**Microbial growth**

 **Microbial growth:** defined as an increase in cellular constituents result an increase in a microorganism size, population number or both.

 Usually, growth of bacterial cell characterize via several changes such as total population numbers using different analysis method such as growth curve of microbial culture.

When M.O are cultivated in liquid medium, they are grown in a batch culture or closed system, because no fresh medium is provided during incubation, nutrient concentrations decline and concentrations of wastes are increased.

The growth of microorganisms reproducing by **binary fission** can be plotted as the logarithm of the number of viable cells versus the incubation time, resulting in curve has four distinct phases.

**Bacterial growth curve**

* When a bacterium is added to a suitable liquid medium and incubated, its growth follows a definite course.
* If bacteria counts are made at intervals after inoculation & plotted in relation to time, a growth curve is obtained and shows 4 phases:
	+ **Lag**
	+ **Log or Exponential**
	+ **Stationary**
	+ **Dead or Decline phase**

**Lag phase**

 When M.O are inoculated into fresh culture medium, will not reproduce immediately, therefore this period is called the lag phase. During the lag phase bacterial cell will not divide into new cells in addition there is no net increase in mass, however the cell is synthesizing new components. A lag phase considers a vital period prior to cell division because of the age of the cell, depleted of ATP molecules, essential cofactors and ribosome. This must be synthesized before growth can begin.

The surrounding medium different from the original where the M.O used to grow previously, new enzyme for different nutrient molecules could involve for consumption different nutrients. After all the previous stages cells begin to replicate their DNA, increase in cell mass and finally divide into two new daughters.

 The lag phase varies in length depending on the surrounding conditions; it may be long if the inoculums are from an old culture or one that has been refrigerated. Inoculation of a chemically different medium also results in a longer lag phase, while fresh exponential phase culture when inoculated into a new batch could reduce the length of lag phase which would be short or absent.

**Exponential (log) phase**

 Microorganisms start replicates their number in logarithmic order at maximal rate. During the log phase, the M.O are growing under constant conditions including (nutrient and division rate) causing a uniform identical new cells at regular intervals.

 The population is most uniform in chemical & physiological characteristic therefore exponential phase cells employing in biochemical and physiological studies. Exponential growth is stable therefore; cellular constituents are manufactured at constant rates relatives to each other. The reduction of nutrient levels and O2 variation in environmental conditions resulting in slowing the growth.

**Stationary phase**

 In the stationary phase the total number of viable microorganisms remains constant, this may result from a balance between cell division & cell death, or the population may simply cease to divide though remaining metabolically active. Microbial population enters the stationary phase for several reasons; nutrient limitation, if an essential nutrient is severely depleted, population growth will slow.

 **Population size** depends on **nutrient availability** & other factors, as well as **the type of M.O** being cultured.

 Aerobic organisms are limited by O2 availability, because O2 resulting depleted quickly therefore only the surface of batch culture continue to grow due to presence of the appropriate level of O2. The cells beneath the surface will not be able to grow unless the culture is shaken or aerated in another way. Growth also may cease due to the accumulation of toxic waste products.

 Bacteria in a batch culture may enter stationary phase in response to **starvation**, this occurs in nature environment when nutrient levels are low.

**Death phase**

 Decline in the number of viable cells result from unsuitable conditions, the death of microbial population may be logarithmic like it’s growth during the log phase, the total cell number remains constant because the cells fail to lyses after dying, only way of deciding whether a bacterial cell is viable is by incubating it in fresh medium, if it does not grow and reproduce, it is assumed to be dead. Death is defined to be the irreversible loss of the ability to reproduce.

**Morphological & Physiological alterations during growth**

* Lag phase – maximum cell size towards the end of lag phase.
* Log phase – smaller cells, stain uniformly.
* Stationary phase – irregular staining, sporulation and production of exotoxins & antibiotics.
* Phase of Decline –involution forms (with ageing).



**Microbial growth curve in a closed system, note the 4 phases**

**Potential Importance of the Growth Curve**

* -Implications in microbial control, infection, food microbiology, and culture technology.
* -Growth patterns in microorganisms can account for the stages of infection.
* -Understanding the stages of cell growth is crucial for working with cultures.
* -In some applications, closed batch culturing is inefficient, and instead, must use a chemostat or continuous culture system.

**Generation time (G.T)**

 During the exponential phase each M.O is dividing at constant intervals, thus the population will double in number during a specific length of time called the **generation time or doubling time**, G.T vary with the species of M.O and environmental conditions, it range from less than 10 minutes for several days, and g.t in nature is much longer than in culture.

 Measurement of microbial growth can be determined by:

1. Measurement of cell numbers
2. Measurement of cell mass.

**Types of bacteria with generation times**

* Coliform bacilli like *E.coli* & other medically important bacteria/ 20 mins.
* *Staphylococcus aureus/* 27-30 mins*.*
* *Mycobacterium tuberculosis/ 792-932* mins.
* *Treponema pallidum/ 1980* mins.

**Types of culture systems**

1. **Batch cultures or closed system:** In which nutrient supplies are not renewed nor wastes removed, exponential growth lasts for only a few generations and soon the stationary phase is reached.
2. **Continuous culture system (open system):** Microorganisms are growing in a system with constant environmental conditions maintained through continual provision of nutrients & removal of wastes; these conditions are met in the lab. by a continuous culture system.

 A microbial population can be maintained in the exponential growth phase and at a constant biomass concentration for extended periods in a continuous culture systems, these systems are very useful because they provide a constant supply of cells in exponential phase and growing at a known rate, they make possible the study of microbial growth at very low nutrient levels, these systems are essential for research; in studies on interactions between microbial species under certain environmental conditions, they also used in food & industrial microbiology.

**The Diauxic Growth Curve**

 The process was discovered in *E. coli*. Since glucose is degraded by constitutive enzymes and lactose is initially degraded by inducible enzymes, what would happen if the bacterium was grown in limiting amounts of glucose and lactose. A plot of the bacterial growth rate resulted in a **diauxic growth** **curve** which showed two distinct phases of active growth. During the first phase of exponential growth, the bacteria utilize glucose as a source of energy until all the glucose is exhausted. Then, after a secondary lag phase, the lactose is utilized during a second stage of exponential growth.

**The Diauxic Growth Curve of *E. coli* grown in limiting concentrations of a mixture of glucose and lactose**