Introduction

In the history of insulin therapy some milestones are present. In 1921, the young physician Frederick Banting (1881-1941) and the fourth year medical student Charles Best (1899-1978), working in the laboratory of Prof. Mclod in the Toronto University, found the final link in a series of studies begun in 1916 by other researchers that guessed the pancreas secretion of substance, already named “insulin”, capable to decrease blood glucose concentrations. The young researchers isolated this pancreas extract that “cured” hyperglycemia in diabetic dogs1-2 and in 1922 they successfully administered it for the first time to a 14-year-old diabetic patient, Leonard Thompson3.

In 1923, the company Lilly began marketing animal insulin. In 1928, the hormone was identified as a protein. Since the insulin of Banting and Best could not function for more than 6 h, much research went into finding ways of prolonging action. In 1936, Hagedorn noted that addition of a basic protein, such as protamine, to the insulin preparation, kept the hormone in suspension at the injection site, delaying absorption and prolonging its action4. In 1946, the first insulin “Isophane NPH” (Neutral Protamine Hagedorn), obtained combining insulin and protamine in stoichiometric quantities (hence the term isophane from the Greek equal and manifest) at neutral pH, was marketed5. In 1952, the first Lente insulin, retarded with zinc and without protamine, was produced in Denmark6. In 1955, Sanger determined the exact formula of bovine insulin7. The sixties saw the development of a radioimmunoassay for insulin8. In the seventies, production of mono-component insulins, obtained by ion exchange chromatography and of monopeak ones, separated by Sephadex G50 column, began. High performance chromatography made it possible to isolate an insulin 99% pure from proinsulin and other islet hormones9.

Abstract. – Although the improvement on insulin therapy since it was first conceived, it is still far from mimicking physiological secretion of pancreatic b-cells and research to find new insulin formulations and new routes of administration continues. Human biosynthetic insulin (rapid-acting, intermediate-acting and long-acting), produced by recombinant DNA technique, is currently available. The pharmacokinetic profile of rapid-acting insulin (regular) does not adequately reproduce the physiological post-prandial insulin response. This has led to the development of molecular analogues with slight modifications that prevent the spontaneous polymerisation underlying delayed absorption. Fast-acting analogues such as Lyspro and Aspart can be injected immediately before the meal, inducing a very fast and substantial peak of insulin, similar to that produced by b-cells, but have the disadvantage of short duration of action. For this reason, and because of the difficulty of obtaining sufficient basal insulin concentrations to control pre-prandial blood glucose levels with current long-acting insulins, analogues known as Glargine and Detemir have been synthesized. They have virtually no plasma peak and acts for about 24 h. These characteristics make it ideal to cover basal insulin requirement. With insulin analogues, it also seems possible to overcome the problem of intra- and inter-individual variability in absorption after subcutaneous injection. This variability is directly proportional to the duration of insulin action. Research into new routes of administration has led to production of inhaled insulin powder, soon to become commercially available. Insulin is absorbed through the lung alveoli. Trials to evaluate efficacy and tolerance have shown that inhaled insulin has a similar kinetic profile to the fast-acting injected analogue and can therefore be used for mealtime requirement, combined with a single daily injection of long-acting insulin. Oral insulin is currently being studied in type 1 diabetes prevention with promising results.

Key Words:
Human insulin, Lyspro, Aspart, Glargine, Inhaled insulin, Oral insulin.
Human Insulin

Human insulin is a globular protein with a molecular weight of about 5,800 kd, consisting of 51 amino acid residues organised in two polypeptide chains (A and B), linked by two disulphide bonds. Chain A consists of 21 residues with an extra disulphide bond between A6 and A11; chain B consists of 30 aminoacids. Complete synthesis of the human insulin molecule was achieved in 196610.

Insulin exists as a monomer only at low concentrations while it shows propensity to aggregate into stable dimers at higher concentrations, in aqueous solution at pH 2-8 and into hexamers in the presence of zinc ions. The hexamer, in which chain A constitutes much of the polar surface, is almost spherical in structure, with a diameter of 5 nm and a height of 3.5 nm. Polymerisation of the hormone has major pharmacological implications.

Before the eighties, insulin was extracted from bovine or pig pancreas. Commercial production of human insulin began in 1982. This milestone in the history of diabetes is of similar scientific relevance to the discovery of insulin itself. Once it became clear that it was impossible to obtain sufficient human insulin for world requirements from cadavers, the techniques for synthesising insulin were developed. Semisynthetic insulin was obtained by an enzymatic method, in which trypsin catalyses the substitution of alanine, in position B30 of pig insulin, with threonine. The insulin thus produced has the same aminoacid sequence that the human one and it was completely free of other pancreatic and gastroenteric hormones11-12.

Genetic engineering has enabled industrial synthesis of human insulin by the technique of recombinant DNA. Fragments of DNA coding for chains A and B are inserted separately in plasmids and then in a special strain of Escherichia coli which acts as a microlaboratory for in vitro synthesis. The insulin chains released into the medium are harvested (together with other bacterial proteins) and reassembled with the appropriate disulphide bonds13-15. To avoid contamination during the various steps of the procedure, plasmids were produced with genetic material coding for the proinsulin molecule, which is isolated from the culture medium and deprived of its connecting peptide16-18. Biosynthetic insulin can be also produced by another technique. The procedure exploits the yeast Saccharomyces cerevisiae as a living laboratory by inserting purified DNA that dictates the synthesis of a single chain precursor different from proinsulin in one of the yeast’s plasmids. The protein, produced by yeast continuously during fermentation, is harvested from the culture medium, isolated by centrifugation and crystallisation, converted to insulin ester by trypsin transpeptidisation and finally hydrolysed. This method guarantees high purity12,19-20.

Human insulins currently on the market include rapid-acting (Regular), intermediate-acting (NPH and Lente) and long-acting (Ultralente) formulations. The former is a clear, colourless, aqueous solution buffered at neutral pH (7-7.8). Meta-cresol is added as preservative, glycerol as tonic stabiliser, as well as zinc chloride. Hexamers, made stable by zinc ions, are the predominant quaternary structure of pharmacological insulin; other structures include dimers and tetramers21.

The molecule tends to aggregate in the vial and in the tissue where it is injected. The hexamers need to be split for the insulin absorption from the subcutaneous injection site. For this reason, Regular insulin enters the general circulation after a lag phase about 30 min after injection. It achieves the plasmatic peak at 2 to 4 h and lasts for 6 h.

Absorption of NPH is delayed by protamine, a protein extracted from the nucleus of fish sperm, where its role is to stabilise DNA. The commercial form is Isophane-NPH insulin, a white suspension of orthorhombic crystals containing 0.9 molecules of protamine and two atoms of zinc per hexamer5. In the crystal, protamine regulates interactions between dimers and hexamers. The vehicle is water buffered at pH 6.9-7.5. Phenol or meta-cresol are added as preservatives21. Insulin crystals are insoluble in water and tend to precipitate to the bottom of the vial, which has to be tipped various times to resuspend them before use22. NPH is longer-acting; in fact its blood absorption begins 1.5 h after subcutaneous injection; it has a peak plasma concentration at 4 to 12 h and disappears within 24 h. The “tail”, however, is relatively ineffective. Although NPH has the most regular absorption of all intermediate- and long-acting insulins, inter- and intra-individual variability is high.
Lente, delayed without addition of protamine, is obtained by precipitating the hormone in the presence of zinc salts. When its molar ratio with respect to insulin is greater than one, the zinc ion reduces the solubility of insulin in neutral solvent. When an acid solution of insulin is brought to pH 7.4 with an excess of zinc ions, the resulting precipitate is amorphous and has moderately delayed absorption after subcutaneous injection. This kind of insulin formulation is known as Semilente, and was produced until several years ago. If insulin is allowed to crystallise at pH 5.5 before zinc is added and pH corrected, the obtained formulation is known as Ultralente. This insulin consists of crystals insoluble in water which remain in suspension and are absorbed very slowly. A 3:7 mixture of amorphous and crystalline insulin, known as Lente, has an intermediate absorption profile. In commercial preparations, methyl-parahydroxybenzoate is used as preservative and the suspension is buffered at pH 7-7.8. It appears in the circulation 2.5 h after injection, has the peak plasma concentration at 7 to 15 h and disappears by 20 h. Both NPH and Lente insulins can be mixed with Regular insulin. However, while NPH and Regular conserve their pharmacokinetic characteristics, Lente/Regular mixture changes in time because the excess of zinc ions binds part of the rapid-acting insulin, transforming it into a form similar to Semilente.

Long-acting insulin (Ultralente) is an aqueous suspension of zinc-insulin crystals at neutral pH, of milky appearance. It has an onset of action after 4 h, achieves a slight but undesirable peak plasma concentration at 7 h and sustains blood insulin levels approximately 8 to 20 h; it inconsistently mimics endogenous basal secretion. Moreover its absorption is completely irregular and it cannot be mixed with Regular insulin in the syringe because the action of the latter would be excessively delayed.

### Insulin Analogues

Since the nineteen-eighties, pharmacological research has been concentrated on analogues of human insulin, seeking to develop molecules with minimal modifications and appropriate kinetics. In 1996, an analogue in which the aminoacids in positions B28 and 29 were inverted, was marketed. The inversion conferred the property of prompt absorption and hence fast action. The analogue is known as Lyspro, from lysine and proline, the aminoacids transposed. The handling of the molecule prevents its natural tendency to hexamers aggregation, since position B28 is crucial for its spatial configuration. Because the interaction between B28 and B23, two molecules of insulin align antiparallel forming a non polar dimer which in turn aggregates into hexamers by linking to zinc in pharmacological preparations. Since modification of aminoacid residue B28 counteracts polymerisation of insulin, the analogue may be injected immediately before the meal, as it acts in a few minutes. Lyspro maintains a plateau for 1 to 2 h and disappears by 3 to 4 h. Need for availability of insulin analogue was dictated by three pharmacokinetic defects typical of Regular insulin:

1. Its relatively slow absorption causes lacking reproduction of physiological profile of the b-pancreatic secretion.
2. The comparison of the insulin curve in a normal subject to that in a type 1 diabetic patient (injected with Regular insulin) shows the following figures: (a) the absence of the first peak, (b) the flat overall profile in the first two hours after the meal (leading to early post-prandial hyperglycemia) and (c) the persisting raised insulin plasma levels during post-absorptive period.
3. This pattern may lead to pre-prandial hypoglycemic events and conditions the prescription of snacks for type 1 diabetic patients.

A lot of Italian, European and American trials carried out after experimentation and consolidated use of insulin Lyspro indicate that the post-prandial insulin profile after the analogue administration is very close to the physiological one. Comparison with the insulin profile after an injection of Regular insulin (even when the patient waits the prescribed 30 min before eating) confirms the higher validity of Lyspro to mimic physiological glucose-stimulated insulin se-
cretion. In clinical practice, this phenomenon should lead to more physiologic glycemia 90-120 min after the meal. However, while the short duration of action allows for better control over post-prandial glucose levels, it does not satisfy, alone, the basal insulin requirement in diabetic patients with low endogenous insulin reserve (type 1). In order to use the fast-acting analogue in the daily therapeutic protocol of the type 1 diabetic subject, a combination of Lyspro and intermediate-acting insulin is therefore necessary. On the other hand, this imposes the use of the syringe to mix insulins or the practice of a simultaneous double injection by the “pen”, which is the simplest and more pleasant actual method for self-administration of insulin. Until Lyspro becomes available in “penfills” with a premixed preparation of the analogue (25, 50 and 75%) and intermediate-acting insulin (neutral protamine Lyspro, NPL, 75, 50 and 25%) (expected soon), it cannot completely replace Regular insulin, but it can only coexist for a more a versatile use. For example, it can be administered to type 2 diabetic patients requiring insulin due to secondary failure of oral hypoglycemic agents. In this case it can be used at mealtimes, in association to intermediate- or long-acting insulin at bed-time, if necessary. The pharmacokinetic of Lyspro suggests that type 2 diabetics, characterised by lacking first peak of insulin secretion, can mostly benefit from the analogue. It may also be successfully used in those patients with iatrogenic diabetes due to steroids, in whom hyperglycemia occurs selectively after meals, or in those patients with diabetes associated to liver disease having the same feature. It may be useful in type 1 diabetic subjects, educated to manage “free” variations in diet, for spot correction of occasionally glucose peaks, detected during self-monitoring (well illustrated in the “pizza, coca-cola and tiramisù” study)36. A nother application is in the first period after diagnosis, when a residual basal insulin secretion is consistent. A interesting indication is provided by patients who have an uncertain food intake (intercurrent pathologies, gastroparesis diabeticorum...) In these subjects, Lyspro can be injected immediately after the meal at a dose calculated on the basis of the quantity of food eaten, without deteriorating post-prandial glucose levels.

In 2000, another analogue of human insulin, Aspart, became available. This analogue is obtained by substitution of the proline residue in position B28 with an aspartic acid residue38. This substitution eliminates the interaction with glicine B23 that triggers polymerisation. Monomer status is also aided by repulsion between the aspartic acid residue and the nearby glutamic acid B21, both positively charged. Since the variation introduced in Aspart is not close to the receptor site, there is no change in receptor binding affinity39 or in the rate of intracellular dissociation40-41 with respect to native insulin. This manipulation has also solved the problem of affinity for IGF-1 receptors. The delay in making Aspart available was due to the intense mitogenic activity (due to excessive affinity for the IGF-1 receptor) of the originally proposed analogue with aspartic acid in B10 position39.

There are many advantages of therapy using fast-acting insulin analogues. The need to limit post-prandial hyperglycemia, both in type 1 and 2 diabetics, emerges from recent studies with large cohorts, the results of which unequivocally correlate poor metabolic control with the development and progression of complications. Specifically, post-prandial hyperglycemia seems a major factor for the cardiovascular manifestations so frequent in type 2 diabetics42-44. This is the rationale for another indication of insulin therapy in type 2 patients, using analogues. Restoration of the early insulin peak seems to be particularly important. In fact, correcting hyperglycemia 90-120 min after the meal, it is possible to reduce late hyperinsulinemia and, as a consequence, the insulin resistance highly related to the development of cardiovascular complications45-46. Pre-prandial and nocturnal hypoglycemia can also be removed by fast-acting insulin37. Finally, the possibility of injecting the hormone immediately before the meal, without having to wait 30 min, greatly improves the quality of life, especially for young patients who are dependent on injections throughout their life. However, type 1 diabetic subjects still require simultaneous administration of intermediate- or long-acting insulin.

The pharmaceutical formulation of fast-acting insulin is the same as that of regular insulin.
Long-Acting Analogues

All available intermediate- and long-acting insulins (NPH, Lente, Ultralente) have two major handling problems. The first is that all the classical formulations have wide variations in absorption between different individuals, and even in different injection sites and at different moments in the same subject, in a manner which is directly proportional to the duration of their action. This phenomenon is small, but present and significant with Regular insulin; it is greater with NPH, Lente and greatest of all with Ultralente, which is rarely used for this reason. The second problem is their time-action profile. After a delay of a few hours, their absorption gives rise to an insulin peak and a lowering effect on blood glucose, the major cause of nocturnal hypoglycemia, often unawareness, which afflicts many insulin-treated patients. The peak is followed by a stable plateau, which, in some subjects, ends in a small, but present and significant with NPH, Lente and greatest of all with Ultralente, which is rarely used for this reason.

The need for an ideal long-acting insulin, that gives rise to a square wave of plasma concentration, or in other words, constant basal insulin levels, has prompted research into long-acting analogues. The aim of the research is to obtain an insulin formulation, which can be combined with a fast-acting analogue to ensure a closer parallel to physiological b-cell secretion.

Candidate analogues for daily use are: (i) soluble analogues, created by varying the electrical charge of the insulin molecule and hence its isolectric point, which affects solubility in vivo (insulin Glargine) and (ii) acyl analogues in which a fatty acid molecule is linked to an aminoacid residue so as to bind with albumin in the blood, forming a "depot" product that gradually releases insulin into the tissues (insulin Detemir).

Glargine

In 2001, insulin Glargine has become commercially available. This long-acting analogue is produced from human DNA recombinant insulin, modified by the addition of two arginine residues, conferring a positive charge, to the C-terminal (B30a and B30b) of the molecule. This addition shifts the isolectric point (pI) from 5.4 to 6.7, making the molecule more soluble at the acid pH of the vial and less soluble at the neutral pH of subcutaneous site. The acidity of the medium in which insulin is conserved dictated another modification of the molecule, namely substitution of asparagine in position A21 with glycine. This prevents deamidation and confers stability. It has been demonstrated that Glargine interacts with insulin receptors in a similar way to human regular insulin and also that its dissociation is similar; it does not have mitogenic effects due to activation of IG-F-1 receptors.

Pharmacological studies have been carried out with the iso- and euglycemic clamp techniques to evaluate the action profile of Glargine in normal and diabetic subjects. Its absorption rate has been measured in type 2 diabetics and compared with that of insulin NPH: the mean 25% disappearance times of 27.9%56. The area under the blood insulin curve between 0 and 6 h after administration of Glargine was significantly less than that of NPH57. Maximum plasma concentration in the first 4 h was 5.11 IU/l for Glargine and 10.8 IU/l for NPH58. A recent study in type 1 diabetic patients without endogenous insulin reserve (C-peptide negative) confirmed that the action of Glargine lasts 24 h and that it gives no plasmatic insulin peak, whereas NPH and Ultralente achieve significantly high plasma levels a few hours after injection. Moreover, NPH lasts only 12 h and the disadvantage of Ultralente is its enormous variability in absorption. Compared to NPH, Glargine therefore shows a constant increase, followed by a stable plateau, which, in some patients, lasts almost 24 h. All these features indicate that the pharmacokinetical characteristics of Glargine meet the criteria necessary to ensure basal insulin requirement in type 1 diabetics with negligible plasma C-peptide concentrations. Many clinical trials show that Glargine provides good control of basal glycemic levels59-64.
Glargine insulin is a clear liquid (like fast-acting insulin) with a pH of 4. After injection, it forms a microprecipitate in the tissues, ensuring slow and steady absorption. Unfortunately, it cannot be mixed with fast-acting formulations and this is a problem since the number of daily injections increases when therapy protocol is based on the introduction of Glargine.

Stability of Insulin in Commercial Preparations

Agents as heat, repeated and strong shaking, exposure to hydrophobic surfaces (such as drip attachments) cause changes in insulin conformation leading to linear aggregation and formation of insoluble fibrils. This phenomenon mainly affects intravenous infusions (drips required in acute situations) and continuous subcutaneous infusions by minipump used for intensive insulin therapy. Many measures have been devised to avoid it, such as addition of glycerol, albumin, patient’s serum or blood, but none of them have proved practical, physiological and safe. At temperatures above 25°C, chemical destabilisation due to deamidation or polymerisation may occur. Current insulin formulations ensure stability of the hormone below 25°C for a month. Vials for daily use can therefore be kept at room temperature and supplies must be kept at 2-8°C, where their physical and chemical characteristics are maintained for a period of about 30 months.

Absorption

Insulin must be administered parenterally because digestive enzymes destroy it. When injected s.c., insulin must be split into monomers before it enters circulation. The lymphatic system only plays a secondary role in absorption of the hormone. Less than 5%, a clinically insignificant amount, is broken down at the injection site. As already mentioned, the problem of intra- and inter-individual variability in insulin absorption is very significant and can cause a variation of 10-52% in the amount of insulin absorbed daily, responsible for excursions in glycemia by as much as 80%. This problem has not yet been solved.

Many factors influence the absorption rate of insulin.

Injection site

The abdominal region offers faster and more reproducible absorption (T/2=87 min) than the arm (140 min), buttocks (155 min) and thighs (165 min)72. This difference is of clinical significance and is linked to regional differences in blood flow. Absorption of insulin monomers is correlated with available capillary area, explaining why muscle contraction (which causes vasodilation) around the site of a recent injection, significantly speeds onset of the glucose lowering effect of the hormone. Similarly, high room temperature or sauna facilitate Regular but not long-acting insulin to be absorbed; the mechanism involved may not be related to changes in blood flow but rather to accelerated diffusion in the tissues. The fast action of analogues is maintained irrespective of injection site25.

Injection depth

If the injection is too deep and does not distribute the insulin solution or suspension in the subcutaneous tissue but rather in the muscle, absorption is faster due to the greater vascularisation of muscle with respect to subcutaneous fat. If the injection is too superficial (in the derma), the process is slower and incomplete.

Concentration

Injection of smaller volumes of insulin solution, corresponding to higher concentrations, improves absorption. The influence of concentration has been eliminated by standardising all commercial preparations to 100 IU/ml.

Massage of injection site

Massage with cotton wool soaked with disinfectant should be avoided because it increases the absorption rate by a mechanism apparently not related to blood flow73.

Distribution and Elimination

Intravenous insulin has a half-life of 4 minutes. If administered subcutaneously, it is distributed in the plasma phase in the free form,
that is, not bound to proteins, as does endogenous insulin, unless the organism has produced a significant quantity of antibodies against it. These antibodies act as a buffer system, binding insulin with a capriciously reversible bond. In normal subjects, insulin has a distribution volume of 85 ml/kg. It reaches specific receptors in target tissues. It is eliminated by breakdown in the liver (60-80%), kidney (10-20%), muscle and adipose tissue (10-20%). A small amount is excreted in the urine. Paradoxically, significant impairment of renal function reduces its clearance more than severe liver disease. The residence time in subcutaneous tissue is brief for fast-acting analogues, making their action profile short. Subcutaneous deposit of Regular and long-acting insulin is significant and is achieved after repeated administrations. The size of deposit is related to the number of injections/die. For example, a dose of 40 IU/die in two injections, is calculated to deposit 22-60 IU; the same dose in multiple doses deposits 12-20 IU and continuous infusion deposits 1-14 IU. This explains why patients taking few doses/die are protected for longer against rises in blood glucose levels, whereas patients undergoing continuous infusion cannot even tolerate brief interruptions without developing ketosis.

**Alternative Routes of Insulin Administration**

Since 1924 researches about alternative routes of insulin administration (oral, nasal, rectal, transdermal) were reported, but until recently no realistically useful preparations had been obtained. Of all these routes, inhalation is the one most tested greatly. The rationale behind inhalation is the passage of protein molecules, including insulin, across the mucosa of the lung alveoli, which have a surface area of 100 m². Insulin has shown a particular aptitude for this route. A pharmacological formulation of the hormone for inhalation will soon become commercially available. It is a dry powder dosed in mg, in blisters containing 1 or 3 mg of insulin in 5 mg of excipient (mannitol, glycine and sodium citrate powder). The blisters are opened and emptied directly into the inhalation apparatus. One or two inhalations (2-6 mg) are generally the appropriate dose for countering post-prandial hyperglycemia in a type 1 diabetic patient with poor endogenous insulin reserve. Pharmacokinetic studies in normal and diabetic subjects show that its absorption rate is similar to that of injected analogue. A plasma peak is achieved 5-60 min after inhalation, compared to 60-80 min for Regular insulin. It is therefore administered immediately before the three meals. Its duration of action is intermediate between analogue and Regular. A mean daily requirement of 12 mg is equivalent to 350 IU of soluble insulin. The divergence between these doses and those usually administered by injection of soluble or suspended insulin is due to a series of factors that cause dispersion.

1) The first loss of hormone is in the inhalation apparatus, but 80-95% reaches the oral cavity.
2) The second loss is the quantity that deposits on the mucosa of the mouth and pharynx on the way to the alveoli.
3) Lung tissue metabolises part of the insulin, which is not absorbed. The efficiency of absorption is calculated at 20-40%, in other words, 6-8 out of 10 insulin molecules do not enter circulation.

Many trials have been conducted with type 1 and 2 diabetic patients, comparing traditional therapy with inhaled insulin at meal-times associated to a subcutaneous Ultralente injection at bedtime. A Ll unequivocally found the similarity in the hypoglycaemic effect of the inhaled insulin in respect to subcutaneous route. In a 3-month study, inhalation was so well tolerated and accepted that 80% of patients chose to continue inhalation therapy for a year. However there are still some doubts as to whether this route is suitable for all patients. The oral route has been neglected for some time because of the difficulty to keep insulin intact in the gastric environment and to get it across the barrier of the intestinal mucosa. It is currently used in experimental studies for
the prevention of type 1 diabetes mellitus, exploiting its property of modifying the immune response underlying insulitis by contact with the tolerant mucosa of the digestive system. Moreover, at the annual meeting of the American Society for Clinical Pharmacology and Therapy, Still recently presented a hexyl-insulin monoconjugate (HIM2) in which an alkyl-polyethylene-glycol group attached to lysine B29 confers stability in the gastric environment. The phase II trials of oral insulin have been presented at American Diabetes Association annual meeting. Oral administration of insulin would be a breakthrough for obvious reasons of patient convenience and also because it restores the primary role of the liver as manipulator of insulin, a role lost with the introduction of subcutaneous administration. Under physiological conditions, b-cell hormone is captured by the liver which metabolises about 40% and then releases the remaining 60% into the peripheral circulation, whereas injected insulin is absorbed directly into the general circulation, causing excessive impregnation of the peripheral tissues. This is a major aspect in which insulin replacement therapy diverges from what happens in nature.

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