# 7.0 : Introduction

# Q.1. What is respiration? Describe different types overspiration.

**Ans:**Definition: Respiration is defined as an intracellular oxidation in which complex organic substances are broken down in step-wise manner with the release of energy which is immediately converted into metabolically usable form of energy, i.e. ATP.

# **Types of respiration :**

i) Aerobic respiration :

When free molecular oxygen is used in respiration to breakdown the organic substrate, the respiration is called aerobic respiration.

It can be represented as follows:

 $\hat{C}_{6}H_{12}O_{6}+6O_{2}+6CO_{2}+6H_{2}O+Energy$ 

# ii) Anaerobic respiration<sup>2</sup>:

When free more used in respiration to breakdown the organic substrate, then it is called an across respiration.

$$O_6H_{12}O_6 \longrightarrow 2C_2H_5OH + 2CO_2 + Energy$$

# Q.2. What are respiratory substrates? Name the most common respiratory substrates.

Ans: The compounds that are oxidized during the process of respiration are known as respiratory substrates. They include carbohydrates, proteins, fats, organic acids, etc. Among these, carbohydrates are the main cource of energy and glucose is the most preferred substrate because it is easily available and acceptable of all kinds of organisms. However, when the carbohydrates are not available, proteins and fats serve as respiratory substrates.

# Q.3. Why is respiration an exergonic and catabolic physio-chemical process ?

**Ans:** A chemical reaction that proceeds with release of energy is called exergonic reaction. Respiration involves the exchange of oxygen and carbon dioxide and the oxidation of glucose inside the mitochondria to produce energy, which is partly stored in the form of ATP molecules. In this reaction, glucose is broken down, hence it is called catabolic process.

# 7.1 : ATP as Currency of Energy

# Q.4. Why is ATP called as energy currency of a cell? Give its significance.

**Ans:**ATP (Adenosine triphosphate) is the energy rich organic compound. Whenever energy is released, ATP synthesis takes place from ADP and Pi, and when energy is required, ATP is hydrolysed and energy released as high energy phosphate bond is broken. This energy is used for metabolic activities in the body. Therefore, ATP is called energy currency of the cell.

# Significance of ATP :

- i) Storage of energy.
- ii) Supply of energy to the cell for metabolic processes.
- iii) It donates one or two phosphate groups and functions as phosphorylating agent.

# Q.5. Write a short note on ATP.

- **Ans:**i) ATP (adenosine tri phosphate) is the biologically usable form of chemical energy. It provides energy for all cellular activities, hence it is called the energy currency of cells.
  - ii) Structurally, the ATP consists of nitrogenous purine base called Adenine, a pentose sugar called Ribose  $(C_5H_{10}O_5)$  and 3 Phosphate molecules.
  - iii) Adenine and ribose sugar together form Adenosine. Adenosine with 1 phosphate, 2 phosphates and 3 phosphates are called AMP, ADY and ATP respectively.
  - iv) Between the 2nd and 3rd molecules of phosphate, a high energy phosphate bond is present. For the formation of these bonds, energy is given by either light (Photophosphorylation) or oxidative process of respiration (oxidative phosphorylation).

- v) Whenever the cell needs energy, ATP is hydrolyzed to produce ADP and Pi (inorganic phosphate) and energy is released as high energy phosphate bond is broken.
- vi) Both, synthesis and breakdown of ATP are regulated by enzymes ATP synthetase and ATPase, respectively.

 $ADP + iP \implies ATP$ 

### Q.6. Describe the structure of ATP.

- Ans:i) Structurally, the ATP consists of nitrogenous purine base called Adenine, a pentose sugar called Ribose  $(C_5H_{10}O_5)$  and 3 Phosphate molecules.
  - ii) Adenine and ribose sugar together form Adenosine Adenosine with 1 phosphate, 2 phosphates and 3 phosphates are called AMP, ADP and ATP respectively.
  - iii) Between the 2<sup>nd</sup> and 3<sup>rd</sup> molecules of phosphate, a **high energy phosphate bond** is present. For the formation of these bonds, energy is given by either light (Photophosphorylation) or oxidative process of respiration (oxidative phosphorylation).



Q.7. More much energy is released when one molecule of ATP is hydrolysed to ADP and Pi ? Any when one molecule of ATP is hydrolysed to ADP and Pi, 7.3 Kcal of energy is released.

Q.8. Describe the ultrastructure of mitochondrion.



- i) Mitochondria are membrane bound cell organelles, essential for aerobic respiration.
- ii) They are present in the eukaryotic cells and absent in prokaryotic cells.
- iii) They are randomly distributed in the cytoplasm.
- iv) Each mitochondrion is covered by two **lipo-protein membranes** which are separated by a narrow, fluid filled space called **outer chamber**.

[Oct 14]

- v) The outer membrane is smooth and continuous, while the inner membrane has a large number of finger-like folds called cristae.
- vi) The space enclosed by the inner membrane is called **inner chamber** and is filled with a colourless fluid called **matrix** which is rich in enzymes and co-enzymes.
- vii) Matrix also contains 70S type of ribosome and single, circular BNA molecule. Thus, mitochondria are self-duplicating, self-autonomous cellorganelles.
- viii) The inner membrane and cristae bear several small particles called **oxysomes** or elementary particles or F<sub>1</sub> particles.
- ix) Each  $F_1$  particle consists of a base, stalk and head.
- x) The spherical head contains enzyme ATP synthetase which catalyzes ATP synthesis.

# Q.9. Give the functions of mitochondria.

# **Ans:** Functions:

- i) The Krebs cycle of respiration takes place in the matrix of mitochondria.
- ii) The oxidative phosphorylation takes place in the F I particles of inner membrane.
- iii) Mitochondria help in the synthesis of ATP.
- iv) These ATP are energy rich compounds and thus mitochondria is called as "Power house of cell".
- v) Synthesis of fatty acids, amino acids takes place in mitochondria.
- vi) It stores and releases calcum.
- vii) It provides intermediates for the synthesis of important biomolecules such as chlorophyll, cytochromes, steroids, etc.

# Q.10."Mitochondria are called power house of cell". Comment.

- **Ans:**i) Oxidative phosphorylation takes place in  $F_1$  particles to generate ATP molecules.
  - ii) About 96 ATP is generated in mitochondria to carry out cellular function.
  - iii) Generated ATP is stored and utilized for active transport, biosynthesis, contraction of muscles and soon. Therefore, mitochondria are called power house of cell.

# 7.2 : Mechanism of Aerobic and Anaerobic Respiration

# Aetohic Respiration

# Q.l1.Describe various steps involved in glycolysis.

Ans: Glycolysis is defined as the stepwise, enzymatic breakdown of hexose sugar (glucose) into two molecules of pyruvate in the cytoplasm. Glycolysis is also known as EMP pathway, as its process was discovered by three German scientists

### Embden, Meyerhof and Parnas.

The EMP pathway is a common pathway for the degradation of glucose in both aerobic and anaerobic respiration.

# Glycolysis consists of two major phases:

- a) Preparatory phase and cleavage.
- b) Oxidative and payoff phase.
- a) Preparatory phase and cleavage :

In this step, glucose molecule is activated by phosphorylation and then cleaved into two molecules of triose phosphates, namely 3-PGAL and DHAP. In the next step, only 3-PGAL participates and hence DHAP gets converted into 3-PGAL. Thus, 2 molecules of 3-PGAL are formed.

It includes the following reactions:

- i) **Phosphorylation I :** A glucose molecule is phosphorylated to glucose 6-phosphate in presence of enzyme hexokinase. In this reaction, ATP supplies the phosphate and gets converted into ADP.
- **ii) Isomerisation :** Glucose-6-phosphate IS isomerised to fructose-6-phosphate in presence of enzyme phosphoglucoisomerase.
- **iii) Phosphorylation II :** Fructose-6-phosphate IS phosphorylated to fructose-l,6-diphosphate. Phosphate is supplied by ATP which gets converted to ADP. Reaction is catalyzed by enzyme phosphofructokinase.
- **iv)** Cleavage: Fructose-l,6-diphosphate splits up into two inter-convertible compounds, i.e. Dihydroxy acetone phosphate (DHAP) and 3-phosphoglyceraldehyde (3-PGAL). Reaction occurs in presence of enzyme aldolase.
- v) **Isomerization :** DHAP is isomerised into 3-PGAL with the help of enzyme phospho-triose isomerase, thus two molecules of PGAL are obtained.

3

#### b. Oxidative and payoff phase :

In this phase, oxidation or removal of hydrogen takes place and then there is ATP generation.

- i) Oxidative Phosphorylation : 3-PGAL undergoes oxidation and phosphorylation to form 1,3diphosphoglyceric acid (1,3-DPGA). Hydrogen released combines with NAD to form NADH<sub>2</sub>. Phosphate is supplied for the reaction by phosphoric  $acid_{1}(H_{3}PO_{4})$ .
- **ii)** ATP generation I I Dephosphorylation I : 1,3-diphosphoglyceric acid is dephosphorylated to 3-phosphoglyceric acid (3-PGA). Phosphate released combines with ADP to form ATP. Reaction takes place in presence of enzyme phosphoglycero-kinase.
- iii) Isomerisation : In presence of enzyme phosphoglycero-mutase, 3-Phosphoglyceric acid IS isomerised to 2-phosphoglyceric acid.
- iv) **Dehydration:** 2-Phosphoglyceric acta loses a water molecule to form phospho enol pyruvic acid. The enzyme enolase catalyses the reaction.
- ATP generation II / Dephosoporylation II : Phosphoenol pyruvic acid is dephosphorylated to pyruvic acid in presence openzyme pyruvate kinase. Phosphate released combines with ADP to form ATP.



## Q.12. Define glycolysis. Give its significance.

**Ans:**Definition: Glycolysis is the process that takes place in the cytoplasm of both eukaryotic and prokaryotic cell in which one molecule of glucose breaks down into two molecules of pyruvic acid.

### Significance of glycolysis :

- i) The net gain of glycolysis is two molecules of ATP, two molecules of NADH2 and two molecules of pyruvic acid per glucose molecule.
- ii) Since each molecule of NADH2 can further give rise to 3 molecules of ATP, each glucose molecule has a net potential of forming 8 ATP (If pyruvic acid enters the mitochondria).

# Q.13.What are the products of cleavage in glycolysic?

Ans:Dihydroxy acetone phosphate (DHAP) and 3-phosphoglyceraldehyde (3-PGAL) are the products of cleavage in glycolysis.

# Q.14.Where does dehydration occur in glyodysis ?

Ans: In glycolysis, dehydration occurs when 2-Phosphoglyceric acid loses a water molecule (dehydration) to form phosphoenol pyruvic acid in presence of enzyme enolase.

# Q.15.What will be the fate of end product of EMP pathway, if free oxygen is not available in the cell?

Ans: If free oxygen is not available, the oxidation will be incomplete and the end product will be  $CO_2$  and ethyl alcohol. Released energy will be less.

# Q.16. Where does glycolysis take place in a cell ?

Ans: Glycolysis takes place in the cytoplasm in a cell.

# Q.17.What is the full form of EMP pathway?

Ans: The full form EMP pathway is Embden-Meyerhof-Parnas pathway.

# Q.18.Describe the connecting link between glycolysis and Krebs cycle.

- **Ans:**i) Keeh pyruvic acid molecule formed in glycolysis undergoes oxidative decarboxylation (i.e. oxidation by dehydrogenation and decarboxylation by loss of  $CO_2$ ), to form Acetyl group (2C), which combines with Co-A to form Acetyl Co-A.
  - Thus, pyruvic acid is converted to Acetyl Co-A. This is called Acetylation of Pyruvate.
  - iii) It involves several reactions, many enzymes and co-factors.
  - iv) Hydrogen released combines with NAD to form reduced co-enzyme  $NADH_2$ , which has the potential to form 3ATP in ETS.
  - v) In this process,  $CO_2$  is released.
  - vi) Acetyl Co-A is a connecting link between glycolysis and Krebs cycle.
  - vii) Summary is represented as :

 $\begin{array}{c} 2PA + 2Co-A + 2 \text{ NAD} \underline{\quad \text{Enzyme Complex (Pyruvic dehydrogenase)}} \\ (3C) \\ \end{array} 2 \text{ Acetyl Co-A} + 2 \text{ CO}_2 \uparrow + 2 \text{ NADH}_2 \\ (2C) \\ \end{array}$ 

### Schematic representation of formation of Acetyl Co-A:



# Q.19. What is TCA cycle? Describe its different steps.

Ans:TCA cycle is a stage of aerobic respiration which takes place in the matrix of the mitochondria. It was discovered by a scientist Sir Hans Krebs, hence called as Krebs cycle. It is also called the TCA cycle, i.e Tricarboxylic acid cycle because the organic acids formed in the cycle contain 3 COOH groups, i.e. carboxylic groups. It is also called the Citric acid cycle because the primary intermediate compound formed is Citric acid.



#### i) Condensation :

Acetyl Co-A (2C) combines with water and Oxaloacetic acid (4C) to form Citric acid (6C). Thus, Citrate/Citric acid is the first stable compound of Krebs cycle. The enzyme citrate synthetase catalyses the reaction..

#### ii) Isomerization :

Citric acid gets converted into its isomer, isocitric acid in presence of enzyme aconitase and Fe<sup>++</sup>. This happens in two steps:

- a) **Dehydration :** Citric acid loses a molecule of water to' form Cis-aconitic acid.
- **b)** Hydration : Cis-aconitic acid forms isocitric acid by the addition of water.

### iii) Oxidation (Dehydrogenation I) :

**Isocitric acid** undergoes oxidation to form **oxalosuccinic acid**. The 2H released combines with NAD to form NADH2.

The enzyme iso-citrate dehydrogenase and Mn<sup>++</sup> are necessary for this reaction.

### iv. Decarboxylation I: Oxalosuccinic acid (6C) is decarboxylated in presence of enzyme decarboxylase to form $\alpha$

Respiration
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ketoglutaric acid (5C). CO, goes out of the mitochondria.

Oxidative decarboxylation (Dehydrogenation II and Decarboxylation II) : v)

 $\alpha$  - ketoglutaric add (5C) undergoes oxidative decarboxylation in the presence of Co-A and forms succinyl Co-A (4C). 2H released combine with NAD to form NADH, Enzyme a-ketoglutarate dehydrogenase catalyses this reaction.

- vi) Hydration and Phosphorylation : Succinyl Co-A loses Co-A and gains a H<sub>2</sub>O molecule to form Succinic acid. During the reaction, GDP forms GTP. GTP reacts with ADP to produce ATP and GDP. The reaction takes place in presence of enzyme succinate thiokinase.
- vii) Oxidation (Dehydrogenation III) : Succinic acid is oxidised by dehydrogenation to form Fumaric acid. The 2H released combines with FAD to form FADH2. The reaction is catalysed by enzyme succinate dehydrogenase.

# viii) Hydration :

Fumaric acid takes a molecule of water and forms Malic acid. This occurs in presence of enzyme Fumarase.

ix) Oxidation (Dehydrogenation IV) : Malic acid undergoes ordation to form oxaloacetic acid. The 2H released combines with NAD to form NADH<sub>2</sub>. The reaction is catalysed by enzyme malate dehydrogenase. [Note : GTP is Guardsine Triphosphate. FAD is Flavin Adenine Dinucleotide]

# Q.20.What is the site of Krebs cycle in mitochondrion ?

Ans: Krebs cycle takes place in the mitochondrial matrix.

Q.21.Name the reaction when a-ketoglutarate is converted to succinyl Co-A in Krebs cycle. [Oct 2013] Ans: During Oxidative decarboxylation, a-ketoglutarate is converted to succinyl Co-A in Krebs cycle.

# Q.22.Why is acetyl Co-A called connecting link between glycolysis and Krebs cycle ?

Ans:i) Sormation of acetyl Co-A from pyruvic acid is called connecting link between glycolysis and Krebs cycle. End product of glycolysis undergoes the process of Acetylation to form acetyl Co-A.

Pyruvic is first decarboxylated by removal of  $CO_2$  and then it is oxidised by removal of  $H_2$  and gets converted into acetyl Co-A.

# Q.23. Give the significance of Krebs cycle.

# Ans: Sigmflcance :

- i) It is a common pathway for break down of carbohydrates and fats.
- ii) It forms reduced co-enzymes like NADH, and FADH,
- iii) Krebs cycle has the potential to produce 24 ATP.
- iv) The CO<sub>2</sub> which is liberated is useful in photosynthesis.
- It forms a number of intermediate products which serves as building blocks for synthesis of other v) complex organic compounds.

# **Q.24.** What are the main steps in aerobic respiration? Where does it take place ?

Ans: It occurs in most of the plant and animals, inside their mitochondria in the presence of molecular oxygen.

- It is divided into four major steps:
- i) Glycolysis
- ii) Oxidative decarboxylation or connecting link or acetylation.
- Krebs or TCA cycle iii)
- iv) Electron Transport System (ETS)

# Q.25. Discuss "The respiratory pathway is an amphibolic pathway".

- In respiration, various respiratory substrates are broken down to release energy by the process of Ans:i) cellular respiration. This is called catabolic pathway or catabolism.
  - ii) However, the breakdown of respiratory substrates provides carbon skeleton for synthesis of various essential plant products like polysaccharides, proteins, fats, nucleic acids, pigments, cytochromes, etc.
  - Thus, same respiratory process acts as catabolic as well as anabolic pathway for synthesis of various iii) intermediate metabolic products. Hence, it is called amphibolic pathway.
  - Some important catabolic processes are as follows : iv)
    - Fats are broken down into glycerol and fatty acid. a)
    - b) Fatty acids would enter the pathway only when degraded to acetyl CoA.
    - Glycerol would enter the pathway only after being converted to PGAL. c)

- d. Proteins get degraded into proteases which after deamination get converted into amino acids.
- v. Many substrates are withdrawn from the pathway for formation of many substrates. Some anabolic processes are as follows:
  - a. Acetyl CoA is withdrawn for formation offatty acids.
  - b. a-ketoglutaric acid is converted into glutamate and there is converted into proteins.
  - c. Oxalo acetic acid is converted into aspartate and then into proteins.
  - d. Glutamate can also form pigments like chlorophylls, phycocyanins, enzymes like catalase, cytochromes, etc.

# Q.26. Explain the mechanism of electron transportation system (ETS) and give its significance.

# [Mar 2013 Old Course]

# Ans:Electron Transport System! Terminal oxidation :

The oxidation of reduced coenzymes into the oxidized forms is known as **terminal oxidation**. This process involves electron transfer through various electron carriers which form electron transport system or respiratory chain. During this process, ATP formation takes place which is called as **oxidative phosphorylation**.

The hydrogen released from the organic compounds during glycolysis and Krebs cycle is accepted by hydrogen acceptors or coenzymes, mostly NAD and sometimes FAD. Thus, NAD and FAD get reduced to form NADH<sub>2</sub> and FADH, respectively. Neither NADH<sub>2</sub> nor FADH2 can donate their electrons directly to  $O_2$  to form  $H_2O$ .

# Mechanism :

- i) NAD<sup>+</sup> accepts hydrogen from an organic compound and gets reduced to NADH<sub>2</sub>.
- ii) The NADH, donates hydrogen to FMN due to which it gets reduced to FMNH<sub>2</sub>. Energy released during his oxidation reduction step, is utilized in the synthesis of a ATP molecule.
- iii) FMNH, further donates hydrogen to Co-enzyme Q (Ubiquinone) and get oxidised to FMN.
- iv) co-enzyme Q. gets oxidised by releasing hydrogen. The released H, splits into protons (2H<sup>+</sup>) and electrons (2e<sup>-</sup>).
- This electron pair (2e<sup>-</sup>) is transferred through cytochrome  $b_{,c_1}c$  and  $a_{,a_3}$  in a sequence. The protons (2H<sup>+</sup>) remain in the matrix.
- vi) Since cytochromes are iron compounds, they are alternatively reduced and oxidized.
- vii) On the acceptance of electrons, ferric ( $Fe^{+++}$ ) is reduced to ferrous ( $Fe^{++}$ ) and on donating the electrons, ferrous ( $Fe^{+++}$ ) is oxidized back to ferric ( $Fe^{+++}$ ).
- viii) During the electron transfer through the cytochromes, energy is released, which is used to synthesize ATP from ADP and Pi.
- ix) The second ATP is formed between cytochrome band  $C_1$ . The third ATP IS formed between cytochrome a and a3.
- x) Therefore, oxidation of NADH<sub>2</sub> produces 3 ATPs.
   When the electrons are transferred from cyt. a<sub>3</sub>, they combine with oxygen and make it ionized.
   When FADH<sub>2</sub> is reoxidised through ETS, two molecules of ATP are formed.

# Significance of ETS :

- i) In this process, energy is released in a controlled and stepwise manner to prevent any damage to the cell.
- ii) By the end of ETS,  $H_2$  is added to molecular oxygen to form water.
- iii) It indicates that each  $H_2$  or electron releases 3 molecules of ATP. In this process, most. of the energy is released, trapped and conserved as ATP.
- iv) Thus, per glucose molecule, 38 ATP molecules are formed.



## Q.27. In which part of the mitochondria is the electron transport chain located ?

Ans: The electron transport chain is located on the cristae of inner mitochondrial membrane.

## Q.28.What is oxidative phosphorylation ?

Ans:Oxidative phosphorylation: this a metabolic pathway that uses energy released by the oxidation of substrates to produce ATP.

- i) Oxidative phosphorylation takes place in the mitochondrial membranes.
- ii) Many interm diate products during respiration are oxidised and release 2H.
- iii) They are trapped by NAD or FAD.
- iv) Such acceptors pass through electron transport system to produce ATP and H<sub>2</sub>O.

## Q.29. What is the significance of step-wise release of energy in respiration ?

Ans:i) Evergy is released in steps and in small amounts, so it prevents sudden increase in body temperature.ii) Wastage of energy is avoided as more time is available to use and store the energy in ATP molecules.

- Energy produced can be regulated according to the needs of the body.
- iv) Intermediate products of cell-respiration can be used in different metabolic pathway. e.g. Acetyl Co-A (in the formation of fatty acids, cutin and isoprenoids);  $\alpha$  -ketoglutaric acid (in the formation of glutamic acid); Oxaloacetic acid (in the formation of aspartic acid, pyrimidines and alkaloids); Succinyl Co-A (synthesis of pyrrol compound of chlorophyll).
- v) Krebs cycle is a common pathway of oxidative breakdown of carbohydrates, fatty acids arid amino acids.

### Q.30. Give the balance sheet of ATP formed in aerobic respiration.

#### Ans: 1. In Glycolysis

	a)	ATI	Ps formed (By s	ubstrate leve	el phosphoryla	tion)	
		i)	1,3-diPGA to	3PGA	2ATPs		
		ii)	PEPA to PA		2ATPs		
				Total	4ATPs		
	b)	ATI	Ps used				
		i)	Glucose to Gl	ucose-6 - PO	$D_4$	1 A	ГР
		ii)	Fructose-6 PC	$D_4$ to Fructos	se-I,6-diPO <sub>4</sub>	1 A	ГР
				Total		2 A	TPs
2.	Kre	ebs (	Cycle				
	ATF	s for	med: (By substi	ate level pho	osphorylation)		
	Suc	cinyl	CoA to Succina	ate	$1 \times 2$		
				Total	2ATPs		
3.	Terr	ninal	Oxidation (thro	ough ETS)			
	i)	Fro	m NADH,: 10N	$\text{NADH}_2 \times 3A$	АТР	30 A	ATPs
	ii)	Fro	m FADH, : 2FA	$ADH_{2} \times 2AT$	ГР	4A7	ſPs
			2	Total		<b>34</b> A	ATPs
		Tota	al ATPs formed	(4+2+34)	)	=	40 ATPs
		Tota	al ATPs used			=	2ATPs
		Net	gain			=	38A TPs

# Q.31.What is meant by the statement "aerobic respiration is more efficient"?

**Ans:** Aerobic respiration leads to complete oxidation of organic substances in presence of  $O_2$  and releases water,  $CO_2$  and large amount of energy. In anaerobic respiration, incomplete oxidation takes place which leads to release ofless amount of energy. Hence, aerobic respiration is more efficient. Anaerobic Respiration

# Q.32.Define anaerobic respiration.Which organisms respire apaerobically ?'

Ans: When free molecular oxygen is not used in respiration to break down the organic substrate, then it is called anaerobic respiration.

OR

Incomplete oxidation of the respiratory substrate in the absence of oxygen to yield CO2 and ethyl alcohol is called anaerobic respiration. Anaerobic respiration takes place in bacteria, yeast and many other microorganisms.

# Q.33.Describe the mechanism of anaerobic respiration.

Ans: Process of anaerobic respiration:

- i) A Glucose molecule (6C) is broken down into 2 molecules of pyruvic acid (3C) by the process of glycolysis.
- ii) Each pyruvic acid (3C) molecule then undergoes decarboxylation, i.e. removal of a  $CO_2$  molecule and forms an acetaldehyde (2C) molecule.
- iii) An **acetaldehyde** notecule undergoes reduction to form a molecule of ethyl alcohol. NADH<sub>2</sub> provides hydrogen for reduction and gets oxidized to NAD.

# Steps involved in anaerobic respiration :

i) Glycolysis :

Glycolysis or EMP pathway is an anaerobic process and occurs in cytoplasm. During glycolysis, each glycose molecule is broken down into two molecules of pyruvate with the formation of two molecules each of NADH2 and ATP.

Glucose + 2NAD + 2ADP + 2iP  $\longrightarrow$  2 pyruvic acid + 2NADH2 + 2ATP

# **(**6C)

# ii. Decarboxylation :

Pyruvate undergoes decarboxylation by removal of a CO2 molecule. It occurs in presence of enzyme pyruvate decarboxylase to give acetaldehyde.

Pyruvic acid  $\longrightarrow$  Acetaldehyde (3C) (2C)

# iii) Reduction :

Acetaldehyde reduces to ethyl alcohol by the reduced Coenzyme NADH2 in presence of enzyme dehydrogenase. NADH2 is oxidised to NAD.

Acetaldehyde  $NADH_2$  NAD Ethyl alcohol (2C) (2C)

Q.34. Name the process which is common to both aerobic and anaerobic respiration.

Ans: Glycolysis is common to both aerobic and anaerobic respiration.

# **Additional Information**

# Q.35. Why do some higher organisms including humans sometimes respire anaerobically? How ?

- **Ans:**i) There are some plants and animals, which can use anaerobic respiration also, but only for a short period of time.
  - ii. Humans performs aerobic respiration, but in skeletal muscles anaerobic respiration takes place during exercise because more energy is needed and the O<sub>2</sub> is insufficient for cellular respiration.
  - iii) The pyruvic acid gets reduced to lactic acid by enzyme lactate dehydrogenase.
  - iv) In plants, sucrose is converted into fructose and glucose by invertase enzyme. Glucose then enters glycolysis.
  - v) End product of glycolysis is pyruvic acid that may not get sufficient  $O_2$  in higher plants which leads to anaerobic respiration.

# Q.36.Why is less energy produced during anaerobic respiration ?

Ans: Anaerobic respiration produces less energy because:

- i) Incomplete breakdown of respiratory substrate takes place.
- ii) Some of the products of anaerobic respiration can be oxidised further to release energy which shows that anaerobic respiration does not liberate the whole energy contained in the respiratory substrate.
- iii)  $NADH_2$  does not produce ATP, as electron transport is absent.

# Q.37. Explain fermentation. Give its significance.

# Ans: Fermentation (Greek word: fervos = bubbling)

- i) It is the process of anaerobic breakdown (oxidation) of an extracellular organic substrate by the action of enzymes secreted by micro-organisms (which respire anaerobically) resulting in the formation of some important products.
- ii) However, substrate breakdown (oxication) in fermentation is extracellular, while in respiration it is always intracellular.

# Mechanism :

It involves glycolysis in which one molecule of glucose (6C) is degraded to two molecules of pyruvic acid (3C), two ATP molecules and two NADH2 molecules. Depending upon the metabolic pathways, the end products of anaerobic respiration are either ethanol and carbon dioxide (as in yeast) or lactic acid (as in animals and some bacteria).

# Significance :

- i) Production of alcohol by fermentation is used in brewing industry for production of whisky, beer and other types of wines.
- ii) CO released during fermentation is used for making bread light in baking industry.
- iii) **Production** of vinegar by acetic acid bacteria.
- iv In milk industry, for production of yoghurt, cheese, etc.
- Synthesis of organic acids like citric acid and malic acid during fermentation by fungi like *Aspergillus, Penicillium and Rhizopus.*
- vi) Tea and tobacco leaves are cured by fermentation with certain bacteria.
- vii) Antibiotics and vitamins are also produced by fermentation.

# Q.38. What are the two types of fermentation ?

**Ans:** Fermentation is mainly of two types:

# i) Alcoholic fermentation :

This process is involved in brewing industry for producing beverages like beer, rum, whisky, etc. It is done by brewing yeast (Saccharomyces cerevisiae). In alcoholic fermentation, sucrose (molasses) is converted into alcohol. Yeast or a group of enzymes collectively called zymase can bring about the alcoholic fermentation of sucrose. The first step is hydrolysis, in which sucrose is hydrolysed into glucose and fructose as follows:

 $C_{12}H_{22}O_{11} + H_2O \rightarrow C_6H_{12}O_6 + C_6H_{12}O_6$ 

Second step is glycolysis followed by decarboxylation and reduction as in anaerobic respiration.

# ii) Lactic acid fermentation :

In case of lactic acid fermentation, milk sugar lactose is converted into lactic acid. The first step is hydrolysis of lactose (disaccharide) into glucose and galactose which is followed by glycolysis and reduction (There is no decarboxylation). Thus, fermentation is similar to anaerobic respiration, but it can be extracellular or intracellular.

# Q.39. Name the process of respiration which does not involve intake of oxygen (02) and release of carbon dioxide. (CO<sub>2</sub>). [Mar 2014]

**Ans:**Lactic acid fermentation is the process of anaerobic respiration which, does not involve intake of oxygen  $(O_2)$  and release of carbon dioxide  $(CO_2)$ ,

# Q.40.Name the products of lactic acid fermentation.

Ans: Lactic acid is the product of lactic acid fermentation.

# Q.41. Distinguish between the following :

i) Glycolysis and Krebs cycle.

#### Ans:

No.	Glycolysis / EMP pathway	Krebs cycle / TCA cycle / Circle acid cycle	
a)	Glycolysis is common in both aerobic and	Kerbs cycle occurs only in aerobic	
	anaerobic respiration.	respiration.	
b)	It takes place in the cytoplasm	It takes place in the mitochondria	
c)	$CO_2$ is not released	$CO_2$ is released.	
d)	Total amount of energy produced = 8 ATP	Total amount of energy produced = 24 ATP	
e)	It is linear pathway.	It is cyclic pathway.	
f)	Pyruvic acid is the end product	$CO_2$ and $H_2O$ are the end products.	

# ii) Aerobic respiration and anaerobic respiration.

# [Sept 2009]

[Oct 2013]

Ans:

No.	Aerobic respiration	Anaerobic respiration
a)	It takes place in higher organisms.	It takes place in lower organisms.
b)	It takes place in cytoplasm and mitochondria.	It takes place in cytoplasm.
c)	It involves the participation of free molecular	It does not involve participation of free
	oxygen.	molecular oxygen.
d)	Oxidation of food is complete.	Oxidation of food is incomplete.
e)	It produces $CO_2$ and $H_2O$ .	It produces $CO_2$ and $C_2H_5OH$ .
f)	Tox'c products are not formed.	Toxic products are formed.
Ð	It releases more energy, i.e. 38 ATP.	It releases less energy, i.e. 2 ATP.
h)	Overall equation :	Overall equation :
	$C_6H_{12}O_6+6O_2 \rightarrow 6CO_2 + 6H_2O + Energy$	$C_6H_{12}O_6 \rightarrow 2C_2H_5OH + 2CO_2 + Energy$

# iii) Photosynthesis and Aerobic respiration.

Ans:

No.	Photosynthesis	Aerobic respiration
a.	It takes place In the cells containing	It takes place in all living cells of higher
	chloroplasts.	organisms.
b.	It occurs in chloroplast.	It occurs in cytoplasm and mitochondria.
c.	It is an energy trapping process.	It is an energy releasing process.
d.	It is an anabolic process.	It is a catabolic process.
e.	This process requires $CO_2$ and $H_2O$ .	This process requires sugar and $O_2$ .
f.	Light is necessary for photosynthesis.	Light is not necessary for aerobic respiration.
g.	End products are carbohydrates and oxygen.	End products are CO2, H20 and energy.
h.	Overall equation :	Overall equation :
	$6\mathrm{CO}_2 + 12\mathrm{H}_2\mathrm{O} \rightarrow \mathrm{C}_6\mathrm{H}_{12}\mathrm{O}_6 + 6\mathrm{H}_2\mathrm{O} + 6\mathrm{O}_2\uparrow$	$C_6H_{12}O_6+6O_2 \rightarrow 6H_2O + 6CO_2 + Energy$

## iv. Chloroplast and Mitochondria.

#### Ans:

No.	Chloroplast	Mitochondria
a. Structural and functional unit of photosynthesis.		Structural and functional unit of aerobic respiration
b. Found in plant cells. Found in all eukaryotic cells, i.e. plants and animals.		Found in all eukaryotic cells, i.e. plants and animals.
c.	It is a energy trapping unit.	It is a energy releasing unit, i.e. power house of cell.
d.	It releases $O_2$ and absorbs $CO_2$ .	It absorbs $O_2$ and releases $CO_2$
e.	In this, organic food is synthesized. In this, organic food is broken down.	

### v) Glycolysis and Fermentation.

No.	Glycolysis	Fermentation	
a.	Works both in aerobic and anaerobic	Works only in anaerobic respiration	
	respiration.	$\frown$	
b.	2 molecules of NADH <sub>2</sub> are produced per	NADH <sub>2</sub> is not formed.	
	glucose molecule.		
c.	ATP is produced.	ATPris not produced.	
d.	End product is Pyruvic acid	End product is ethyl alcohol / lactic acid/ acetic	
		acid.	

# vi) Oxidative phosphorylation and Photophosphorylation.

Ans:

Ans:

No.	Oxidative phosphorylation	Photophosphorylation			
a.	It occurs during respiration	It occurs during photosynthesis.			
b.	It takes place in $F_1$ particle of mitochondria.	It takes place in grana of chloroplast.			
c.	It is dependent on Q2	It is independent of O2.			
d.	Released hvdrozen is accepted by NAD and FAD.	Released hydrogen is accepted by NADP.			

# vii) Respiration and Combustion.

Ans			
Ans.	No.	Respiration	Combustion
	a.	Biochemical and stepped process.	Physiochemical and spontaneous process.
	b	Occurs inside the cells.	It is a non-cellular process.
	N	Energy is released in steps.	Large amount of energy is released at a time.
5	d.	No light is produced.	Light may be produced.
	e.	It is controlled by enzymes.	It is not controlled by enzymes.
	f.	A number of intermediates are produced.	No intermediates are produced.

### Exchange of gases

# Q.42. How does pJant anatomy help in exchange of gases ?

**Ans:**Plants are well adopted for gaseous exchange as during photosynthesis, large amount of gases are exchanged. In plants, each living cell is located quite close to the surface of the plants. Thus, when cells photosynthesize, availability of  $O_2$  is not a problem. Stems also posses openings called lenticels. The loose packing of parenchyma cells in stems and roots provide interconnected network of air spaces and transport of gases is facilitated.

# 7.3: Respiratory Quotient

### Q.43. What is R.Q.? Write down R.Q. for different nutrients.

**Ans:**i) Respiratory quotient (R.Q.) is the ratio of volume of  $CO_2$  given out to the volume of oxygen consumed in respiration in the given period of time at standard temperature and pressure.

$$\mathbf{P}_{\mathbf{O}}$$
 Volume of  $\mathrm{CO}_2$  evolved

ii) R.Q.  $\overline{\text{Volume of } O_2 \text{consumed}}$ 

- iii) The value of respiratory quotient depends upon the nature of respiratory substrate used during respiration. The value varies as the substrate changes. RQ. value for different nutrients is as follows:
  - a) Carbohydrates (R.Q. is 1)

When carbohydrates are used as substrate, equal volumes of  $CO_2$  and  $O_2$  are evolved and consumed respectively, thus its R.Q. is 1.

 $C_6 H_{12}O_6 + 6O_2 \rightarrow 6 CO_2 + 6H_2O$ 

R.O. 
$$=\frac{6CO_2}{6O_2}=1$$

### b) Fats (R.Q. is less than 1)

Substrates like fats are poorer in oxygen than carbohydrates. Thus, more oxygen is utilized for its complete oxidation.

$$2(C_{51}H_{98}O_6) + 145O_2 \rightarrow 102CO_2 + 98H_2O$$
  
R.O. =  $\frac{CO_2}{O_2} = \frac{102}{145} = 0.7$ 

# c. Protein respiration (R.Q. is less than 1)

When proteins serve as respiratory substrate, they are first degraded to amino acids. Then, amino acids are converted into various intermediates of carbohydrates. However, amino acids have low proportion of O2 as compared to carbohydrates. Thus, they require more O2 during their complete oxidation and value of RQ. becomes less than 1. In case of proteins, the RQ. is approximately 0.5.

d. Anaerobic respiration (RQ is infinity)

In anaerobic respiration, CO2 is evolved but O2 is not consumed, therefore RQ. is infinite. C.H.,  $O_{c} \rightarrow 2C.H.OH + 2CO_{c}$ 

$$R.O = \frac{CO_2}{O_2} = \frac{CO_2}{O_2} = Infinity$$

# 7.4 : Significance of Respiration

## Q.44. Give the significance of respiration.

- Ans:i) It converts chemical energy into metabolically usable form of energy, i.e. ATP.
  - ii) Along with photosynthesis, it helps to maintain the carbon cycle in nature, i.e. CO2 released during the respiration is used in photosynthesis.
  - ii) Anaerobic respiration or fermentation is applied in the preparation of a number of commercially
  - important products like alcohol, antibiotics and vitamins.
  - iv) The intermediate products formed during respiration are also used in the synthesis of organic compounds like fats and proteins.
  - v) Respiration provides source of energy for cell division, growth and repair, replacement of worn out parts, movements, locomotion, etc.

# **Additional Theory Questions**

- Q.1. Define respiration. Write the overall reaction of aerobic and anaerobic respiration. Refer Q.1.
- Q.2. What is respiration ? Give the overall equations of aerobic and anaerobic respiration. Refer Q.1.
- Q.3. Draw a neat and labelled diagram showing ultrastructure of mitochondrion. [Mar 2013 Old Course]

[Oct 2013, Mar 2014] Refer Q.8.

- Q.4. Describe briefly the EMP pathway of respiration. Refer Q.11.
- Q.5. Explain the common pathway for aerobic and anaerobic respiration. Refer Q.11.

Q.6. Describe the steps involved in formation of pyruvic acid from glucose molecule. Refer Q.11.

Q.7. Give the schematic representation of EMP pathway. Refer Q.11.

Q.8. Give the schematic representation of glycolysis. Refer Q.11.

Q.9. Describe the formation of Acetyl Co-A in respiration. Refer Q.18.

Q.10.Describe Krebs cycle. Refer Q.19.

Q.11.Describe Citric acid cycle. Refer Q.19.

Q.12. Give the schematic representation of an overall view of Krebs cycle. Refer Q.19.

Q.13. Give an account of ATP generation steps during ETS. Refer Q.26.

Q.14.Illustrate the mechanism of electron transport system. Refer Q.26.

Q.15.Explain ETS. Refer Q.26.

- Q.16. Explain terminal oxidation. Refer Q.26.
- Q.17. Give an account of net gain of ATP during aerobic respiration. Refer Q.30.
- Q.18. Explain the process of anaerobic respiration. Refer Q.33.
- Q.19.What is fermentation?

[Oct 2014] Refer Q.37.

- Q.20.Distinguish between Glycolysis and Citric acid cycle. Refer Q.41. (i)
- Q.21. Define R.Q. What is its value for fats? Refer Q.43 (i) and (b).
- Q.22.What is respiratory quotient (RQ)? Why RQ in an aerobic respiration is infinite? [Mar 2013] ARNING Refer Q.43
  - (i, ii, iii) and (d).

# **Quick Review**

*				
No.	Steps(s)	Substrate	Enzyme	End product(s)
i)	Phosphorylation	Glucose + ADP	Hexokinase	Glucose – 6 - Phosphate + ADP
ii)	Isomerisation	Glucose - 6 - Phosphate	Phosphoglucoisomerase	Fructose – 6 – Phosphate
iii)	Phosphorylation	Fractose – 6- Phosphate	Phosphofructokinase	Fructose 1,6 – Diphosphate + ADP
iv)	Cleavage	Fructose 1,6 – Diphosphate	Fructose diphosphate aldolase	3 – Phosphoglyceraldehyde + Dihydroxy acetone phosphate

(Note: 3 -PGAL and Dihydroxy acetone p	hosphate are interconv	ertible in presence of triose-
No posspin(6) isomerase). Substrate	Enzyme	End product(s)

	1		, and the second s	
v) -	Phosphorylation	3-PGAL+ H <sub>3</sub> PO <sub>4</sub>	Phosphoglycerokinase	1,3 Diphosphoglyceraldehyde.
	Dehydrogenation	1,3 -DPGAL+ NAD <sup>+</sup>	Glyceraldehyde-	1,3 Diphosphoglyceric acid+
			phosphate	NADH+H <sup>+</sup>
			dehydrogenase	
vi)	Dephosphorylation	1, 3-DPGA + ADP	Diphosphoglycerokinase	ATP +
			(Mg2+)	3 - Phosphoglyceric acid.
vii)	Rearrangement	3-PGA	Phosphoglyceromutase	2-Phosphoglyceric acid
viii)	Dehydration	2-PGA	Enolase	Phosphoenol Pyruvic acid $+$ $H_2O$
ix)	Dephosphorylation	Phosphoenol Pyruvic acid	Pyruvate kinase	Pyruvic acid + ATP
		+ADP		

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*
    Various steps involved in Krebs cycle :
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No.	Step(s) Substrate		Enzyme	End product(s)		
i)	Condensation	Acety I-CoA + Oxalo - acetic acid + $H_2O$	Citric acid synthetase	Citric acid (6C) + Coenzyme-A		
iii)	Rearrangement	It is again completed in two steps : a.Citric acid b. Cis-Aconitic acid + H <sub>2</sub> O	Aconitase . Aconitase	Cis- Aconitic acid + $H_2O$ Iso - citric acid (6C) Iso-citric acid (6C)		
iii)	Dehydrogenation (-2H)	Isocitric acid + NAD <sup>+</sup>	Scitrate dehydro genase	Oxalo succinic acid(6C) + NADH+H <sup>+</sup>		
iv)	Decarboxy lation $(-CO_2)$	Oxalo succinic acio	Oxalosuccinate decarboxylase	$\alpha$ - ketoglutaric acid(5C) + CO <sub>2</sub>		
v)	Oxidative Decarboxylation. (-2H) (-CO <sub>2</sub> ).It is also completed in	<ul> <li>a. α -Ketoglutaric acid</li> <li>H, α + NAD<sup>+</sup> +</li> <li>Coenzyme A</li> <li>b. Succinyl- Co A</li> </ul>	+ α - Ketoglutarate dehydrogenase complex	Succinyl- CoA(4C) + NADH +H <sup>+</sup> CO <sub>2</sub>		
	two steps:	+ GDP+iP	Succinate thiokinase	Succinic acid(4C) + Coenzyme-A + GTP		
vi)	Dehydrogeoation (-2H)	Succinic acid + FAD (H-acceptor)	Succinate dehydro - genase	Fumaric acid $(4C) + FADH_2$		
vii)	Hydration(+H <sub>2</sub> O)	Fumaric acid + $H_2O$	Fumarase	Malic acid (4C)		
viji	behydrogenation (-2H)	Malic acid + NAD+ (H-acceptor)	Malate dehydrogenase	Oxaloacetic acid + NADH+ H <sup>+</sup> (4C)		



# • Scientists and their Contribution

No.	Scientists	Contribution		
i)	Embden, Meyerhof and Pamas	Worked out various steps and reactions of glycolytic pathway hence, glycolysis is also called EMP pathway.	-	
ii)	Sir Hans Krebs	TCA cycle was discovered by him. He was awarded Nobel Prize in Biochemistry in 1953.	1937	

16

()	Respi	ratior	1 <b>17</b>
	Multiple Choice Question	13.	During Krebs cycle, fumaric acid gets converted
1.	Respiration is regarded as a process.		a) decarboxylation b) dehydrogenation
	a) catabolic b) anabolic		c) dehydration d) hydration
	c) reduction d) synthetic	14.	The net gain of energy from a molecule of glucose
2.	Mitochondria are regarded as semi autonomous		in the perceptic respiration is
	organelles, due to the presence of		a) 38 b) 35
	a) Cristae b) RNA		c) 70 d) 76
	c) DNA d) ribosomes	15.	Mitochondria are called
3.	Which of the following is a 5C compound	Ó	a) store house of cell
	produced in Krebs cycle?	$\mathbf{x}$	b) power house of cell
	a) Citric acid	9	c) energy store of cell
	b) Oxalosuccinic acid		d) repair mechanism of cell
	c) a-ketoglutaric acid	16.	The amount of energy lost in respiration in the
	d) Succinic acid		form of heat is about
4.	In glycolysis, dehydration occurs during formation		a) 40% b) 50%
	of		c) 60% d) 70%
	a) 3-PGA c) PEPA	17.	Membranes of mitochondria are made up of
	b) 2-PGA d) DNAP		a) glycoprotein
5.	Which of the following is an aerobic respiration?		b) lipoprotein
	a) Glycolysis (b) Decarboxylation		c) phospho protein
	c) Reduction d) Phosphorylation	10	d) chromoprotein
6.	Electron carriers of oxidative phosphorylation are	18.	$F_1$ particles are present on
•	present on		a) cristae b) inner membrane
	a) outer membrane of mitochondria	10	c) both a) and b) d) oxysomes
	b) inner membrane of mitochondria	19.	Glycolysis is also called
	c) thy lakoid membrane of chloroplast		a) HSK pathway b) Discumthatic pathway
	a matrix of mitochondria		b) Biosynthetic pathway
7.	During anaeropic respiration decarboxylation		c) EMP pathway d) C, pathway
•	reaction occurs in	20	d) $C_3$ painway The first compound formed in glycolysis is
	a) cytonlasm	20.	a) Chucose 6 phosphate
	b) mitochondrial matrix		a) Glucose 1.6 biphosphate
	c) cristae		c) Fructose-6-nhosnhate
	d) E particles		d) Pyruvic acid
8	Which of the following derives maximum energy	21	Which of the following compounds in last step of
0.	ner molecule of glucose?	41.	glycolysis gives pyruvic acid?
	a) Glycolysis in liver cells		a) 3 - PGAL b) DHAP
	b) Alcoholic fermentation		c) PEPA d) 2- PGA
	c) Lactic acid fermentation	22.	Oxidation of pyruvic acid occurs in
	d) Aerobic respiration unicellular organisms		a) cytoplasm b) matrix
D	In Krebs cycle, dehydration of substrate occurs		c) F, particles d) Golgi bodies
	a) once	23.	Krebs cycle is also called TCA cycle because
	a) once c) times		a) the first compound formed is citric acid.
10	Which of the following steps generate ATP		b) it was discovered by Sir Hans Krebs.
10.	which of the following steps generate ATF		c) organic acids formed have 3 carboxylic acid
	without $E I S ?$		groups.
	a) Fyluvic actu $\rightarrow$ Acetyl Co-A		d) acetyl Co-A is formed
	b) $\alpha$ - Kelogiularate $\rightarrow$ Succinic acid	24.	Which of the following compound is the acceptor
	c) Iso - citric acid $\rightarrow$ Oxalosuccinic acid		of Acetyl Co-A in Krebs cycle ?
11	a) Succinyl Co-A $\rightarrow$ Succinic acta		a) Oxalo acetic acid
11.	i ne cytochrome which donates de-energised		b) Fumaric acid
	electron to oxygen is		c) Malic acid
	a) cytochrome-a b) cytochrome-b		d) Oxalo succinic acid
	c) cytochrome-a, d) cytochrome-c	25.	During the conversion of succinic acid to fumaric
12.	In Krebs cycle, the acid which undergoes		acid, the hydrogen is accepted by
	oxidative decarboxylation is		a) NAD b) FAD
	a) citric acid b) oxalosuccinic acid		c) $\alpha$ - KGA d) Isocitric acid
	c) malic acid d) a-ketoglutaric acid		

<ul> <li>26. Which of the following compounds is formed in Krebs cycle from fimmaric acid?</li> <li>a) Oxalo acetic acid b) Malic acid</li> <li>c) a - KGA</li> <li>d) Citric acid</li> <li>a) cytoplasm</li> <li>b) F- particles</li> <li>c) matix</li> <li>d) oxysomes</li> <li>28. Each molecule of NADH, through ETS yields</li> <li>a) ATP</li> <li>b) 2ATP molecules</li> <li>d) 4ATPs</li> <li>29. One glucose molecule, through ETS yields</li> <li>a) 2ATP molecules</li> <li>d) 4ATP molecules</li> <lid) 4atp<="" th=""><th><math>\bigcirc</math></th><th>Respi</th><th>ratio</th><th></th><th>18</th></lid)></ul>	$\bigcirc$	Respi	ratio		18
Krebs cycle from fumaric acid ?a) Oxalo acetic acid d?a) Oxalo acetic acid b) Malic acida) alcoholb) lactosec) $\alpha \times KGA$ d) Citric acidc) actaldehyded) glucose27. Enzymes for Krebs cycle are found ina) alcoholb) Pyruvic acida) atrixd) oxysomes38.28. Each molecule of NADH, through ETS yieldsa) 1ATPb) 2ATPsc) 3ATPd) 4ATPsc) actaldehydeb) glucose29. One glucose molecule, through ETS yieldsa) 2ATP moleculesb) 3ATP moleculesc) 34ATP moleculesd) 3ATP moleculesc) actaldehydeb) matrixc) 24.8.6d) 68.24c) coxysomed) ribosome31. In which of the following scare dehydrogenatioa. 86.24b) 8.24.6c) 24.8.6d) 68.24c) Both a) and b) d) Succinic acid31. In which of the following scare dehydrogenatioa. Co.Q $\rightarrow$ cyto.b.c $\rightarrow$ cyto.c. $\rightarrow$ cyto.c. $a_{1}$ a) Clucose - Glucose - almosphateb) PGAL - 1.40PGA32. The common pathway for both acerobic and anacrobre cespiration isa) XFPS cyclea) Cherobaly 5-C compound produced in Krebs cycle isc) Co.Q $\rightarrow$ cyto.b.c $\rightarrow$ cyto.c. $a_{2}$ cyto.b.c $\rightarrow$ cyto.c. $a_{3}$ 33. The compound produced in Krebs cycle isd) Terminal oxidation34.11c) 3b) 2d) 435. The compound produced in Krebs cycle is caids $a_{2}$ d) Carbohydrates are used as respiratory substrates.36.12b) 2d) 437. The compound produced in Krebs cycle is caids $a_{2}$ d) Arr in small stepwise units38. Freb conyces i	26.	Which of the following compounds is formed in	37.	Lactic acid is formed from	m
a) Oxalo acetic acid b) Malic acid c) $\alpha$ - KGA d) Citric acid 27. Enzymes for Krebs cycle are found in a) cytoplasm b) F <sub>1</sub> particles c) matrix d) oxysomes 28. Each molecule of NADH, through ETS yields a) 1ATP b) 2ATPs c) 3ATP molecules b) 3ATP molecules c) 3ATP molecules b) 3ATP molecules c) 34ATP molecules d) 3ATP molecules c) 248,6 d) 6.8,2 31. In which of the following sche dehydrogenation cocurs? a) Glucose - Glucose b mosphate b) 3-PGA - 2-PG c) PEPA _ pyrtyme d) PGAL - 1.1. dPGA 32. The computo pathway for both aerobic and anaerobic respiration is a) Krebs cycle b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid c) Succinate d) Oxalo-acetate 34. During Krebs cycle, decarboxylation occurs times. a) 1 c) 3 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid a) 1 c) 0.9 d) 0.1 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid a) 1 c) 0.9 d) 0.1 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid a) 1 marge acids are used as respiratory substrates. b) Organic acids are used as respiratory substrates. c) Thm axiation of respiratory substrates. b) Organic acids are used as respiratory substrates. b) Organic acids are used as respiratory substrates. b) Organic acids are used as respiratory substrates. c) The oxidation of respiratory substrates. b) Organic acids are used as respiratory substrates. c) The oxidation		Krebs cycle from fumaric acid ?		a) alcohol b)	lactose
c) α - KGA d) Citric acid 37. Enzymes for Krebs cycle are found in a) cytoplasm b) F, particles c) matrix d) oxysomes 38. The compound common to both aerobic and anaerobic respiration is a) glucosc b) pyruvic acid a) algucosc b) pyruvic acid a) glucosc b) pyruvic acid a) glucosc b) pyruvic acid a) glucosc b) actia delayde b) lactic acid b) actiadehyde b) lactic acid c) actaldehyde b) lactic acid c) consument b) actic acid c) actaldehyde b) lactic acid c) actaldehyde b) lactic acid c) consument b) archares c) 3ATP molecules c) 3ATP molecules c) 3ATP molecules c) 3AATP molecules c) 3ATP and b) d) Succinic acid d) Co.Q → cyto.b.c, → cyto.c.→ cyto.a.a, → cyto.c. c) Co.Q → cyto.b.c → cyto.a.a, → cyto.c. c) carbohydrates c) and for action malic acid to oxalo acetic acid c) carbohydrates c) analate reductase b) malate thiokinase c) fumarase c)		a) Oxalo acetic acid b) Malic acid		c) acetaldehyde d)	glucose
<ul> <li>27. Enzymes for Krebs cycle are found in a cytoplasm b) F, particles c) matrix d) oxysomes</li> <li>28. Each molecule of NADH, through ETS yields a) 1ATP b) 2ATPs c) 3ATP molecules b) 2ATPs d) 4ATPs</li> <li>29. One glucose molecule, through ETS yields a) 2ATP molecules b) 3ATP molecules c) 34ATP molecules d) 38ATP molecules c) 34ATP molecules d) 68.24</li> <li>30. The adenosine triphosphate (ATP) gain through 27D and an accurate d) 0x3comet for the following scate dehydrogenation occurs ?</li> <li>a) Glucose - Glucose to mosphate b) 3-PGA - 2-PGA (c) PEPA pyrotect c) SATS d) Terminal oxidation 33 PGA - 2-PGA (c) PEPA pyrotect c) Succinate d) Oxalo-acetate d) Citrate b) α - ketoglutarate c) Succinate d) Oxalo-acetate 34. During Krebs cycle, decarboxylation occurs fitmes.</li> <li>a) 1 c) 3 b) 2 d) 4</li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by a) malate thiokinase c) fumarase d) malate dehydrogenes 36. Acetyl CoA produced from pyruvate by a) oxidative decarboxylation</li> <li>36. Acetyl CoA produced from pyruvate by a) oxidative decarboxylation</li> <li>37. The conversion of malic acid to oxalo acetic acid is catalyzed by a) malate thiokinase c) fumarase d) malate dehydrogenes 36. Acetyl CoA produced from pyruvate by a) oxidative decarboxylation</li> <li>36. Acetyl CoA produced from pyruvate by a) oxidative decarboxylation</li> <li>37. The</li></ul>		c) $\alpha$ - KGA d) Citric acid	38.	The compound commo	n to both aerobic and
a) cytoplasm b) F, particles c) matrix d) oxysomes 28. Each molecule of NADH, through ETS yields a) 1ATP b) 2ATPs c) 3ATPs d) 4ATPs 29. One glucose molecule, through ETS yields a) 2ATP molecules b) 3ATP molecules c) 34ATP molecules b) 3ATP molecules d) Columbra and Kreb cycle respectively are c) 248,6 d) 6.8,23 31. In which of the following sease dehydrogenation occurs? a) Glucose - Glucose to hosphate b) 3-PGA - 2-PG c) PEPA pyruche b) 3-PGA - 2-PG c) PEPA pyruche c) Succinate b) EMP pathway c) HES d) Terminal oxidation anaareotor espiration is a) Krobs cycle b) EMP pathway c) HES d) Terminal oxidation 33. The only 5-C compound produced in Krebs cycle is c) Cirtate b) $\alpha$ - ketoglutarate c) firmarase d) malate thiokinase c) firmarase d) malate theolydrogenase 36. Acetyl CoA produced from pyruvate by a) oxidative decarboxylation	27.	Enzymes for Krebs cycle are found in		anaerobic respiration is	
c) matrix d) oxysomes 28. Each molecule of NADH, through ETS yields a) 1ATP b) 2ATPs c) 3ATP molecules b) 2ATP molecules c) 3ATP molecules d) 3ATP molecules c) 24,8,6 d) 6,8,23 31. In which of the following second hydrogenation occurs? a) Glucose - Glucose to thosphate c) PEA _ pyrupare d) PGAL - 1, EdPGA 32. The compute pathway for both aerobic and anaerobre espiration is a) Kebsr cycle b) EMP pathway c) FMS d) Terminal oxidation 33. The conversion of malic acid to oxalo acetic acid a) 1 c) 3 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid a) 1 c) 3 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid a) 1 c) 3 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid a) 1 c) 3 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid a) 1 c) 3 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid a) 1 c) 3 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid a) 1 c) 3 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid a) 1 act reductase b) malate thiokinase c) furmarase d)		a) cytoplasm b) F. particles		a) glucose b)	pyruvic acid
<ul> <li>28. Each molecule of NADH, through ETS yields <ul> <li>a) 1ATP</li> <li>b) 2ATPs</li> <li>c) 3ATP molecules</li> <li>d) 4ATPs</li> </ul> </li> <li>29. One glucose molecule, through ETS yields <ul> <li>a) 2ATP molecules</li> <li>b) 3ATP molecules</li> <li>c) 34ATP molecules</li> <li>b) 3ATP molecules</li> <li>c) 34ATP molecules</li> <li>d) 38ATP molecules</li> <li>d) 38ATP molecules</li> <li>d) 38ATP molecules</li> <li>d) 38ATP molecules</li> <li>d) 4ATP gain 4Try 414 alcohol</li> <li>Acetylation takes place in <ul> <li>a) 2ATP molecules</li> <li>d) 3ATP molecules</li> <li>d) 3ATP molecules</li> <li>d) 3ATP molecules</li> <li>d) 3ATP molecules</li> <li>d) 4ATP gain 4Try alcohol</li> <li>d) condecarboxylation, pyruvate gives are actaldehyde b) lactic acid</li> <li>d) ethyl alcohol</li> <li>d) ethyl alcohol</li> <li>d) methyl alcohol</li> <li>Acetyl Acohol</li> <li>d) methyl alcohol</li> <li>Acetyl Acohol</li> <li>d) 4ATP in 4Cohol</li> <li>d) Condecarboxylation, pyruvate gives are actaldehyde b) lactic acid</li> <li>d) Condecarboxylation respiratory auditic acid</li> <li>d) Condecarboxylation for 4D and b)</li> <li>d) Condecarboxylation fore 5D and and b)</li> <li>d) Condecarboxylation for 4D</li></ul></li></ul></li></ul>		c) matrix d) oxysomes		c) acety CoA d)	free oxygen
a) IATP b) 2ATPs c) 3ATPs d) 4ATPs 29. One glucose molecule, through ETS yields a) 2ATP molecules b) 3ATP molecules c) 34ATP molecules d) 38ATP molecules c) 24,86 d) 68,24 31. In which of the following schedenydrogenation occurs ? a) Glucose - Glucose chosphate b) 3-PGA - 2-PGA c) PEPA _ pyrnote d) PGAL - 1.1 cuPGA 32. The common pathway for both aerobic and anaaroth cespiration is a) Kros cycle b) EMP pathway c) Citrate b) $\alpha$ - ketoglutarate c) Succinate d) Oxalo-acetate 34. During Krebs cycle, decarboxylation occurs times. a) 1 c) 3 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid is catalyzed by a) malate reductase b) malate thiokinase c) fumarase d) malate dehydrogenase 36. Acetyl CoA produced from pyruxate by a) oxidative decarboxylation	28.	Each molecule of NADH through ETS yields	39.	On devarboxylation, pyru	ivate gives
c) $3ATPs$ d) $4ATPs$ 29. One glucose molecule, through ETS yields a) $2ATP$ molecules b) $3ATP$ molecules c) $34ATP$ molecules d) $38ATP$ molecules 30. The adenosine triphosphate (ATP) gain forming glycolysis, connecting link and Kreb cycle respectively are 10Cf 2013] a) $8,624$ b) $8,24,6$ c) $24,8,6$ d) $6,8,24$ 31. In which of the following schedehydrogenation occurs ? a) Glucose - Glucose to hosphate b) $3-PGA - 2-PGA$ c) PEPA pyrtuent d) PGAL - 1,1 diPGA 32. The commod pathway for both aerobic and anaerobic respiration is a) Krebs cycle b) EMP pathway c) HTS d) Terminal oxidation 33. The only 5-C compound produced in Krebs cycle is c) Succinate d) Oxalo-acetate c) Succinate d) Oxalo-acetate 34. During Krebs cycle, decarboxylation occurs times. a) 1 c) 3 b) 2 d) 4 35. The conversion of malica cid to oxalo acetic acid is catalyzed by a) malate reductase b) malate thiokinase c) furmarase d) malate dehydrogenase 36. Acetyl CoA produced from pyruvate by a) oxidative decarboxylation	-0.	a) 1ATP b) 2ATPs		a) acetaldehyde b)	lactic acid
<ul> <li>29. One glucose molecule, through ETS yields <ul> <li>a) 2ATP molecules</li> <li>b) 3ATP molecules</li> <li>c) 34ATP molecules</li> <li>d) 3ATP molecules</li> <li>d) 44. Acetylation takes place in <ul> <li>a) cytoplasm</li> <li>b) matrix</li> <li>c) cysome</li> <li>d) ribosome</li> </ul> </li> <li>44. Acetylation takes place in <ul> <li>a) cytoplasm</li> <li>b) matrix</li> <li>c) cysome</li> <li>d) ribosome</li> </ul> </li> <li>44. Acetylation takes place in <ul> <li>a) cytoplasm</li> <li>b) matrix</li> <li>c) cysome</li> <li>d) ribosome</li> </ul> </li> <li>44. Acetylation takes place in <ul> <li>a) cytoplasm</li> <li>b) matrix</li> <li>c) cysome</li> <li>d) ribosome</li> </ul> </li> <li>44. Acetylation takes place in <ul> <li>a) cytoplasm</li> <li>b) matrix</li> <li>c) cysome</li> <li>d) ribosome</li> </ul> </li> <li>44. Acetylation takes place in <ul> <li>a) cytoplasm</li> <li>b) matrix</li> <li>c) cysome</li> <li>d) respiratory automage takes and bio discome takes place in <ul> <li>a) cytoplasm</li> <li>b) addition</li> </ul> </li> <li>44. Acetylation takes place in <ul> <li>a) cytoplasm</li> <li>b) addition</li> <li>c) concordination</li> <li>d) co</li></ul></li></ul></li></ul></li></ul>		c) 3ATPs d) 4ATPs	5	(c) ethyl alcohol d)	methyl alcohol
<ul> <li>a) 2ATP molecules b) 3ATP molecules c) 34ATP molecules b) 3ATP molecules c) 34ATP molecules d) 38ATP molecules c) 34ATP molecules d) 38ATP molecules d) 78ATP molecules d) 78</li></ul>	29	One glucose molecule, through ETS yields	49.	Acetylation takes place in	n
<ul> <li>c) 34ATP molecules d) 38ATP molecules</li> <li>30. The adenosine triphosphate (ATP) gain forming glycolysis, connecting link and Kreb cycle respectively are 10CC 2013]</li> <li>a) 8,6,24 b) 8,24,6</li> <li>c) 24,8,6 d) 6,8,24</li> <li>31. In which of the following steps dehydrogenation occurs ?</li> <li>a) Glucose - Glucose 6 bihosphate</li> <li>b) 3-PGA - 2-PGA</li> <li>c) PEPA _ pyrovate</li> <li>d) PGAL - 1.1 dIPGA</li> <li>32. The commony pathway for both aerobic and anaerobrocespiration is</li> <li>a) Krops cycle b) EMP pathway</li> <li>c) Mark and b conversion of malic acid to oxalo acetic acid</li> <li>d) PGAL - 1.1 dIPGA</li> <li>33. The only 5-C compound produced in Krebs cycle is</li> <li>d) The correct source d) Oxalo-acetate</li> <li>d) During Krebs cycle, decarboxylation occurs_times.</li> <li>a) 1 c) 3</li> <li>b) 2 d) 4</li> <li>d) During Krebs cycle, decarboxylation occurs_times.</li> <li>c) fumarase d) malate thiokinase</li> <li>c) fumarase d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> </ul>	27.	a) 2 ATP molecules b) 3 ATP molecules	2	a) cytoplasm b)	matrix
<ul> <li>41. Which compound is found both in respiration and photosynthesis?</li> <li>33. The denosine triphosphate (ATP) gain form gruptor glycolysis, connecting link and Kreb Cycle respectively are (0.02 × 10.00 × 10.0</li></ul>		c) 3/ATP molecules d) 38ATP molecules		c) oxysome d)	ribosome
<ul> <li>30. The additional tripulsphate (ATT) gain forming glycolysis, connecting link and Krebs cycle is espectively are interfaced and by the system is the correct sequence of electron carriers during C 24,8,6 d) 6,8,24</li> <li>31. In which of the following steps dehydrogenation occurs ? <ul> <li>a) Glucose - Glucose to phosphate</li> <li>b) 3-PGA - 2-PGA</li> <li>c) PEPA _ pyrtweffe</li> <li>d) PGAL - 1, 40PGA</li> </ul> </li> <li>32. The common pathway for both aerobic and anaerobic espiration is <ul> <li>a) Khots cycle</li> <li>b) EMP pathway</li> <li>c) Ref a _ b) PGAL</li> <li>c) Both a) and b) d) Succinic acid</li> </ul> </li> <li>33. Freeonly 5-C compound produced in Krebs cycle is c) Succinate d) Oxalo-acetate</li> <li>34. During Krebs cycle, decarboxylation occurs_times. <ul> <li>a) 1</li> <li>b) 2</li> <li>d) 4</li> </ul> </li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by <ul> <li>a) malate reductase b) malate thiokinase c) fumarase d) malate dehydrogenasa</li> <li>36. Acetyl CoA produced from pyruvate by a) oxidative decarboxylation</li> </ul> </li> </ul>	30	The adenosine triphosphate (ATP) gain orrang	41.	Which compound is found	d both in respiration and
<ul> <li>a) PGA b) PGAL</li> <li>c) Both a) and b) d) Succinic acid</li> <li>d) Co.Q → cyto.b.c → cyto.c → cyto.a.a, → cyto.c</li> <li>d) Co.Q → cyto.b.c → cyto.a.a, → cyto.c</li> <li>d) Co.Q → cyto.c → cyto.a.a, → cyto.c</li> <li>d) Co.Q → cyto.b.c → cyto.a.a, → cyto.c</li> <li>d) Co.Q → cyto.c → cyto.a</li></ul>	50.	alwoolvsis, connecting link and Krahovala		photosynthesis?	
a) 86,24 b) 8,24,6 c) 24,8,6 d) 6,8,24 31. In which of the following separately drogenation occurs? a) Glucose - Glucose comosphate b) 3-PGA - 2-PGA c) PEPA _ pyrnythe d) PGAL - 1,1 dPGA 32. The common pathway for both aerobic and anaerofic respiration is a) Krobs cycle b) EMP pathway c) HFS d) Terminal oxidation 33. The only 5-C compound produced in Krebs cycle is a) Citrate b) $\alpha$ - ketoglutarate c) Succinate d) Oxalo-acetate 34. During Krebs cycle, decarboxylation occurs times. a) 1 c) 3 b) 2 d) 4 35. The conversion of malic acid to oxalo acetic acid is catalyzed by a) malate reductase b) malate thiokinase c) fumarase d) malate dehydrogenase 36. Acetyl CoA produced from pyruvate by a) oxidative decarboxylation		respectively are		a) PGA b)	PGAL
<ul> <li>a) 6,0,24</li> <li>b) 6,8,24</li> <li>c) 24,8,6</li> <li>d) 6,8,24</li> <li>d) Co.Q → cyto.c. → cyto.c. → cyto.c.</li> <li>d) Co.Q → cyto.b.c → cyto.c. → cyto.a.a, → cyto.c</li> <li>d) Co.Q → cyto.b.c → cyto.a.a, → cyto.c</li> <li>d) ATP in large oxidation reaction</li> <li>a) ATP in large oxidation reaction</li> <li>a) for consumed more oxygen than the amount of CO. released.</li> </ul>		1000000000000000000000000000000000000		c) Both a) and b) d)	Succinic acid
<ul> <li>31. In which of the following steps dehydrogenation occurs?</li> <li>a) Glucose - Glucose c phosphate</li> <li>b) 3-PGA - 2-PGA</li> <li>c) PEPA _ pyrtyche</li> <li>d) PGAL - 1, 1 diPGA</li> <li>32. The common pathway for both aerobic and anaerobic respiration is</li> <li>a) Krobs cycle b) EMP pathway</li> <li>c) FIS d) Terminal oxidation</li> <li>33. The only 5-C compound produced in Krebs cycle is</li> <li>a) Citrate b) α - ketoglutarate</li> <li>c) Succinate d) Oxalo-acetate</li> <li>34. During Krebs cycle, decarboxylation occurs_times.</li> <li>a) 1 c) 3</li> <li>b) 2 d) 4</li> <li>35. The conversion of malic acid to oxalo acetic acidi is catalyzed by</li> <li>a) malate reductase b) malate thiokinase</li> <li>c) fumarase d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> </ul>		a) $6,0,24$ b) $6,24,0$ c) $24,86$ d) $6,824$	42.	The correct sequence of	electron carriers during
<ul> <li>31. In which of the following seeps dehydrogenation occurs ?</li> <li>a) Glucose - Glucose c phosphate</li> <li>b) 3-PGA - 2-PGA</li> <li>c) PEPA _ pyrtware</li> <li>d) PGAL - 1,1-diPGA</li> <li>32. The common pathway for both aerobic and anaeroptic respiration is</li> <li>a) Krobs cycle</li> <li>b) EMP pathway</li> <li>c) HYS</li> <li>d) Terminal oxidation</li> <li>33. Areonly 5-C compound produced in Krebs cycle is</li> <li>c) Succinate</li> <li>d) Oxalo-acetate</li> <li>d) Oxalo-acetate</li> <li>d) 1</li> <li>c) 3</li> <li>d) 1</li> <li>c) 3</li> <li>d) 1</li> <li>c) 3</li> <li>d) 4</li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by</li> <li>a) malate reductase</li> <li>b) malate thiokinase</li> <li>c) fumarase</li> <li>d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> </ul>	21	(0) 24,0,0 $(0) 0,0,0,0$		ETS is	
<ul> <li>a) Glucose - Glucose Chhosphate</li> <li>b) Co.Q → cyto.a.a<sub>3</sub> → cyto.c<sub>1</sub></li> <li>c) PEPA _ pyruvate</li> <li>d) PGAL - 1,1 dPGA</li> <li>32. The common pathway for both aerobic and anaerobic espiration is</li> <li>a) Krobs cycle</li> <li>b) EMP pathway</li> <li>c) HTS</li> <li>d) Terminal oxidation</li> <li>33. Knoonly 5-C compound produced in Krebs cycle is</li> <li>a) Lico 3</li> <li>b) 2</li> <li>d) Oxalo-acetate</li> <li>34. During Krebs cycle, decarboxylation occurs_times.</li> <li>a) 1</li> <li>c) 3</li> <li>b) 2</li> <li>d) 4</li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by</li> <li>a) malate reductase b) malate thiokinase</li> <li>c) fumarase</li> <li>d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> </ul>	31.	In which of the following steps denydrogenation		a) Co.Q $\rightarrow$ cyto.b.c <sub>1</sub> $\rightarrow$	$cyto.c \rightarrow cyto.a.a_3$
<ul> <li>a) Glucose - Cyto.a.a<sub>3</sub> → cyto.c d) Co.Q → cyto.b.c → cyto.b.c → cyto.a.a<sub>3</sub> → cyto.c d) Co.Q → cyto.b.c → cyto.b.c → cyto.aa<sub>3</sub> → cyto.c d) Co.Q → cyto.b.c → cyto.b.c → cyto.b.c → cyto.b.c → cyto.bc → cyto</li></ul>				b) Co.Q $\rightarrow$ cyto.a.a <sub>3</sub> $\rightarrow$	$cyto.b.c \rightarrow cyto.c_1$
<ul> <li>b) 3-PGA - 2-PGA</li> <li>c) PEPA _ pyruvate</li> <li>d) Co.Q → cyto.c → cyto.b.c → cyto.a.3</li> <li>d) Co.Q → cyto.b.c → cyto.a.3</li> <li>d) Co.Q → cyto.b.c → cyto.b.c → cyto.a.3</li> <li>d) Co.Q → cyto.b.c → cyto.b.c → cyto.a.3</li> <li>d) Co.Q → cyto.c → cyto.b.c → cyto.a.3</li> <li>d) Co.Q → cyto.b.c → cyto.b.c. → cyto.b.c. → cyto.b.c. → cyto.b.c. → cyto.b.c.</li></ul>		a) Glucose - Glucose - nosphate		c) Co.Q $\rightarrow$ cyto.b.c $\rightarrow$	$cyto.a.a_3 \rightarrow cyto.c$
<ul> <li>c) PEPA _ pyreate</li> <li>d) PGAL - 1,1 dPGA</li> <li>32. The common pathway for both aerobic and anaerobic respiration is</li> <li>a) Krots cycle b) EMP pathway</li> <li>c) Wrst d) Terminal oxidation</li> <li>33. Preonly 5-C compound produced in Krebs cycle is</li> <li>a) Citrate b) α - ketoglutarate</li> <li>c) Succinate d) Oxalo-acetate</li> <li>34. During Krebs cycle, decarboxylation occurs times.</li> <li>a) 1 c) 3</li> <li>b) 2 d) 4</li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by</li> <li>a) malate reductase b) malate thiokinase</li> <li>c) fumarase d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> <li>43. The overall goal of glycolysis, Krebs cycle and electron transport system is the formation of</li> <li>a) ATP in large oxidation reaction</li> <li>b) sugars</li> <li>c) nucleic acids</li> <li>d) ATP in small stepwise units</li> <li>44. Respiratory quotient is one in case of</li> <li>a) fatty acids b) nucleic acids</li> <li>c) carbohydrates d) organic acids</li> <li>45. The respiratory quotient (R.Q.) of carbohydrate is</li> <li>is catalyzed by</li> <li>a) malate reductase b) malate thiokinase</li> <li>c) fumarase d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> </ul>		b) 3-PGA - 2-PGA	10	d) Co.Q $\rightarrow$ cyto.c $\rightarrow$ c	$cyto.b.c \rightarrow cyto.a.a_3$
<ul> <li>d) PGAL - 1, 4-0PGA</li> <li>32. The common pathway for both aerobic and anaerobic respiration is <ul> <li>a) Kros cycle</li> <li>b) EMP pathway</li> <li>c) HAS</li> <li>d) Terminal oxidation</li> </ul> </li> <li>33. The only 5-C compound produced in Krebs cycle is <ul> <li>a) Citrate</li> <li>b) α - ketoglutarate</li> <li>c) Succinate</li> <li>d) Oxalo-acetate</li> </ul> </li> <li>34. During Krebs cycle, decarboxylation occurs_times. <ul> <li>a) 1</li> <li>b) 2</li> <li>d) 4</li> </ul> </li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by <ul> <li>a) malate reductase</li> <li>b) malate thiokinase</li> <li>c) fumarase</li> <li>d) malate dehydrogenase</li> </ul> </li> <li>36. Acetyl CoA produced from pyruvate by <ul> <li>a) oxidative decarboxylation</li> </ul> </li> </ul>		c) PEPA pyruvae	43.	The overall goal of glyce	olysis, Krebs cycle and
<ul> <li>32. The common pathway for both aerobic and anaerobic respiration is <ul> <li>a) Kreps cycle</li> <li>b) EMP pathway</li> <li>c) Has</li> <li>d) Terminal oxidation</li> </ul> </li> <li>33. The only 5-C compound produced in Krebs cycle is <ul> <li>a) Citrate</li> <li>b) α - ketoglutarate</li> <li>c) Succinate</li> <li>d) Oxalo-acetate</li> </ul> </li> <li>34. During Krebs cycle, decarboxylation occurs_times. <ul> <li>a) 1</li> <li>b) 2</li> <li>d) 4</li> </ul> </li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by <ul> <li>a) malate reductase</li> <li>c) fumarase</li> <li>d) malate thiokinase</li> <li>c) fumarase</li> <li>d) malate dehydrogenase</li> </ul> </li> <li>36. Acetyl CoA produced from pyruvate by <ul> <li>a) oxidative decarboxylation</li> </ul> </li> </ul>		d) PGAL - I, J-dPGA		electron transport system	i is the formation of
<ul> <li>anaerofic respiration is</li> <li>a) Kros cycle b) EMP pathway</li> <li>c) Kros cycle d) Terminal oxidation</li> <li>33. Pro only 5-C compound produced in Krebs cycle is</li> <li>a) Citrate b) α - ketoglutarate</li> <li>c) Succinate d) Oxalo-acetate</li> <li>34. During Krebs cycle, decarboxylation occurs_times.</li> <li>a) 1 c) 3</li> <li>b) 2 d) 4</li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by</li> <li>a) malate reductase b) malate thiokinase</li> <li>c) fumarase d) malate thiokinase</li> <li>c) fumarase d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> </ul>	32.	The common pathway for both aerobic and		a) ATP in large oxidation	reaction
<ul> <li>a) Kreos cycle</li> <li>b) EMP pathway</li> <li>c) Has</li> <li>d) Terminal oxidation</li> <li>33. Are only 5-C compound produced in Krebs cycle is</li> <li>a) Citrate</li> <li>b) α - ketoglutarate</li> <li>c) Succinate</li> <li>d) Oxalo-acetate</li> <li>34. During Krebs cycle, decarboxylation occurs_times.</li> <li>a) 1</li> <li>b) 2</li> <li>c) 3</li> <li>b) 2</li> <li>d) 4</li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by</li> <li>a) malate reductase</li> <li>b) malate thiokinase</li> <li>c) fumarase</li> <li>d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> </ul>		anaeropic respiration is		b) sugars	
<ul> <li>a) 1 c) 3</li> <li>b) 2 d) 4</li> <li>c) a - ketoglutarate c) Succinate d) Oxalo-acetate</li> <li>d) Oxalo-aceta</li></ul>		a) Kreps cycle b) EMP pathway		d) ATD in small stonwige	unita
<ul> <li>33. The only 5-C compound produced in Krebs cycle is C is cally zero by α - ketoglutarate c) Succinate d) Oxalo-acetate</li> <li>34. During Krebs cycle, decarboxylation occurs</li></ul>		c) Ers d) Terminal oxidation	11	a) AIF III sinali stepwise Respiratory quotient is or	rain assa of
<ul> <li>a) Citrate b) α-ketoglutarate c) Succinate d) Oxalo-acetate</li> <li>34. During Krebs cycle, decarboxylation occurs times.</li> <li>a) 1 c) 3</li> <li>b) 2 d) 4</li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by</li> <li>a) malate reductase b) malate thiokinase</li> <li>c) fumarase d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> </ul>	33.	ne only 5-C compound produced in Krebs cycle is	44.	a) fatty acids b)	nucleic acids
<ul> <li>34. During Krebs cycle, decarboxylation occurs_times.</li> <li>a) 1 c) 3</li> <li>b) 2 d) 4</li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by</li> <li>a) malate reductase b) malate thiokinase</li> <li>c) fumarase d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> </ul>		a) Citrate b) $\alpha$ - ketoglutarate		c) carbohydrates d)	organic acids
<ul> <li>34. During Krebs cycle, decarboxylation occurs</li></ul>	~ ^	c) Succinate d) Oxalo-acetate	45	The respiratory quotient (	$(\mathbf{R} \mathbf{O})$ of carbohydrate is
<ul> <li>a) 1 c) 3</li> <li>b) 2 d) 4</li> <li><b>35.</b> The conversion of malic acid to oxalo acetic acid is catalyzed by <ul> <li>a) malate reductase</li> <li>c) fumarase</li> <li>d) malate thiokinase</li> <li>c) fumarase</li> <li>d) malate dehydrogenase</li> </ul> </li> <li><b>36.</b> Acetyl CoA produced from pyruvate by <ul> <li>a) oxidative decarboxylation</li> </ul> </li> <li>a) 0.7 b) 1</li> <li>b) 1</li> <li>c) 0.9 d) 0.1</li> </ul> <li><b>46.</b> If R.O. is less than 1 in a respiratory metabolism, it would mean that <ul> <li>a) Carbohydrates are used as respiratory substrates.</li> <li>b) Organic acids are used as respiratory substrates.</li> <li>c) The oxidation of respiratory substrates.</li> <li>c) The oxidation of respiratory substrates.</li> </ul></li>	34.	During Krebs cycle, decarboxylation occurs	ч	The respiratory quotient (	[Oct 2014]
<ul> <li>a) 1 c) 3</li> <li>b) 2 d) 4</li> <li>c) 0.9 d) 0.1</li> <li>c) 0.9 d) 0.1</li> <li>d) 4</li> <li>d) 6.7 c) 0.9 d) 0.1</li> <li>d) 0.1</li> <li>d) 6.7 c) 0.9 d) 0.1</li> <li>d) 6.8 c) 0.9 c) 0.1</li> <li>d) 6.8 c) 0.9 c) 0.1</li> <li>d) 6.8 c) 0.1 c) 0.1</li> <li>d) 6.8 c) 0.1 c</li></ul>		times.		a) 0.7 b)	1
<ul> <li>b) 2 d) 4</li> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by <ul> <li>a) malate reductase</li> <li>b) malate thiokinase</li> <li>c) fumarase</li> <li>d) malate dehydrogenase</li> </ul> </li> <li>36. Acetyl CoA produced from pyruvate by <ul> <li>a) oxidative decarboxylation</li> </ul> </li> <li>46. If R.O. is less than 1 in a respiratory metabolism, it would mean that <ul> <li>a) Carbohydrates are used as respiratory substrates.</li> <li>b) Organic acids are used as respiratory substrates.</li> <li>c) The oxidation of respiratory substrates.</li> <li>c) The oxidation of respiratory substrate.</li> </ul></li></ul>		a) 1 c) 3		(0, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0	01
<ul> <li>35. The conversion of malic acid to oxalo acetic acid is catalyzed by</li> <li>a) malate reductase b) malate thiokinase</li> <li>c) fumarase d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> <li>iii RCO. IS less than 1 in a respiratory inclusion in a respiratory substrates.</li> <li>b) Organic acids are used as respiratory substrates.</li> <li>c) The oxidation of respiratory substrate consumed more oxygen than the amount of CO. released.</li> </ul>		b) 2 d) 4	46	If $R \cap is$ less than 1 in a	respiratory metabolism
<ul> <li>is catalyzed by</li> <li>a) malate reductase</li> <li>b) malate thiokinase</li> <li>c) fumarase</li> <li>d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> </ul>	35.	The conversion of malic acid to oxalo acetic acid	т.,	it would mean that	i ospitator y metabolism,
<ul> <li>a) malate reductase b) malate thiokinase</li> <li>c) fumarase d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> <li>b) Organic acids are used as respiratory substrates.</li> <li>c) The oxidation of respiratory substrates</li> <li>c) The oxidation of respiratory substrate</li> </ul>		is catalyzed by		a) Carbohydrates are used	as respiratory substrates
<ul> <li>c) fumarase d) malate dehydrogenase</li> <li>36. Acetyl CoA produced from pyruvate by</li> <li>a) oxidative decarboxylation</li> <li>c) fumarase d) malate dehydrogenase</li> <li>c) The oxidation of respiratory substrate</li> <li>c) The oxidation of respiratory substrate</li> <li>c) c) organic actus are used as respiratory substrates.</li> <li>c) The oxidation of respiratory substrate</li> <li>c) c) organic actus are used as respiratory substrates.</li> <li>c) The oxidation of respiratory substrate</li> <li>c) The oxidation of respiratory substrate</li> <li>c) c) organic actus are used as respiratory substrates.</li> <li>c) The oxidation of respiratory substrate</li> <li>c) c) organic actus are used as respiratory substrates.</li> <li>c) The oxidation of respiratory substrate</li> <li>c) c) consumed more oxygen than the amount of CO. released.</li> </ul>		a) malate reductase b) malate thiokinase		h) Organic acids are used	as respiratory substrates
<b>36.</b> Acetyl CoA produced from pyruvate by a) oxidative decarboxylation		c) fumarase d) malate dehydrogenase		c) The oxidation of r	espiratory substrate
a) oxidative decarboxylation CO, released.	36.	Acetyl CoA produced from pyruvate by		consumed more oxyg	en than the amount of
		a) oxidative decarboxylation		CO, released.	Sen than the unbuilt of

- b) oxidative photophosphorylationc) oxidative hydrogenation

( )

d) oxidative photorespiration

d) The oxidation of respiratory substrate consumed less oxygen than the amount of  $CO_2$ released

# **Answer Keys**

<b>1.</b> a)	<b>2.</b> c)	3. c)	4. c)	5. c)	<b>6.</b> b)	7. d)	<b>8.</b> d)	<b>9.</b> a)	<b>10.</b> d)
11. c)	<b>12.</b> d)	<b>13.</b> d)	14. a)	<b>15.</b> b)	16. c)	17. b)	18. c)	<b>19.</b> c)	<b>20.</b> a)
<b>21.</b> c)	<b>22.</b> b)	23. c)	<b>24.</b> a)	<b>25.</b> b)	<b>26.</b> b)	27. c)	<b>28.</b> c)	<b>29.</b> c)	<b>30.</b> a)
<b>31.</b> d)	<b>32.</b> b)	<b>33.</b> b)	<b>34.</b> b)	<b>35.</b> d)	<b>36.</b> a)	<b>37.</b> b)	<b>38.</b> b)	<b>39.</b> a)	<b>40.</b> a)
<b>41.</b> c)	<b>41.</b> a)	<b>43.</b> d)	<b>44.</b> c)	<b>45.</b> b)	<b>46.</b> c)				

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