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An Address

DIABETES AND INSULIN

BEING THE NOBEL LECTURE DELIVERED AT STOCKHOLM ON SEPTEMBER 15TH, 1923

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Gentlemen:

I VERY deeply appreciate the honour which you have conferred upon me in awarding the Nobel prize for 1923 to me and Professor J. J. R. Macleod.—I am fully aware of the responsibility which rests upon me to deliver an address in which certain aspects of the work on insulin may be placed before you. This I propose to do to-day and I regret that an earlier opportunity has not been afforded me of satisfying this obligation.

Since von Mering and Minkowski proved that removal of the pancreas produced severe and fatal diabetes in dogs, physiologists and clinicians have frequently endeavoured to obtain from the pancreas an internal secretion which would be of value in the treatment of diabetes mellitus. Beginning with Minkowski himself many observers tried various forms of extracts of the pancreas. Among the extracts used were water, saline, alcohol and glycerin. The extracts thus obtained were administered by mouth, subcutaneously, intravenously or by rectum, both to experimental animals and humans suffering from diabetes. Little or no improvement was obtained and any favourable results were overshadowed by their toxic effects. In 1908, Zuelzer tried alcoholic extracts on six cases of diabetes mellitus and obtained favourable results, one case of severe diabetes becoming sugar free. His extracts were then tried by Forschbach in Minkowski's clinic with less favourable results, and the investigation was abandoned by this group of workers. Rennie found that the islet cells ex-

isted separate from the acinar cells in certain bony fishes and in conjunction with Fraser extracts of the principal islet cells were tried both on animals and on the human. Their results, however, were not sufficiently convincing to warrant clinical application. The problem of the extraction of the antidiabetic principle from the pancreas was then taken up for the most part by physiologists among whom were Scott, Paulesco, Kleiner and Murlin.

While these efforts were being made by the physiologists valuable knowledge was being gained on carbohydrate metabolism. Lewis and Benedict, Folin and Wu, Schaffer and Hartman, and Ivar Bang had elaborated methods whereby the percentage of sugar in a small sample of blood might be accurately estimated. At the same time a vast amount of knowledge was accumulating on basal metabolism. Special attention was being given to the relative importance of various foodstuffs, and emphasis was being put on dietetic treatment of diabetes. Guelpa, Van Nordam, Allen, Joslin and Woodyatt, had elaborated systems of diabetic diet.

On October 30, 1920, I was attracted by an article by Moses Baron, in which he pointed out the similarity between the degenerative changes in the acinus cells of the pancreas following experimental ligation of the duct, and the changes following blockage of the duct with gallstones. Having read this article the idea presented itself that by ligating the duct and allowing time for the degeneration of the acinus cells, a means might be provided for obtaining an extract of the islet cells free from the de-



stroying influence of trypsin and other pancreatic enzymes.

On April 14, 1921, I began working on this idea in the Physiological Laboratory of the University of Toronto. Professor Macleod allotted me Dr. Charles Best as an associate. Our first step was to tie the pancreatic ducts in a number of dogs. At the end of seven weeks these dogs were chloroformed. The pancreas of each dog was removed and all were found to be shrivelled, fibrotic, and about one-third the original size. Histological examination showed that there were no healthy acinus cells. This material was cut into small pieces, ground with sand, and extracted with normal saline. This extract was tested on a dog rendered diabetic by the removal of the pancreas. Following the intravenous injection the blood sugars of the depancreatized dogs were reduced to a normal or subnormal level, and the urine became sugar free. There was a marked improvement in the general clinical condition as evidenced by the fact that the animals became stronger and more lively, the broken down wounds healed more kindly, and the life of the animal was undoubtedly prolonged.

The beneficial results obtained from this first type of extract substantiated the view that trypsin destroyed the antidiabetic principle and suggested the idea that by getting rid of the trypsin an active extract might be obtained. The second type of extract was made from the pancreas of dogs in which acinus cells had been exhausted of trypsin by the long continued injection of secretin. Although many of the extracts made in this manner produced marked lowering of blood sugar and improvement in the general clinical condition it was not always possible to completely exhaust the gland, consequently toxic effects frequently resulted.

The third type of extract used in this series of experiments was made from the pancreas of foetal calves of less than four months development. Laguesse had found that the pancreas of new-born contained comparatively more islet cells than the pancreas of the adult. Since other glands of internal secretion are known to contain their active principle as soon as they are differentiated in their embryological development, it occurred to me that trypsin might not be present since it is not used till after the birth of the animal. Later I found that Ibrahim had shown that trypsin is not present till

seven or eight months of intrauterine development. Foetal extracts could be prepared in a much more concentrated solution than the former two varieties of extract. It produced marked lowering of blood sugar, urine became sugar free and there was marked clinical improvement. Its greatest value however was that the abundance in which it could be obtained enabled us to investigate its chemical extraction.

Up to this time saline had been used as an extractive. We now found that alcohol slightly acidified extracted the active principle, and by applying this method of extraction to the whole adult beef pancreas active extracts comparatively free from toxic properties were obtained.

Since all large scale production methods for the preparation of insulin to-day have the acid-alcohol extraction as the first step in the process it may be well to elaborate on the methods of preparation at this stage. Insulin was prepared by the extraction of fresh glands with faintly acid alcohol. The concentration of alcohol in the original experiments varied from forty to sixty per cent. The alcoholic solution of pancreas was filtered and the filtrate concentrated by evaporation of the alcohol and water in vacuo or in a warm air current. Lipoid material was removed by extracting the residue with toluene or ether. The resulting product was the original whole gland extract. We were able to show that the active material contained in this extract was practically insoluble in 95 per cent alcohol. The extracts prepared in this way were tried on depancreatized dogs and in all cases the blood sugar was lowered. In one early case hypoglycæmic level was reached and the dog died from what we now know to be a hypoglycæmic reaction.

It had been known that depancreatized dogs were unable to store glycogen in the liver, and that glycogen disappears in three or four days after pancreatectomy. We found that by the administration of glucose and extract the diabetic dog was enabled to store as much as 8 per cent to 12 per cent glycogen. Diabetic dogs seldom live more than twelve to fourteen days. But with the daily administration of this whole gland extract we were able to keep a depancreatized dog alive and healthy for ten weeks. At the end of this time the dog was chloroformed and a careful autopsy failed to reveal any islet tissue.

The extract at this time was sufficiently purified to be tested on three cases of diabetes mellitus in the wards of the Toronto General Hospital. There was a marked reduction in blood sugar and the urine was rendered sugar free. However the high protein content rendered the continuous use undesirable, due to formation of sterile abscesses.

At this stage in the investigation, February, 1922, Professor Macleod abandoned his work on anoxæmia and turned his whole laboratory staff on the investigation of the physiological properties of what is now known as insulin. Dr. Collip took up the biochemical purification of the active principle and ran the scale of fractional precipitation 70 per cent to 95 per cent alcohol and succeeded in obtaining a more improved end product. But unfortunately his method was not applicable to large scale production. Dr. Best then took up the large scale production and contributed greatly to the establishment of the principles of production and purification. This work was carried out in the Connaught Laboratories under Prof. Fitzgerald who is kind enough to be here to-day. It had been found that the final product obtained by the earlier methods was not sufficiently pure for prolonged clinical use and efforts were made to secure a better product. The benzoic acid method of Maloney and Findlay which depends upon the fact that insulin is absorbed from watery solutions by benzoic acid, was successfully used in Connaught Laboratories for several months.

Professor Shaffer of Washington University, St. Louis, and his collaborators, Somogyi and Doisy, introduced a method of purification which is known as the isoelectric process. This method depends upon the fact that if a watery solution of insulin is adjusted to approximately $\text{pH } 5$ a precipitate settles out which contains much of the potent material and relatively few impurities. Dudley has found that insulin was precipitated from water solutions by picric acid and he made use of this fact to devise a very ingenious method for the purification of the active material.

Best and Scott who are responsible for the preparation of insulin in the Insulin Division of the Connaught Laboratories have tested all the available methods and have appropriated certain details from many of these, several new

procedures which have been found advantageous have been introduced by them. The yield of insulin obtained by Best and Scott at the Connaught Laboratories, by a preliminary extraction with dilute sulphuric acid followed by alcohol, is 1,800 to 2,200 units per kg. of pancreas.

The present method of preparation is as follows. The beef or pork pancreas is finely minced in a large grinder and the minced material is then treated with 5 c.c. of concentrated sulphuric acid, appropriately diluted, per pound of glands. The mixture is stirred for a period of three or four hours and 95 per cent alcohol is added until the concentration of alcohol is 60 to 70 per cent. Two extractions of the glands are made. The solid material is then partially removed by centrifuging the mixture and the solution is further clarified by filtering through paper. The filtrate is practically neutralized with NaOH . The clear filtrate is concentrated in vacuo to about one-fifteenth of its original volume. The concentrate is then heated to 50°C which results in the separation of lipid and other materials, which are removed by filtration. Ammonium sulphate (37 gms. per 100 c.c.) is then added to the concentrate and a protein material containing all the insulin floats to the top of the liquid. The precipitate is skimmed off and dissolved in hot acid alcohol. When the precipitate has completely dissolved, ten volumes of warm alcohol are added. The solution is then neutralized with NaOH and cooled to room temperature, and kept in a refrigerator at 5°C for two days. At the end of this time the dark-coloured supernatant alcohol is decanted off. The alcohol contains practically no potency. The precipitate is dried in vacuo to remove all trace of the alcohol. It is then dissolved in acid water, in which it is readily soluble. The solution is made alkaline with NaOH to $\text{pH } 7.3$ to 7.5 . At this alkalinity a dark-coloured precipitate settles out, and is immediately centrifuged off. This precipitate is washed once or twice with alkaline water of $\text{pH } 9.0$ and the washings are added to the main liquid. It is important that this process be carried out fairly quickly as insulin is destroyed in alkaline solution. The acidity is adjusted to $\text{pH } 5.0$ and a white precipitate readily settles out. Tricesol is added to a concentration of 0.3 per cent in order to assist in the isoelectric precipitation and to act as a preservative. After

standing one week in the ice chest the supernatant liquid is decanted off and the resultant liquid is removed by centrifuging. The precipitate is then dissolved in a small quantity of acid water. A second isoelectric precipitation is carried out by adjusting the acidity to a P_H of approximately 5.0. After standing overnight the resultant precipitate is removed by centrifuging. The precipitate, which contains the active principle in a comparatively pure form, is dissolved in acid water and the hydrogen ion concentration adjusted to P_H 2.5. The material is carefully tested to determine the potency and is then diluted to the desired strength of 10, 20, 40, or 80 units per c.c. Tricesol is added to secure a concentration of 0.1 per cent. Sufficient sodium chloride is added to make the solution isotonic. The insulin solution is passed through a Mandler filter. After passing through the filter the insulin is retested carefully to determine its potency. There is practically no loss in berkefelding. The tested insulin is poured into sterile glass vials with aseptic precautions and the sterility of the final product thoroughly tested by approved methods.

The method of estimating the potency of insulin solutions is based on the effect that insulin produces upon the blood sugar of normal animals. Rabbits serve as the test animal. They are starved for twenty-four hours before the administration of insulin. Their weight should be approximately 2 kg. Insulin is distributed in strengths of 10, 20, 40 and 80 units per c.c. The unit is one-third of the amount of material required to lower the blood sugar of a 2-kg. rabbit which has fasted twenty-four hours from the normal level (0.118 per cent) to 0.045 per cent over a period of five hours. In a moderately severe case of diabetes one unit causes about 2.5 grammes of carbohydrate to be utilized. In earlier and milder cases, as a rule, one unit has a greater effect, accounting for three to five grammes of carbohydrate.

With the improvement in the quality of insulin, the increased knowledge of its physiological action and the increased quantities at our disposal, we were now prepared for more extensive clinical investigation. In May 1922, a clinic was established in association with Dr. Gilchrist, at Christie Street Hospital for returned soldiers. Following this, a clinic was established in the Toronto General Hospital in asso-

ciation with Drs. Campbell and Fletcher, and at Toronto Hospital for Sick Children in association with Dr. Gladys Boyd. In general the routine followed in all these clinics was as follows.

After a careful history had been taken, the patient was given a complete physical examination. Special attention was directed to the finding of foci of possible infection. The teeth, tonsils, accessory sinuses, chest and digestive system were examined clinically, as well as by x-ray. Special consideration was given to biliary tract infection, constipation and chronic appendicitis. If any source of septic absorption was located it was appropriately treated, since such conditions may lower carbohydrate tolerance. If indicated the eye grounds were examined for a possible diabetic retinitis or neuro-retinitis. The daily routine urinalysis included the volume of the twenty-four hour specimen, the specific gravity, the reaction, and tests for albumen by heat or nitric acid. The acetone bodies were estimated by means of the Rothera and ferric chloride tests. Sugar determinations were done by means of the Benedict qualitative and quantitative solutions. In addition to the above, the blood sugars were estimated by means of the Schaffer-Hartman method and the respiratory quotients with the Douglas bag and Haldane gas analysis apparatus. At first the patient continued on the same diet as that previous to his admission to hospital in order to obtain some idea of the severity of his case, and to avoid complications from sudden change of diet. Coma will be discussed separately. On the second or third day he was placed upon a diet the caloric value of which was calculated on his basal requirement. This was determined from Dubois' chart and Aub-Dubois' table. It has been estimated by Marsh, Newberg, and Holly that the body requires two-thirds of a gram of protein per kilogram, (1 kilo = 2.2 pounds), of body weight per day, in order to maintain nitrogenous equilibrium. The remaining calories must be supplied by carbohydrate and fats in a ratio that will prevent the production of ketone bodies.

The patient remained on this basal requirement diet at least a week. During this time, blood sugar was estimated before, and three hours after, breakfast, in order to determine the fasting level and the effect of food. The quantity of sugar excreted was estimated daily, and

this amount subtracted from the available carbohydrate ingested gives approximately the utilization. The available carbohydrate includes fifty-eight per cent of the protein, ten per cent of the fat, and the total carbohydrate in the diet. It may be noticed that when a patient was placed upon a diet in which the protein, fat and carbohydrates were balanced, that the amount of sugar excreted soon approached a fairly constant amount, whereas if the diet was not well adjusted to the patient's requirements, there was wide variation in the amounts of sugar excreted. If a patient became sugar-free and blood sugar normal on a basal requirement diet the caloric intake was gradually increased until sugar appeared in the urine. The tolerance was thus ascertained. If a patient remained sugar-free and had a normal blood sugar when on a diet containing five hundred calories above his basal requirement he was not considered sufficiently severe for insulin treatment, since five hundred calories over and above the basal requirement are sufficient for daily activities. If, however, he was unable to metabolize this amount, insulin treatment was commenced.

Diabetes mellitus is due to a deficiency of the internal secretion of the pancreas. The main principle of treatment is, therefore, to correct this deficiency. If it is found that the patient is unable to keep sugar-free on a diet that is compatible with an active, useful life, sufficient insulin is administered to meet this requirement. In severe cases insulin was administered subcutaneously three times a day, from one-half to three-quarters of an hour before meals. This was done so that the curve of hypoglycæmia produced by the Insulin was superimposed on the curve of hyperglycæmia produced by the meal. In rare cases a small fourth dose was given at bed time to control nocturnal glycosuria. The less severe cases could be satisfactorily treated on a morning and evening dose or a single dose before breakfast. When the insulin treatment was established, if sugar was present in the twenty-four hour specimen of urine, the dosage was gradually raised till the patient became sugar-free. If he was not receiving sufficient food for maintenance, diet and dosage of insulin were gradually raised. If small quantities of urinary sugar persist, it was desirable to find out at what period of the day this was excreted. In order to do this, each spec-

imen in the twenty-four hours was analysed separately. An increase in the dose previous to the appearance of glycosuria will prevent its occurrence.

In severe cases it was found preferable to give the largest dose of insulin in the morning, and reduced doses throughout the day. For example, a patient may receive fifteen units in the morning, ten units at noon and ten units at night. If three equal doses are given there may be morning glycosuria and evening hypoglycæmia, whereas the extremes of blood sugar causing these conditions may be prevented by the above distribution. The effect of the same dosage of extract on different individuals was found to vary considerably. Five patients, whose weights varied from forty-six to sixty-seven kilograms, each received two cubic centimetres of the same lot of Insulin, and in four hours the blood sugars had decreased 0.012 per cent, 0.044 per cent, 0.128 per cent, 0.146 per cent, and 0.0180 per cent respectively. It was found, however, that one patient would persistently give marked decreases in blood sugar after insulin, while in another the fall in blood sugar was persistently less. In our experience, the more marked decreases in blood sugar occurred in the milder cases. The blood sugars of some of the patients were followed throughout the twenty-four hours and it was found that it was possible to gauge the dosage of insulin so as to keep the blood sugar within normal limits and still avoid the dangers of hypoglycæmia.

Coincident with the maintenance of the blood sugar at normal level the cardinal symptoms of the disease disappear. The patient loses the irritating thirst and dryness of the mouth and throat, and does not desire the large amounts of fluid with which he had previously tried to combat these symptoms. The lowered fluid intake diminishes the polyuria and from a twenty-four hour excretion of three to five litres the output falls to normal. The appetite which has been voracious is now satisfied with a normal meal, the carbohydrate of which is utilized, and the patient loses the persistent craving for food.

We found that when a patient was given too large a dose of insulin there was a marked reaction, and the hypoglycæmia which developed gave rise to symptoms which were very similar to those observed in animals. The reaction began in from one and a half to six hours after

the patient received the overdose. The average time was three to four hours. The interval varied with the individual, the dosage, and the food ingested. The first warning of hypoglycæmia was an unaccountable anxiety and a feeling of impending trouble associated with restlessness. This was frequently followed by profuse perspiration. The development of this symptom was not affected by atmospheric conditions. It appeared while the patient was in a frosty outside atmosphere, or in a heated room, and was independent of physical or mental activity. At this time there was usually a very great desire for food. No particular foodstuff was desired, but bulk of any kind seemed to give satisfaction. At times the appetite is almost unappeasable.

At this stage of the reaction the patient noticed a certain sensation as of clonic tremor in the muscles of the extremities. This could be controlled at first. Coordination, however, was impaired for the more delicate movements. Coincident with this there was a marked pallor of the skin with a rise in pulse rate to one hundred or one hundred and twenty beats per minute, and a dilatation of the pupils. The blood pressure during this period fell about fifteen to twenty-five millimetres of mercury, and the patient felt faint. The ability to do physical or mental work was greatly impaired. In a severe reaction there was often a considerable degree of aphasia, the patient having to grope for words. The memory for names and figures became quite faulty.

The onset of hypoglycæmic symptoms depends not only on the extent, but also on the rapidity of fall in blood sugar. The level at which symptoms occur is slightly higher in the diabetic with marked hyperglycæmia than in a patient whose blood sugar is normal. When the blood sugar is suddenly reduced from a high level, premonitory symptoms may occur with a blood sugar between the normal levels of 0.100 per cent and 0.080 per cent while the more marked symptoms of prostration, perspiration, and incoordination develop between 0.080 per cent and 0.042 per cent. As a patient becomes accustomed to a normal blood sugar the threshold of these reactions becomes lower. One patient who formerly had premonitory symptoms of hypoglycæmia at 0.096 per cent now has no reaction at 0.076 per cent but symptoms commence between this level and 0.062 per cent. The ingestion of carbo-

hydrate, in the form of orange juice, (four to eight ounces), or of glucose, relieves these symptoms in from one-quarter to one-half hour. If the reaction is severe, or if coma or convulsions occur, epinephrin or intravenous glucose should be given. The former acts in from three to ten minutes, but in order that the symptoms should not recur glucose must be given by mouth as soon as the patient has sufficiently recovered. The patients were warned that when these reactions occurred they were to obtain carbohydrate immediately.

“Fats only burn in the fire of carbohydrate.” The ability of the severe diabetic to burn glucose is markedly impaired, therefore the excess of fat is incompletely oxidized, giving rise to ketone bodies. These appear in the blood and urine as acetone, diacetic and betaoxybutyric acids. Insulin causes increased carbohydrate metabolism, and consequently fats are completely burned. This is substantiated by the fact that acetone and sugar disappear from the urine almost simultaneously following adequate amounts of insulin. When insulin is discontinued in these cases, acetone bodies and sugar reappear in the urine.

Since the Rothera test is exceedingly delicate, (sensitive to 1 part of aceto-acetic acid in 30,000), patients on a high fat diet may be sugar-free and still show traces of acetone bodies. A comparison with the ferric chloride test, (which is sensitive to only 1 part in 7,000) is, therefore, desirable. The persistence of ketone bodies in amounts which can be determined by the ferric chloride test necessitates either an increase in the carbohydrate or a decrease in fat of the diet.

When the production of acetone bodies is more rapid than the excretion they accumulate in the blood, giving rise to air hunger, drowsiness, and coma. The need of insulin is then imperative. After its administration the utilization of carbohydrate by the body gives complete combustion of the fats. When a patient was admitted to hospital in coma the blood sugar tests and a urinalysis were done as soon as possible. (The urine was obtained by catheterization if necessary.) While these tests were being carried out the large bowel was evacuated with copious enemata. If the blood sugar was high and acetone present in large amounts in the urine, from thirty to fifty units of insulin were

given subcutaneously. Blood and urinary sugar were frequently estimated because of the danger of hypoglycæmia. To prevent this from thirty to fifty grammes of glucose in ten per cent solution were given intravenously. If the patient was profoundly comatose the insulin was administered intravenously with the glucose.

The patient usually regained consciousness in from three to six hours. From this time on, fluids and glucose were administered by mouth if retained. The patient was urged to take at least two hundred cubic centimetres of fluid per hour. In from eight to ten hours the ketone bodies were markedly reduced. On the following day protein was given every four hours as the white of one egg in two hundred cubic centimetres of orange juice. In two to three days, when ketone bodies had disappeared from the urine, fat was cautiously added, and the patient was slowly raised to a basal requirement diet. He was then treated as an ordinary diabetic. During the period of coma the patient was kept warm and toxic materials eliminated from the bowel by purgation and repeated enemata. A large amount of fluid was given to dilute the toxic bodies and promote their elimination. This was administered intravenously, subcutaneously, or per rectum. If signs of circulatory failure developed these were treated by appropriate stimulation.

Striking results were obtained with the above procedure. However, it was found that the longer the period of untreated coma the more grave was the prognosis and the slower the recovery if it occurred. Cases complicated by severe infection, gangrene, pneumonia, or intestinal intoxication may recover from acidosis and coma, but succumb to the complication. Marked lipæmia was present in three cases. This disappeared in the course of a week to ten days after the patient was placed on insulin and on a diet in which the fat was restricted. The urine of one patient became acetone-free while lipæmia persisted.

The severe diabetic, whose ability to burn carbohydrate is markedly impaired, has a persistently low respiratory quotient, from 0.7 to 0.8, which is but little raised by the ingestion of glucose: when glucose and insulin are given together the respiratory quotient is markedly increased, showing that carbohydrate is being metabolized. The highest values have been ob-

tained when pure glucose was used with insulin. Less extensive rises have been secured when the patient, while on a mixed diet, received insulin. All the patients gained in weight on the additional calories. There was an increase in sexual vigour and there was a greater ability to do mental and physical work. Nearly all of the patients have returned to their former employment, and while still under supervision, they administer their own insulin and arrange their own diets with satisfactory results.

All diabetics who have not an adequate knowledge of the dietetic treatment of their disease should be admitted to hospital in order that they may receive instruction in the preparation of their calculated and weighed diet—that they may learn the qualitative tests for sugar and acetone in the urine — that their carbohydrate tolerance may be accurately determined; and that the use of insulin, if required, may be safely instituted. Mild cases, especially if over fifty years of age can be controlled by diet. Cases that cannot be adequately controlled by dietetic treatment alone should be given sufficient insulin to enable them to attain to a diet on which they may “carry on.”

One of the commonest complications of diabetes especially in untreated patients over fifty is gangrene. It is often associated with varying degrees of sclerosis of the leg arteries, which makes it extremely difficult to obtain healing. This may be accomplished by the use of insulin, but when permanent impairment has occurred it is advisable to amputate. Amputation is also advisable when an infection is so severe that the life of the patient is in jeopardy. Treatment of these cases is difficult because, due to the infection, there is a marked variation in the daily production of insulin by their own pancreas. But with careful treatment they can be rendered free from acetone and sugar and their general condition improved. Operation is then performed preferably under nitrous oxide and oxygen anæsthetic. If the blood sugar is maintained normal and acidosis is prevented the wound heals kindly, provided that the amputation has been high enough to assure a good blood supply. For varying periods after the operation the patient remains on insulin treatment. In nearly all cases at the end of three or four weeks mild hypoglycæmic reactions indicate an overdose of insulin. It is

then necessary to increase the diet or decrease the insulin. In some cases the tolerance improves sufficiently to warrant the discontinuance of insulin.

Diabetic patients requiring major operations, such as appendectomy, cholecystectomy and tonsillectomy, or removal of teeth, are first rendered sugar and acetone-free unless the severity of symptoms demand immediate attention. Patients formerly considered bad surgical risks, if given proper dietetic treatment with insulin may be protected from the acidosis, hyperglycæmia and glycosuria which otherwise usually result from the anæsthetic. In the diabetic, infections such as boils and carbuncles and also intercurrent infections such as bronchitis, influenza, and fevers are favourably influenced by the normal blood sugar and increased metabolism which the administration of insulin permits. In the diabetic with tuberculosis insulin allows the administration of proper nourishment to combat the tubercle infection.

During the past year and a half I have not been in active practice but have remained associated with the clinics. I have also kept in personal touch with the first fifteen patients who received insulin treatment. These patients were all extremely severe diabetics for whom diet had done its best. Of these fifteen patients seven were children under fifteen years. It has been possible through the intelligent co-operation of the parents to continue a proper balance between diet and insulin dosage and to maintain six of the seven children sugar-free. None of these have had to return to hospital, and all have gained in tolerance and require from one-half to one-third less insulin than when they first began treatment. They have all gained in height and weight and for the most part have developed into healthy normal children. The one child whose diet and insulin has not been properly controlled has been back in hospital repeatedly and is steadily losing in tolerance. Of the remaining eight cases there were four women and three men whose ages ranged from twenty-five to thirty-five years. The weight of the women varied from seventy-four to seventy-nine pounds. Two of the women, although they have gained to normal or overweight and now have no symptoms of disease, have not shown any increase in tolerance, due perhaps, to the fact that they have not kept sugar-free. All the others, both

men and women, have been able to reduce their dose of insulin from two-thirds to one-fifth of the original requirement. The one remaining case was admitted for amputation. She had had diabetes for six years, and at the time of admission her blood sugar was 0.350 per cent and large amounts of acetone and sugar were being excreted in the urine. She was rendered sugar- and acetone-free by means of insulin before the operation was performed. Amputation was done at the middle third of the thigh. The stump was entirely healed in three weeks. Within six weeks of her operation insulin was discontinued and her diet was increased without the return of diabetic symptoms. It is now three years since her operation and she is sugar-free on a liberal diet without insulin.

It may be of interest to mention a few cases in greater detail to further illustrate the improvement in carbohydrate tolerance following insulin treatment.

Case 1.—Male, aged twenty-nine years had suffered from chronic appendicitis. The urine of the patient in December, 1916, was sugar-free. About the middle of March, 1917, he suddenly developed polyuria, polyphagia and polydipsia, and lost fourteen pounds in weight in a fortnight. There was marked weakness. Urinary sugar was discovered to be as high as eight per cent at this time. On April 4th, the patient was placed on Allen treatment, and slowly regained a tolerance of about two hundred grammes available carbohydrate. He returned to his army duties in September 1917, and was able to carry on uninterruptedly until March, 1919. His tolerance had decreased during this time to about one hundred and fifty grammes. Following discharge from the army in March, 1919, the course of the patient was slowly downhill until October, 1921, when a particularly severe form of influenza shattered his tolerance. Up to this time the patient was maintained practically sugar-free, but following the attack of influenza his tolerance fell to about sixty-six grammes of available carbohydrate. He began to lose weight rapidly. Thirst, hunger and polyuria returned. His strength diminished and, owing to mental and physical lassitude, he found it impossible to continue his work. Glycosuria became persistent and acetone bodies made their appearance and steadily increased. A distinct

odour of acetone was at times distinguishable on the patient's breath.

On February 11, 1922, this patient was taken to the Physiology Department of the University of Toronto, and the respiratory quotient was found to be 0.74, and unchanged by the ingestion of thirty grammes of pure glucose. Then 5 c.c. of insulin were given subcutaneously, and within two hours the patient's respiratory quotient had risen to 0.90. The urine was sugar-free and he had shaken off his mental and physical torpor. Following this experiment the patient did not again receive insulin until May 15th, as the product was being further improved. Since the latter date, the patient has been constantly on insulin.

During the first six months of insulin treatment it was impossible to maintain him sugar-free, although he received about 120 units per day. However, he gained in weight and his clinical condition improved. About January 1923 with the improvement in the quality of insulin, the patient became sugar-free and has remained sugar-free with the exception of one or two occasions. During the first nine months he required no reduction in the dose of insulin, but since that time on the average of every two months he has had a series of hypoglycæmic reactions which necessitated the reduction of the dose. One exception to this occurred in June 1924 at which time appendectomy was performed following a mild attack of appendicitis. An increased dose was required to maintain him sugar-free during this period. At the present time he requires but 20 units of insulin, or one-sixth of his original requirement. His diet has been practically constant during the whole period of observation. All symptoms attributable to diabetes have long since disappeared. He has gained twenty-five pounds in weight and apart from the necessity of taking insulin and controlling his diet he leads an active normal life.

This case is a striking example of the fact that it is only in cases who are maintained sugar-free over long periods of time that an improvement in tolerance is obtained with a consequent reduction in the dose of insulin.

Case 2.—Female, age fifteen years. In the autumn of 1918 the patient had polydipsia and polyuria and complained of weakness. During the winter she suffered from pains in the legs

and back and from insomnia. In March, 1919, these symptoms became more severe. The appetite became excessive and there was some pruritus. The weight by this time had fallen from seventy-five pounds to sixty-two pounds. Glycosuria was discovered and she was placed under the care of Dr. F. M. Allen, to whom we are very much indebted for complete record of the case from April 1919 till August 1922. During this period the diet was controlled so as to maintain the urine free from sugar. Despite this careful dietetic regime the patient's condition became progressively worse.

When she came under my care on August 16, 1922, the examination showed patient emaciated; skin dry; slight œdema of ankles. Hair brittle and thin. Abdomen prominent. Marked weakness. The patient was brought on a stretcher and weighed forty-five pounds. Nothing of note in the respiratory, cardiovascular, digestive or nervous system.

At this time she was receiving a diet of protein 50 gm., fat 71 gm., carbohydrate 20 gm. (919 calories). Insulin treatment was started immediately. At this early stage the unit of insulin had not been worked out and it is therefore difficult to accurately estimate the dosage she received. The diet was increased daily so that, at the end of two weeks, she was receiving protein 63 gm., fat 208 gm., carbohydrate 97 gm., (2512 calories). This diet was continued up to January 1, 1923. Insulin was given fifteen to thirty minutes before the morning and evening meals. A sufficient amount was given to maintain the urine free of sugar. Each specimen of urine was examined and the dose was increased slightly if traces of sugar appeared. When hypoglycæmia occurred orange juice or glucose candy was given. Between August 16th and January 1st, the urine was sugar free except on ten occasions when traces of sugar appeared, and on two other occasions when less than 2 gm. was excreted. Acetone was absent from the urine.

On this treatment the patient gained rapidly in strength and was soon able to take vigorous exercise. Her weight increased from forty-five to one hundred and five pounds in the first six months. The diet included such foodstuffs as cereals, bread, potato, rice, corn, tapioca, corn starch, and even honey.

At present (June 1925) she is in the best of

health and to use her own words "never felt better in all my life." She has grown four inches and weighs one hundred and thirty-four pounds. Her present diet which is only approximate because she has dispensed with the weighing of food, is 125 gm. carbohydrate, 50 gm. protein, 50 gm. fat. This diet is practically the same as that of December, 1922. The insulin required to maintain her sugar free has been reduced about one-third.

Dr. Gladys Boyd, who is now in charge of the diabetics at the Hospital for Sick Children, Toronto, has been able to follow a number of cases of children under insulin treatment. She has estimated the insulin requirement per 10 gm. of carbohydrate in a number of cases and in general her results show a decided increase in tolerance in all cases in which glycosuria and hyperglycemia are adequately controlled. To illustrate—Case 1, which required 6.9 units per 10 gm. carbohydrate in March 1923, only required 2.6 units in January, 1924. Case 2, which required 7.8 units per 10 gm. in January, 1925, in June 1925, required only 2.8 units. Case 3, which required 6.5 units per 10 gm. in April 1922, required only 3.7 units in January 1925.

From a review of the work Dr. Boyd has found that all the patients had had hyperglycemia or even glycosuria at times, but if such occurrences were only transitory and infrequent, improvement in tolerance occurred. Even short periods of rest to the pancreas by means of balanced diet and insulin resulted in improvement in tolerance. Two of our earliest cases, Fanny Z. and Elsie N. are the only exceptions to this rule. Fanny is to all appearances in the best of health with a blood sugar of 0.3 per cent to 0.4 per cent. She has been admitted in coma four times. During her stay in the hospital she improves but does as she chooses on discharge. Her tolerance is becoming less all the time. Elsie keeps in touch with us but is looked after by another physician. He purposely allows her to have glycosuria at night. She is fine physically, but requires much more insulin than formerly.

Dr. Boyd has also found that in those cases who can handle sufficient food without insulin, although the disease has been kept under control there has not been such striking increase in tolerance. The best evidence that there is regeneration of the pancreas with insulin treat-

ment is provided by Drs. Boyd and Robinson. The following is the case reported by them.

Clinical History: B. N. white, male, aged nine years. Family History: Father and one maternal uncle have diabetes. Diabetes diagnosed in this child when he was two years old. He was placed on a suitable Allen diet, which was strictly adhered to, and for a time did well except for recurrent attacks of dysentery, which lowered his tolerance. Failure to gain in stature or weight in any way commensurate with his age was noted and the general condition became worse each year until he was more or less a chronic invalid with increasingly frequent attacks of acidosis during the last year before starting insulin.

He was admitted to the Hospital for Sick Children, Toronto, the end of December 1922. At this time he was an emaciated dwarf, more or less drowsy and unhappy. His weight was thirty pounds, and his height thirty-nine inches. His tolerance to carbohydrate had decreased until he was unable to utilize 15 gm. of such food. Insulin treatment was started at once and his diet increased to a diet suitable for a boy of his age. Sufficient insulin was given to keep him sugar-free and his blood sugar normal. He was discharged on an adequate diet plus insulin. Progress both in general condition and in improvement of pancreatic function was steady. His tolerance to carbohydrate trebled in the year as shown either by the fact that 30 units of insulin controlled the disease as adequately as 90 units a year before, or, stated in another way, without insulin he could now handle 54 gm. of carbohydrate instead of 15. From a chronic invalid in 1922 he became "the leader of the gang" in 1923. He was killed by fracturing his skull when sleigh-riding. He lived for about three hours after receiving the injury and an immediate post-mortem examination was made. The pancreas was removed within thirty minutes of death.

From this clinical history one might expect the pancreas to show marked degeneration. However, on section there was little sign of degeneration, but on the other hand there was strong evidence to support the view of active regeneration both of acinar and islet tissue. These regenerative changes were more marked in the periphery and smaller lobules of the pancreas than in the central area.

The acinar cells were found to be actively proliferating in cords and clusters forming small lobules in some areas and were in close association with newly formed functioning ducts.

The islets were greatly increased in number, particularly in the periphery, there being about four times as many per field as in the central area. These cells were large but might be overlooked with an ordinary stain. However, they could be identified as islet cells by Bowie's special granule stain. This stain also demonstrated that these cells were almost entirely beta cells and were probably concerned in the increased carbohydrate tolerance. On the other hand those islets in the central areas showed an increased number of cells all in an active state of nutrition, but closely packed together. The special stain showed a normal ratio of alpha and beta cells.

These sections were studied by Bensley, Opie, Allen and others who concurred in the opinion of Drs. Boyd and Robinson. Dr. F. M. Allen, Morristown, N.J., after using insulin for three years states as his belief, "That there has been improvement of tolerance in some cases beyond what was possible without insulin." "This observation is trustworthy only in cases where prolonged strict control of symptoms by diet was previously employed. On the other hand the marked increase of tolerance is limited to a minority of cases and has not proved to be continuous in any of them. In other words the improvement always stops short of a cure. There is certainly no decline of tolerance with the passage of time provided the case is kept under proper control."

This summary is the belief of the most conservative of the outstanding clinicians in the United States engaged in diabetic work on a large scale.

Dr. E. P. Joslin, Boston, Mass., who has one of the largest diabetic clinics in the world, has also found that, "The diabetic who is able to reduce his insulin is the diabetic who is absolutely faithful to diet and restricts gain in weight to a moderate degree."

Joslin and his associates have carefully analysed the gain in weight and height of their thirty-two diabetic children under fifteen years of age. Their conclusions are:

1. The gain in weight of the diabetic child treated with insulin resembles that of the nor-

mal child, but the diabetic child is still under weight for his age, though often not for his height.

2. The increase in height of the diabetic child treated with insulin, though occasionally normal, is usually below that of the normal child. So far he has not grown tall like the normal child, either at the expense of growing thin or while being well nourished.

Of the one hundred and thirty children treated with insulin one hundred and twenty are still living, while of the one hundred and sixty-four who did not receive insulin there are one hundred and fifty-two dead. Of the one hundred and twenty still living forty per cent have either not increased or have actually decreased their insulin. Dr. Joslin believes that if the sixty per cent who have had to increase their insulin had received similar treatment they too would have been able to reduce their insulin.

Sixteen children under ten years of age who have taken insulin under Dr. Joslin's care for an average of two years are all alive and now their duration of life is more than three times the duration of life of diabetic children of similar age treated by Dr. Joslin prior to 1915.

Regardless of the severity of the disease, it has been found that by carefully adjusting the *diet and the dose of insulin, all patients may be artificially administered insulin*. In fact, it is to be strongly advocated, because we have abundant evidence for the belief that there is *regeneration of the islet cells of the pancreas* when the strain thrown upon them by a high blood sugar is relieved. The *increase in tolerance* is evidenced by the *decreasing dosage of artificially administered insulin*. In fact, in some moderately severe cases, the tolerance has increased sufficiently that they no longer require insulin.

Diabetes mellitus may be considered fundamentally as a *disordered metabolism, primarily of carbohydrates, and secondarily of protein and fat*. It is indisputably proven that for normal metabolism of carbohydrate in the body, adequate amounts of insulin are essential. It follows, therefore, that the treatment consists in giving just sufficient insulin to make up for the deficiency in the patient's pancreas. Insulin enables the severe diabetic to burn carbohydrate as shown by the rise in the respiratory quotient following the administration of glucose and in-

sulin. It permits glucose to be stored as glycogen in the liver for future use. The burning of carbohydrate enables the complete oxidation of fats, and acidosis disappears. The normality of blood sugar relieves the distressing thirst and consequently there is a diminished intake and output of fluid. Since the tissue cells are properly nourished by the increased diet, there is no longer the constant calling for food, hence *hunger pain* of the severe diabetic is replaced by *normal appetite*. On the increased caloric

intake, the patients *gain rapidly in strength and weight*. With the relief of the symptoms of his disease, and with the increased strength and vigor resulting from the increased diet, *the pessimistic, melancholy diabetic becomes optimistic and cheerful*. Insulin is not a cure for diabetes; it is a treatment. It enables the diabetic to burn sufficient carbohydrates, so that proteins and fats may be added to the diet in sufficient quantities to provide energy for the economic burdens of life.

Crystalline Insulin—Another Chemical Triumph.—One of the greatest fields of effort study and preparation of pure principles of the for the constructive chemist is the isolation, organs of internal secretion. An understanding of the action of the important glands in the body is never adequate until the chemistry of the active principle (or principles) is known. Although Schäfer and Oliver, Szymonowicz and Cybulski had noted the presence of a vasoconstrictor principle in the suprarenal glands, the fuller appreciation of the secretion came after Abel had succeeded in isolating epinephrine. Subsequently, its chemical structure was determined and the product prepared synthetically, and the science of the therapeutics has been benefited immensely by the availability of pure solutions of definite composition.

The researches of E. C. Kendall leading to the isolation and identification of thyroxin have given new insight into the action of the thyroid; like epinephrine, the pure substance is at the disposal of the physician. The artificial preparation of thyroxin is only a question of time and effort, and "one more outpost of the bulwarks of life will thus have been conquered by the chemist." The extract from the pituitary gland is still the subject of attack; steady progress leading to separation and isolation of the pure substance is attested by the reports of Abel and others who have obtained extremely active solutions. The attack on the ovary, particularly the work of Doisy, lends hope that, even for this gland, chemistry will throw much light on its functions as a secretory organ. Following the fundamental researches of others, Collip has succeeded in preparing

a solution from the parathyroid containing some substances of marked effect on calcium metabolism. A close parallel to the history of epinephrine is that of insulin. The far-reaching publications of Banting, Best and their co-workers on the preparation and use of active extracts of the pancreas are still fresh in our minds. Pharmaceutic houses have been constantly improving the product; yet none has been unconscious of the presence of undesirable impurities of the fact that the methods of standardization at best indicate only approximately the relatively minute amount of that chemical which is the ultimate goal.

In a significant address—the sixth annual Pasteur lecture—recently given before the Chicago Institute of Medicine, John J. Abel made public announcement that he had obtained a crystalline form of insulin and exhibited a small specimen of the product. Lantern slide pictures showed the crystals to be rhombohedral, but other data were not presented at that time. The chemical and medical world will await with great interest further developments, particularly the proof of the identification and chemical constitution of the product. The crystallization of the pure principle "insulin," or a compound of it, may well be considered an outstanding accomplishment in the life of a man already distinguished by his conquests in biochemistry. If following this a better method of standardization is worked out, and the product can be prepared synthetically it will have contributed in great measure to the better understanding of modern physiology, to the increased value of medicine, and to a more enjoyable and prolonged life of the patient with diabetes.—*Jour. A. M. A.*, Jan. 30, 1926.