

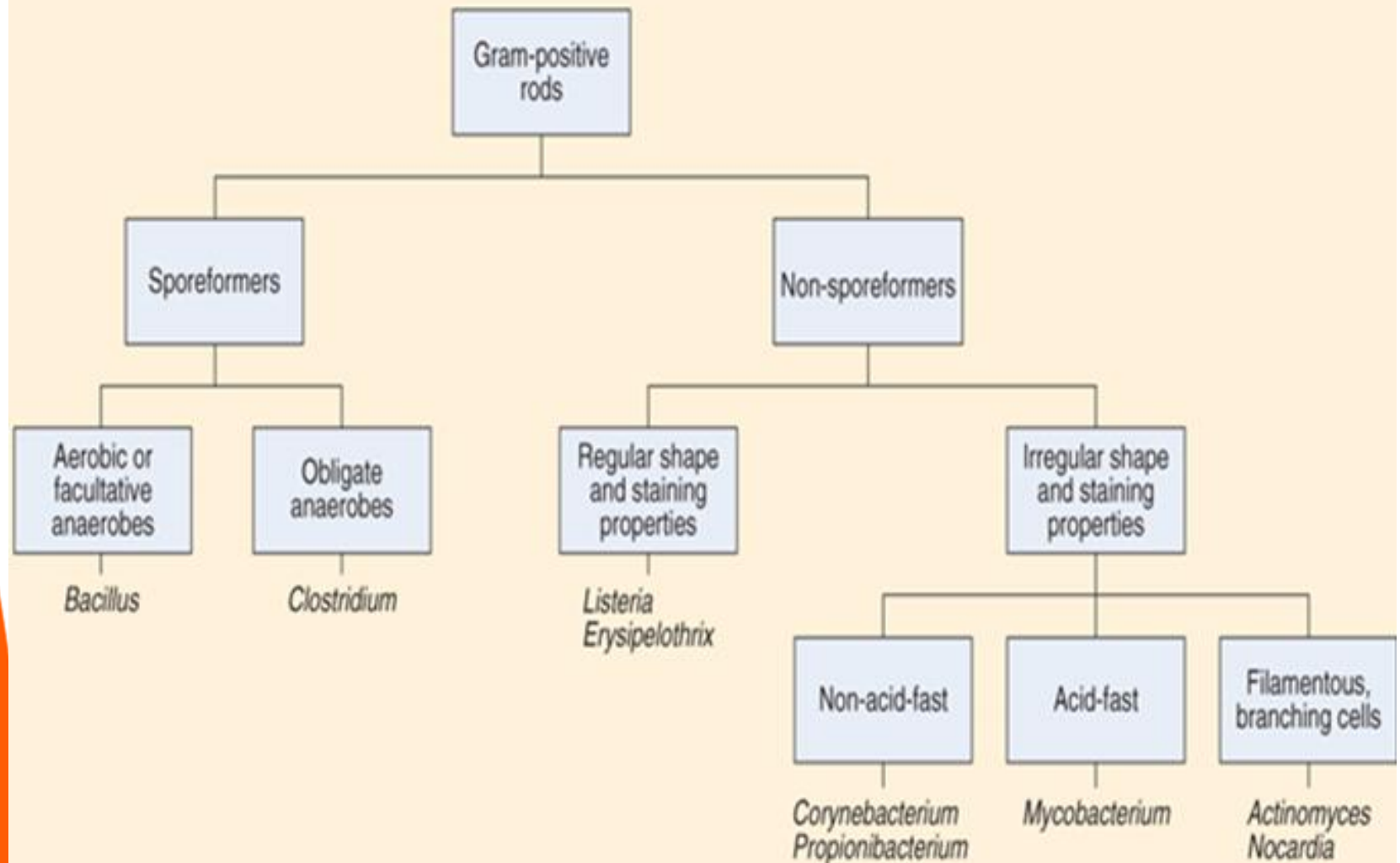
Practical Medical Bacteriology

Lab 6

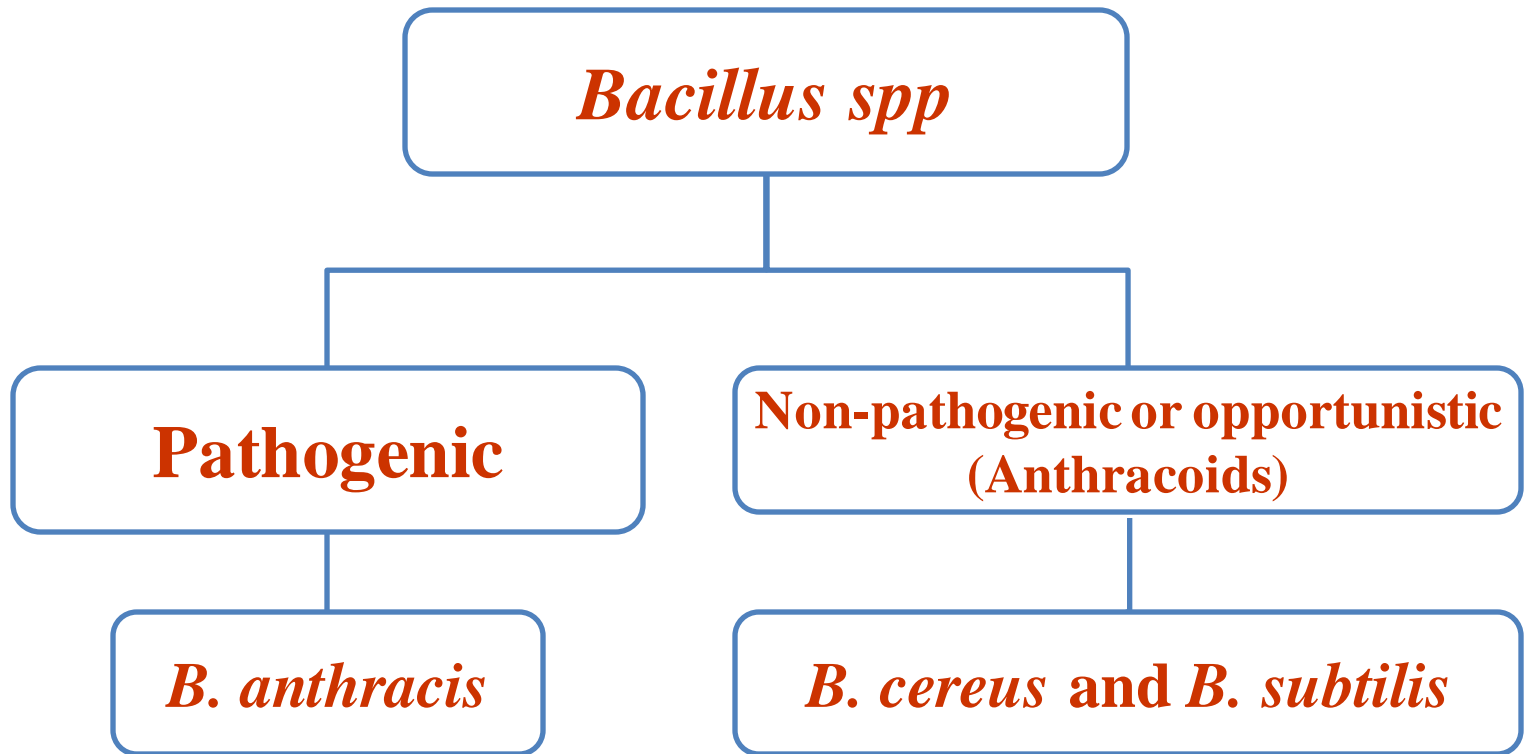
Laboratory Diagnosis of *Bacillus* spp



BLE 19.1 Scheme for Differentiating Gram-Positive Bacilli



Aerobic Spore Forming *Bacillus* spp



General Characters of *Bacillus* spp

- Large Gram positive spore forming bacilli.
- Most are saprophytic in soil, water, and air and called anthracoides such as *Bacillus cereus* and *Bacillus subtilis*.
- *Bacillus anthracis* as a major pathogen.
- All are motile except *B. anthracis*
- Aerobic or facultative anaerobic bacteria
- Catalase positive



Significant *Bacillus* spp

Bacillus anthracis

- large, non-motile, encapsulated, spore-forming Gram-positive bacilli
- Cause anthrax which infect herbivorous animals such as sheep.
- Humans acquire infection by contamination of wound or ingestion or inhalation of spores

Bacillus cereus

- Large, motile, saprophytic bacillus
- Non-capsulated
- A normal inhabitant of soil also isolated from food
- Causes food poisoning

Bacillus subtilis

- Common laboratory contaminant
- Tolerates very high temperatures

Bacillus stearothermophilus

- Used as indicator for efficacy of autoclave



Laboratory diagnosis

1. Specimen:

➤ *Bacillus anthracis*

- Pastular exudates in malignant pustule (skin)
- Sputum
- Stool (stool specimen is emulsified and heating to 80 °C to kill non-spore forming microorganisms).

➤ *Bacillus cereus*

- Normal stool flora, to diagnose food poisoning must culture suspected food NOT **stool**.

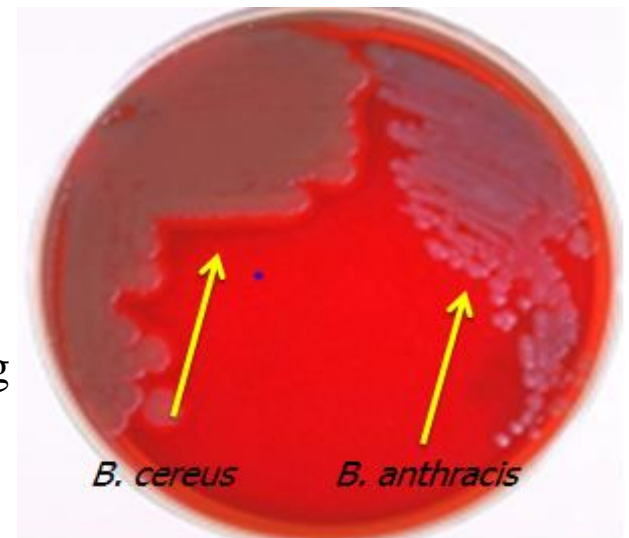


Laboratory diagnosis

2. Cultural characteristics

➤ On blood agar

- *B. anthracis*: Non-hemolytic, raised, large, grayish-white colonies with irregular, fingerlike edges described as beaten egg whites.
- *B. cereus*: β hemolytic; large, feathery, spreading colonies.



➤ On nutrient agar:

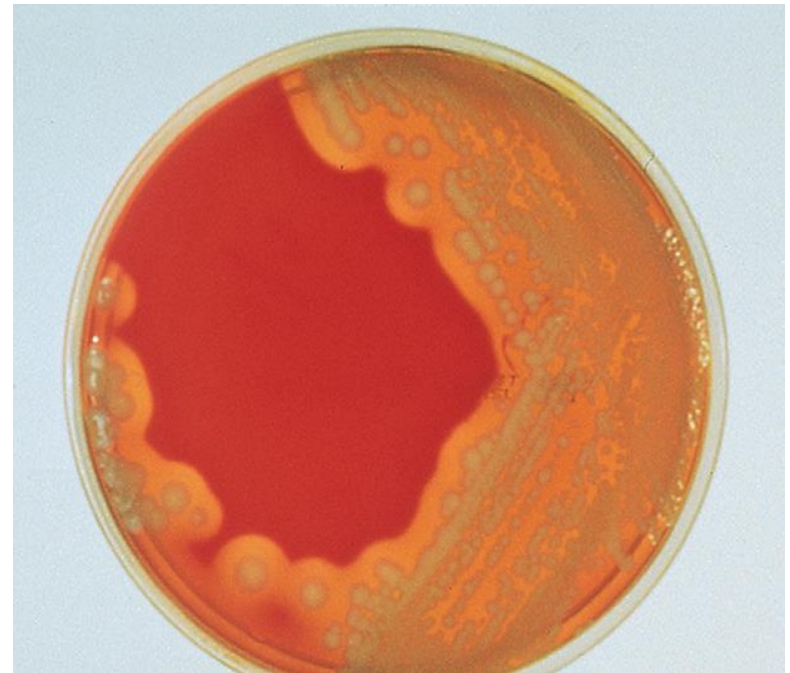
- *Bacillus spp* grow aerobically on nutrient agar at 37°C with characteristic mucoid or smooth colonies, which indicates the pathogenicity of organism (presence of capsule).



Laboratory diagnosis



***B. anthracis* on blood agar**



***B. cereus* on blood agar**

Laboratory diagnosis

3. Microscopical Morphology:

A. Gram stain:

- *Bacillus anthracis*: large, square-ended Gram-positive rods to Gram-variable; may appear end to end giving a "**bamboo appearance**".
- *Bacillus cereus*: large, Gram-positive rods, can stain Gram-variable or Gram-negative.

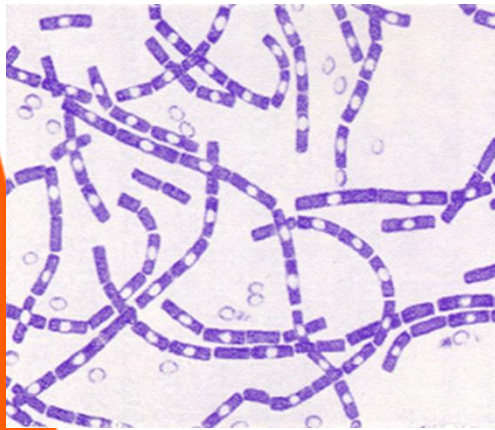
B. Spore Stain:

- *Bacillus spp* spores are oval, central and non-bulging
- By spore staining technique (Malachite green and safranine), the spore appears green while the vegetative cells appear red.

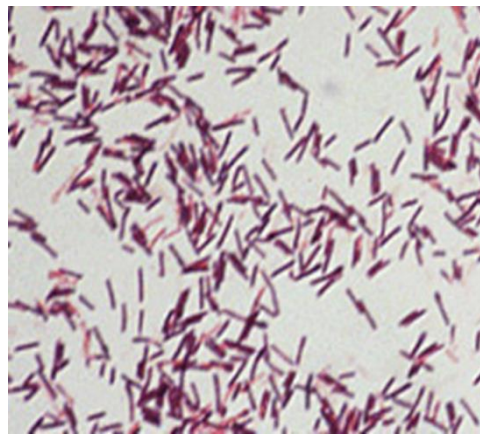


Laboratory diagnosis

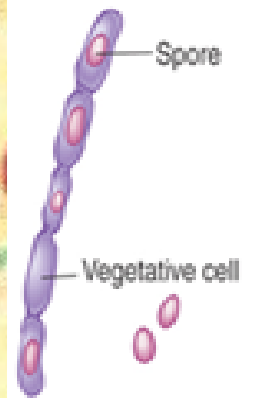
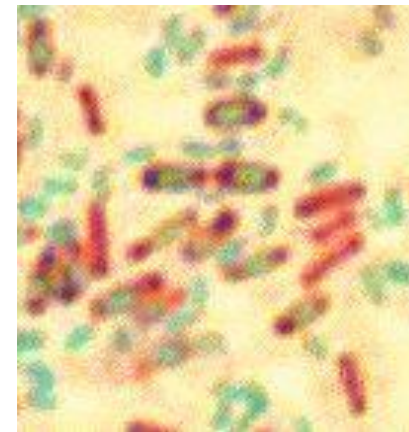
3. Microscopical Morphology:



Gram stain
B. anthracis



Gram stain
B. cereus

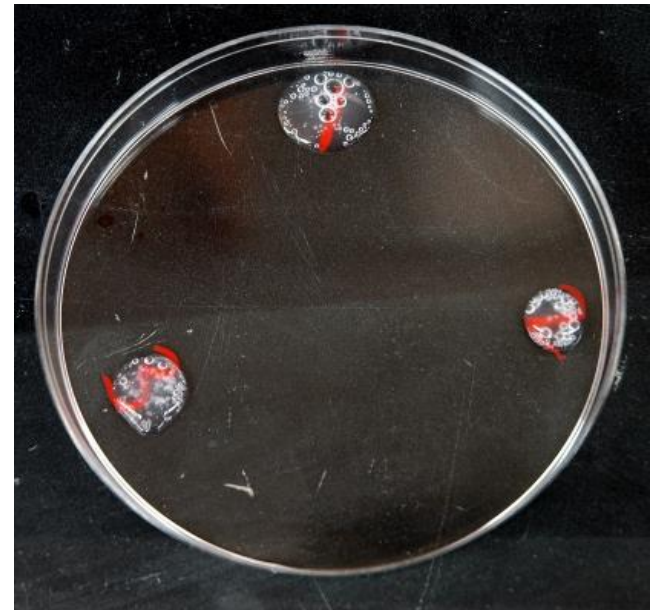
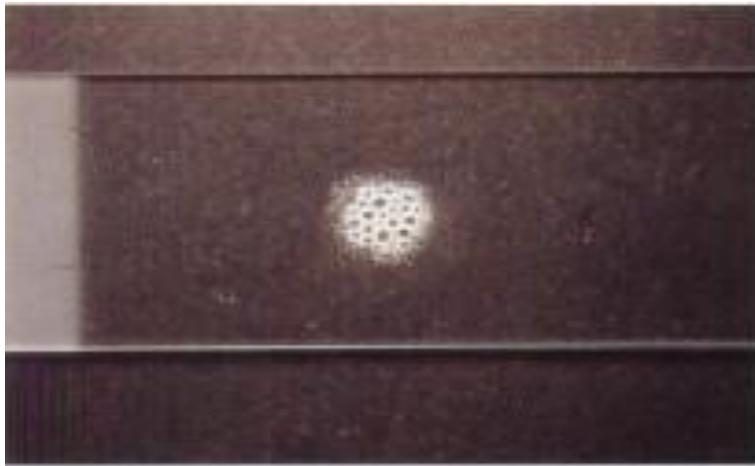


Spore stain

Laboratory diagnosis

4. Preliminary tests:

- **Catalase test:** all *Bacillus* species are catalase positive
- **Motility test:** all *Bacillus* species are motile except *B. anthracis*.



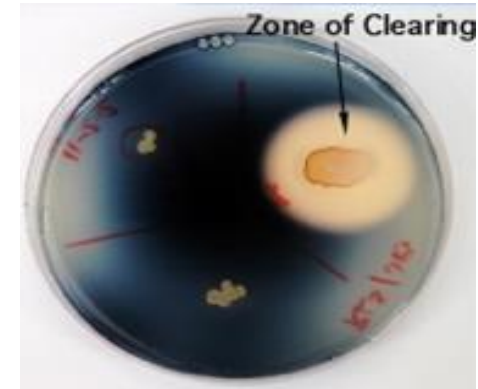
Laboratory diagnosis

5. Biochemical tests:

A. Starch Hydrolysis (Amylase Activity)

Principle:

- Starch + Iodine → blue color
- Glucose + Iodine → No reaction



Nutrient Agar containing Starch + M.O $\xrightarrow{\text{Amylase}}$ Glucose $\xrightarrow{\text{iodine}}$ Appearance of colorless zone around the growth.

Procedure:

1. Inoculate nutrient agar plate containing 1% Starch with the M.O.
2. Incubate the plate at 37 °C for overnight.
3. After incubation, flood the plate with Iodine solution.

Results:

- Activity of amylase is indicated by a clear zone around the growth while the rest of the plate gives blue color after addition of iodine.



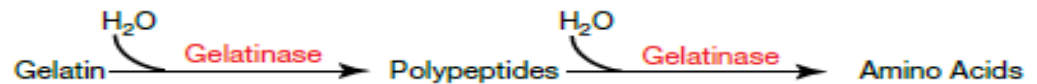
Laboratory diagnosis

5. Biochemical tests:

B. Gelatinase hydrolysis test:

Principle:

Gelatin hydrolysis test is used to determine the ability of a microbe to produce gelatinase as extracellular enzyme that hydrolyze the gelatin.



Procedure:

1. Tubes of nutrient gelatin is stab-inoculating with microorganisms.
2. Incubate the tubes for 7 days at 37° C. 7-days incubation period is usually sufficient to see liquefaction of the medium.



Results:

1. Activity of gelatinase is indicating by a liquefaction medium
2. *B. cereus* is gelatinase-positive while *B. anthracis* is gelatinase-negative



Differential characteristics of *B. anthracis* & *B. cereus*

Tests	<i>B. anthracis</i>	<i>B. cereus</i>
Haemolysis on BA	-	+ (β)
Motility	Non motile	Motile
Starch hydrolysis	+	+
Gelatin hydrolysis	-	+
Penicillin susceptibility	S	R
API 20E system for identification of <i>Bacillus</i> species		

