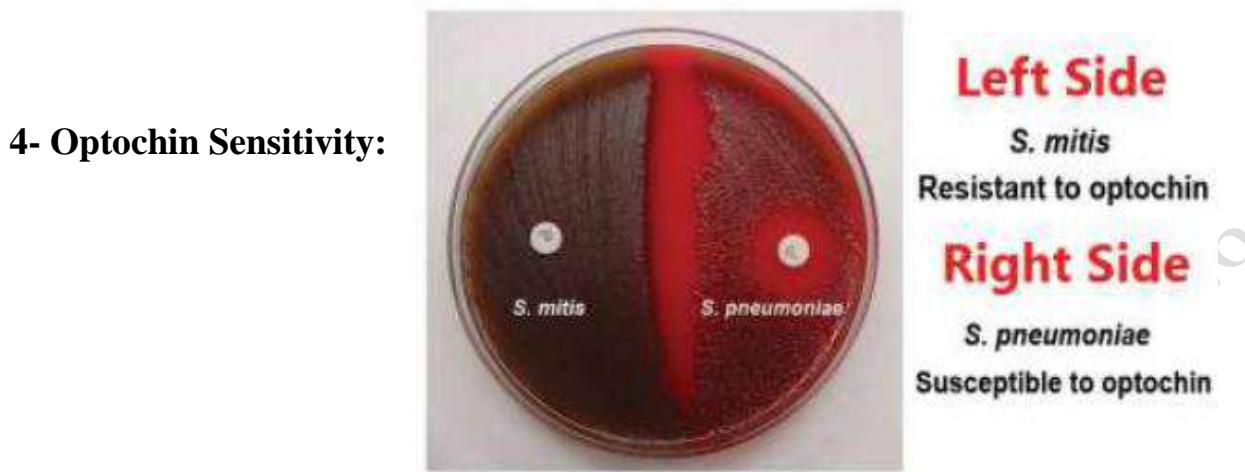


### 2. Bile Solubility Test: Procedure:

- 1- Grow the isolate to be tested for 18 hours at 37°C in 5 ml serum, digest broth or infusion.
- 2- While still warm, add 0.5 ml of 10 percent, bile salt (sodium deoxycholate solution) and re-incubate at 37°C. Pneumococci are lysed within 15 minutes and the initially turbid culture becomes clear and transparent. Pneumococci are soluble in bile; viridans and other streptococci are not.



3. Pneumococci are **Catalase** and **Oxidase** negative.



Comparison between *Streptococcus pneumoniae* and Viridans Streptococci

Features	<i>S. pneumoniae</i>	Viridans streptococci
Colony	Draughtsman	Dome shaped
Bile solubility test	Positive	Negative
Optochin (ethyl hydrocupreine hydrochloride) 5 $\mu$ g sensitivity test	Sensitive	Resistant
Inulin fermentation	Positive	Negative
<b>Pathogenicity</b>	<b>Positive</b>	<b>Negative</b>
Capsule	Present	Absent
Quellung reaction	Positive	Negative

## Laboratory Diagnosis

**1. Specimens:** Sputum, lung aspirate, pleural fluid, cerebrospinal fluid (CSF) or blood are collected according to the site of lesion. Sputum specimens must be mucus from the lungs rather than samples of saliva.

### 2. Microscopy and Antigen Detection

Gram stain of sputum specimens is a rapid way to diagnose pneumococcal disease. If the smears are gram-positive lancet-shaped diplococci, a presumptive diagnosis of pneumococcal pneumonia may be made. A centrifuged deposit of the CSF should be examined immediately in a Gram film in case of meningitis and presumptive diagnosis may be made by finding gram-positive diplococci.

**3. Culture:** Specimen is inoculated on plates of blood agar and heated blood agar (chocolate agar) and incubated in air with 5-10% CO<sub>2</sub> for 18-24 hours.

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### *Pseudomonas aeruginosae*

It is gram negative, motile and rod shaped. It occurs as single bacteria, in pairs, and occasionally in short chains.

**Specimens:** Specimens depend on the site of infection including skin lesions, pus, urine, blood, spinal fluid, sputum, and other material should be obtained by different procedures.

**Culture:** Pseudomonads grow readily on most culture media. It does not ferment lactose and is easily differentiated from the lactose- fermenting bacteria.

*P. aeruginosa* is an obligate aerobe but can grow anaerobically if nitrate is available, that grows readily on many types of culture media, sometimes producing **a sweet or grapelike** or corn taco-like odor. Some strains **hemolyze blood**.

*P aeruginosa* forms smooth round colonies with a **fluorescent greenish color**. It often produces the; 1- non-fluorescent **bluish pigment pyocyanin**, which diffuses into the agar. Many strains produce the 2- **fluorescent pigment pyoverdin**, which gives a greenish color to the agar. Other strains produce the 3- **dark red pigment pyorubinor**, or the 4- **black pigment pyomelanin**.

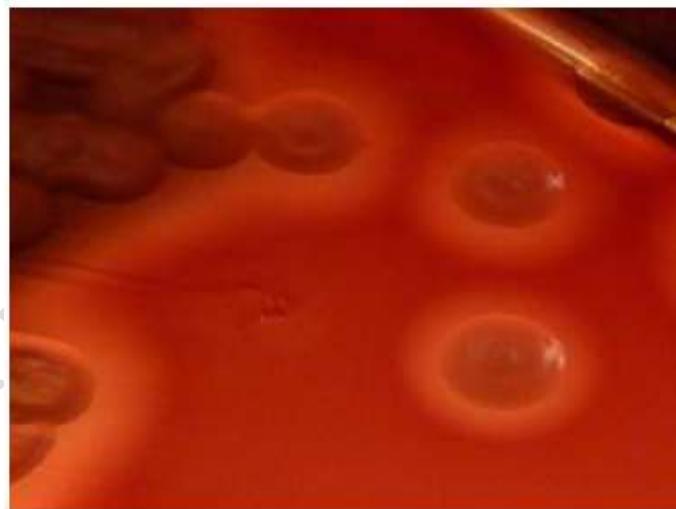
### *Pseudomonas aeruginosa* on Citrimide medium



**On MacConkey agar plates (as shown below)** it produces non-lactose fermenting colonies (**compared with *E. coli* or *Klebsiella***) and the pigments are often poorly observed.



**On blood agar plates**, it is surrounded by a zone of hemolysis (**as shown below**), while in broth culture it forms a dense turbidity with a surface **pellicle**.



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Revised by Prof. Dr. M. I. Mughal