

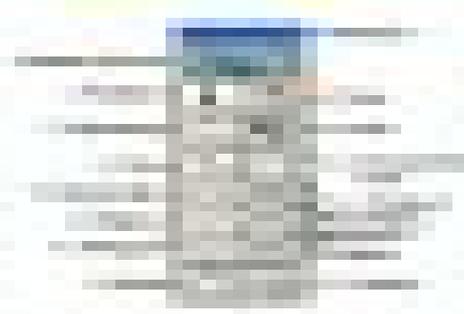
Pharmacokinetics of Insulin 7



Insulin is secreted by the beta cells of the pancreas. The release of insulin is stimulated by the presence of glucose in the blood. Insulin then acts on target tissues to promote the uptake of glucose and the synthesis of glycogen and fat. The half-life of insulin in the blood is approximately 5-10 minutes. Insulin is rapidly degraded by the enzyme insulinase in the liver and kidneys. The overall clearance of insulin is high, resulting in a short half-life. This necessitates the frequent administration of insulin to maintain blood glucose levels.

The pharmacokinetics of insulin are influenced by several factors, including the route of administration, the formulation of the insulin, and the individual's metabolic state. Insulin administered subcutaneously has a slower onset of action compared to intravenous insulin. The onset of action of subcutaneous insulin is typically 30-60 minutes. The duration of action of subcutaneous insulin is typically 2-4 hours. The peak concentration of subcutaneous insulin is typically reached 1-2 hours after administration. The overall pharmacokinetics of insulin are highly variable, and individualized dosing is often required to achieve optimal glycemic control.

Insulin is a peptide hormone that is secreted by the beta cells of the pancreas. The release of insulin is stimulated by the presence of glucose in the blood. Insulin then acts on target tissues to promote the uptake of glucose and the synthesis of glycogen and fat. The half-life of insulin in the blood is approximately 5-10 minutes. Insulin is rapidly degraded by the enzyme insulinase in the liver and kidneys. The overall clearance of insulin is high, resulting in a short half-life. This necessitates the frequent administration of insulin to maintain blood glucose levels.



The subcutaneous route of administration is the most common route for insulin. It is a relatively painless and easy-to-use route. The insulin is absorbed into the bloodstream through the subcutaneous tissue. The rate of absorption is slower than intravenous administration but faster than oral administration. The overall pharmacokinetics of subcutaneous insulin are highly variable, and individualized dosing is often required to achieve optimal glycemic control.

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