



# Examining Metabolic Enzyme Pathways in Cancer Biology

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## Description

Cancer is a complex and multifaceted disease that arises when normal cellular processes, such as growth and division, become dysregulated. One of the key factors contributing to cancer development is the alteration of metabolic pathways, which provide the energy and building blocks necessary for tumor growth. Metabolic enzymes play a central role in these processes and recent research has highlighted their importance in the progression of cancer. By examining metabolic enzyme pathways in cancer biology, scientists are uncovering new therapeutic targets for cancer treatment.

Cancer cells exhibit altered metabolism, often referred to as the "Warburg effect." This phenomenon, first described by Otto Warburg in the 1920s, involves the preference of cancer cells for glycolysis the breakdown of glucose into pyruvate over oxidative phosphorylation, even in the presence of oxygen. This shift allows cancer cells to generate ATP and biomass at a faster rate, supporting their rapid proliferation. One key enzyme involved in glycolysis is hexokinase, which catalyzes the first step in the conversion of glucose to glucose-6-phosphate. Overexpression of hexokinase has been observed in many cancers, making it a potential target for therapy.

Another important metabolic enzyme in cancer is Pyruvate Kinase M2 (PKM2), which regulates the final step of glycolysis. PKM2 is often upregulated in cancer cells and plays a difficult role in controlling the balance between energy production and biosynthesis. By altering the activity of PKM2, cancer cells can divert glucose metabolism toward the production of macromolecules required for cell division, such as nucleotides and lipids. Inhibiting PKM2 could disrupt cancer cell metabolism and limit tumor growth.

In addition to glycolysis, cancer cells often exhibit alterations in other metabolic pathways, such as the Tricarboxylic Acid (TCA) cycle and fatty acid metabolism. The TCA cycle, which takes place in the mitochondria, is responsible for generating energy and metabolic intermediates. In cancer, the TCA cycle is often dysregulated, with some tumors showing increased production of lactate, a byproduct of anaerobic metabolism, even in the presence of oxygen. This phenomenon is thought to support the anabolic needs of rapidly growing tumor cells.

Fatty acid metabolism is also altered in cancer, as tumor cells require an increased supply of lipids for membrane synthesis and energy storage. Fatty Acid Synthase (FASN) is an enzyme involved in the synthesis of fatty acids and its overexpression has been linked to several types of cancer. Targeting FASN could reduce lipid availability and slow tumor growth. The alteration of metabolic enzyme pathways in cancer biology provides an opportunity for novel therapeutic strategies. By targeting key enzymes involved in metabolic processes, researchers are developing drugs that can selectively disrupt cancer cell metabolism. These therapies aim to "starve" tumors of the nutrients and energy they need to grow, while sparing normal cells. Overall, the study of metabolic enzyme pathways in cancer biology has opened up new avenues for cancer treatment. By understanding how metabolic alterations drive tumor growth, scientists are identifying new ways to target cancer cells and improve patient outcomes. As research in this area continues, it is likely that metabolic therapies will become an integral part of the cancer treatment landscape.

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