

# Cellular respiration: The process powering life.

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## Introduction

Cellular respiration is a fundamental biochemical process that powers nearly all life forms on Earth. It involves the conversion of nutrients into adenosine triphosphate (ATP), the primary energy currency of cells. This complex sequence of reactions not only generates energy but also produces essential metabolic intermediates and maintains cellular homeostasis. In this article, we will explore the intricacies of cellular respiration, its key stages, and its significance for both cellular and organismal health [1].

Cellular respiration is the process by which cells break down organic molecules, such as glucose, to produce ATP. It occurs in three main stages: glycolysis, the citric acid cycle (Krebs cycle), and oxidative phosphorylation. These stages are intricately linked and occur in different cellular compartments, primarily in the cytoplasm and mitochondria.

The phosphorylated glucose is then split into two three-carbon molecules, which are further processed to produce four ATP molecules (a net gain of two ATP) and two molecules of NADH, a carrier of electrons [2].

Glycolysis yields two molecules of pyruvate, two ATP molecules, and two NADH molecules per glucose molecule. oxidative phosphorylation consists of two interconnected processes: the electron transport chain and chemiosmosis.

High-energy electrons from NADH and FADH<sub>2</sub> are transferred through a series of protein complexes embedded in the inner mitochondrial membrane. As electrons move through the chain, they cause protons (H<sup>+</sup> ions) to be pumped from the mitochondrial matrix into the intermembrane space, creating an electrochemical gradient [3].

This proton gradient drives protons back into the mitochondrial matrix through ATP synthase, a protein complex that synthesizes ATP from ADP and inorganic phosphate (Pi). The final electron acceptor in the chain is oxygen, which combines with electrons and protons to form water [4].

ATP generated through cellular respiration powers almost all cellular activities, including muscle contraction, biosynthesis, and active transport. the byproducts of cellular respiration, including intermediates from the citric acid cycle, are used in various biosynthetic pathways to produce amino acids, nucleotides, and lipids [5].

Some energy from cellular respiration is released as heat, which helps maintain body temperature in warm-blooded animals.

cellular respiration is tightly regulated to meet the energy demands of the cell. Key regulatory mechanisms include the availability of glucose, oxygen, and other substrates influences the rate of cellular respiration [6].

Specific enzymes involved in glycolysis, the citric acid cycle, and oxidative phosphorylation are regulated by various factors, including allosteric effectors and covalent modifications [7].

Hormones such as insulin and glucagon regulate the availability of glucose and other nutrients, thereby influencing cellular respiration. dysregulation or impairment of cellular respiration can lead to various diseases conditions like diabetes and lactic acidosis are associated with disruptions in glycolysis and the citric acid cycle [8].

Genetic mutations affecting mitochondrial DNA or nuclear genes involved in mitochondrial function can lead to disorders such as mitochondrial myopathy and neurodegenerative diseases [9].

Cancer cells often exhibit altered cellular respiration, relying more on glycolysis (a phenomenon known as the Warburg effect) even in the presence of oxygen [10].

## Conclusion

Cellular respiration is a critical process that underpins life by converting nutrients into usable energy. Understanding its stages—glycolysis, the citric acid cycle, and oxidative phosphorylation—provides insight into how cells generate ATP and maintain homeostasis. Insights into cellular respiration also pave the way for advancements in treating metabolic disorders, mitochondrial diseases, and cancer. As research continues to unravel the complexities of cellular respiration, it holds promise for improving our understanding of health and disease.

## References

1. Gnaiger E, Steinlechner-Maran R, Méndez G, et al. Control of mitochondrial and cellular respiration by oxygen. *J Bioenerg Biomembr.* 1995;27(6):583-96.
2. Wikstrom M, Sharma V, Kaila VR, et al. New perspectives on proton pumping in cellular respiration. *Chem rev.* 2015;115(5):2196-221.
3. Babcock GT, Wikström M. Oxygen activation and the conservation of energy in cell respiration. *Nature.* 1992;356(6367):301-9.

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Received: 04-Aug-2024, Manuscript No. AACBM-24-144832; Editor assigned: 06-Aug-2024, PreQC No. AACBM-24-1448325(PQ); Reviewed: 20-Aug-2024, QC No AACBM-24-1448325; Revised: 22-Aug-2024, Manuscript No. AACBM-24-1448325(R); Published: 28-Aug-2024, DOI:10.35841/aacbm-6.4.221

4. Lobritz MA, Belenky P, Porter CB, et al. Antibiotic efficacy is linked to bacterial cellular respiration. *Proc Natl Acad Sci.* 2015;112(27):8173-80.
5. Songer CJ, Mintzes JJ. Understanding cellular respiration: An analysis of conceptual change in college biology. *J Res Sci Teach.* 1994;31(6):621-37.
6. Wegrzyn J, Potla R, Chwae YJ, et al. Function of mitochondrial Stat3 in cellular respiration. *Science.* 2009;323(5915):793-7.
7. Jafri MS, Dudycha SJ, O'Rourke B. Cardiac energy metabolism: models of cellular respiration. *Annu Rev Biomed Eng.* 2001;3(1):57-81.
8. D'Amico G, Lam F, Hagen T, et al. Inhibition of cellular respiration by endogenously produced carbon monoxide. *J Cell Sci.* 2006;119(11):2291-8.
9. Hill GE. Cellular respiration: the nexus of stress, condition, and ornamentation. *Integr Comp Biol.* 2014;54(4):645-57.
10. Brunori M, Giuffre A, Sarti P, et al. Nitric oxide and cellular respiration. *Cell Mol Life Sci.* 1999;56:549-57.

**Citation:** Paszti E. Cellular respiration: The process powering life. *J Cell Biol Metab.* 2024;6(4):221.