

MONITORING OF THE REGENERATION PROCESS IN PURULENT-NECROTIC PROCESSES OF THE LOWER EXTREMITIES

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Abstract

Currently, the treatment of diabetes mellitus and its complications remains an urgent problem. Patients with diabetic foot syndrome, a complication of diabetes mellitus, are increasingly losing their quality of life and ability to work. The treatment of diabetes mellitus and its complications affects the economic and financial efficiency of our country. Therefore, the new drug Reomannisol, produced in our country, helps to stop the purulent-necrotic process of the foot, which is a complication of diabetes mellitus.

Keywords: Experimental model of diabetic foot, purulent-necrotic process, experimental animals, alloxan, Reomannisol.

Actually. Diabetes mellitus is a chronic disease caused by the absolute absence of insulin production by the pancreas, which leads to metabolic disorders in the body [6,7,8,17]. Due to the high mortality of patients, scientists around the world are currently developing and using various drugs [13,14,15]. For the use of newly created drugs in animals, the first question is to create their models [14,19,20,21]. One such model is the alloxan diabetes model. The study of pathomorphological aspects of chronic diabetic foot ulcers and purulent-necrotic changes in humans is difficult and not always controllable [8,13,14,15]. The healing of chronic wounds is subject to the general principles and pathophysiological aspects of reparative processes. This process depends on the stage of development of a chronic wound, the depth of the wound, the basic structure of the damaged organ, the general condition of the body, and the type of treatment [14,20,21,22]. Along with diabetes mellitus, the wound microflora plays a role in the reparative and regenerative processes of a purulent wound. What type of local treatment should be used for a chronic purulent ulcer that has developed against the background of diabetes mellitus remains a question [8,13,14,15,16].

After the development of the alloxan model of diabetes, the goal was to develop a new drug to correct violations of hepatoprotective and antioxidant processes in target organs [6,7,19,20]. Reomannisol is a complex drug with antihypoxic, antioxidant, antishock, rheological, detoxifying, diuretic properties. The main active ingredient is succinic acid and mannitol.

Target. Drawing up a technological algorithm for the treatment of purulent-necrotic lesions of the foot caused by diabetes mellitus at the stage after the complex treatment of experimental diabetic foot syndrome.

Materials and methods. Experiments From 2021 to 2022, experimental studies were carried out on 100 white sterile male rats weighing 180-200 g, kept in the vivarium of the Tashkent Medical Academy. All rats were observed every 12 hours day and night, they were given water to drink and kept at a constant temperature of 25-28°C. The experimental animals were divided into 4 groups: group 1 intact (unchanged group); 2nd control group - creation of an experimental model of a diabetic foot using traditional complex treatment against the background of alloxan diabetes; 3rd experimental group - according to the experimental model of diabetic foot - traditional treatment and rheosorbilactyl; group 4 - experimental group No. 2 - traditional treatment and rheomannisol [13].

Results. In the experimental group, the scab was difficult to detach, tissues in places with a slight overlay of fibrin. Swelling in the area of the foot wound continued in the rats of the control group on the 7-8th day, but in the rats of the experimental group it decreased already on the 5th day after the injury. on the 7th day after the wound in rats from the experimental group when using the drug, a significant decrease in the area of wounds ($4.3 \pm 0.05 \text{ mm}^2$; $p < 0.001$) was noted, 2 times than its original wound, and 1.5 times than in rats control group ($6.5 \pm 0.07 \text{ mm}^2$), due to wound contraction and marginal epithelization. The crust from the wound was removed with great difficulty, under it there was a brightly granulating wound defect with pronounced signs of marginal epithelization, far ahead of the control group. Also, in the rats of the experimental group, almost complete closure of the wound defect is observed, it is noted that the wound bleeds well, this indicates an improvement in blood circulation in the wound, which exceeds this indicator over the control group. When examining the histological section of the wound under a microscope, there is a restriction and rejection of the destructive-necrotic tissue on the surface of the wound.

At the same time, in the circumference of the wound, the tissue of the epidermis is compacted and the focus of destruction of the dermis is delimited from the underlying healthy elements of the tissue of the dermis. As part of the destructive tissue, the development of the process of disintegration and disorganization of both cellular and fibrous structures is determined. In the underlying connective tissue of the dermis, a neoplasm of hair follicles, compaction of fibrous structures and an increase in the proliferative activity of histiocytic cells are determined.

On the 7th day after treatment with the traditional method of an experimental model of a diabetic foot in the control group, at the bottom and circumference of the diabetic wound, necrobiotic tissue and acute inflammatory infiltrate are preserved in the form of a structureless tissue mass, in the circumference of which it is infiltrated with leukocyte cells, and a picture of the formation of poorly differentiated (granulation) is also found. tissue, the process of vasculogenesis is very weak (Picture 2). As part of the structureless mass, there are foci of edema and tissue vacuolization, as well as hematoxylin conglomerates and calcification. Thus, the development of microangiopathy phenomena took place, which makes it possible to consider diabetic microangiopathy as one of the morphofunctional manifestations of diabetes

mellitus. By the 10th day in the experimental group, who received traditional treatment and Reomannisol, an independent almost complete closure of the wound defect and hair growth around the wound were noted, vacuolization and increased proliferative activity of cells in the basal layers.

The connective tissue of the dermis is compacted, without signs of an acute inflammatory process, the fibrous structures are transformed into dense, homogeneous, fibrous masses due to disorganization and fusion with each other. In them, cellular structures are localized along the periphery and around the vessels, and they are in a proliferative active state. On the 10th day after the treatment of diabetic foot with Reomannisol, regeneration of the integumentary stratified epithelium is noted in the form of sliding of epithelial cells onto the surface of the wound. At the same time, the epidermis is represented in the regeneration zone by several layers of active and hyperchromic epithelial cells. Under the epidermis, the formation of a basement membrane from loose fibrous structures is determined. In the tissue of the dermis, the presence of foci of reparative regeneration is determined in the form of excessive formation of fibrous, cellular structures that differ from the surrounding tissue by the presence of many vessels and proliferatively active histiocytic cells. Both the epidermis and the dermis show no signs of an acute inflammatory process. In the control group, on the 10th day, a wound defect of about 4.9 ± 0.05 mm² remains. At the bottom and in the circumference of the wound, the formation of inflammatory granulation tissue with foci of necrosis and hemorrhage is noted. The inflammatory infiltrate is represented by proliferative active lymphoid and histiocytic cells, randomly arranged fibrous structures consisting of argyrophilic and collagen fibers. In the center of the inflammatory infiltrate, a defect surrounded by fibrinoid necrosis and a homogeneous fibrous mass is determined. In the circumference of the inflammatory infiltrate, the connective tissue is subjected to edema, loosening with the breakdown of fibrous structures, and cell necrobiosis. The wound healed on the 14th day of the experiment. Thus, hyperglycemia causes microvascular complications due to impaired angiogenesis, which leads to a prolongation of the inflammatory effect of the wound and the period of wound healing. The use of standard wound treatments (iodine and levomekol) and the drug rheomanisol, indicators of planimetric control over the course of the wound process in animals of this group (experimental) are at a consistently high level.

Finding. The rate of healing of wound defects in rats with diabetic foot syndrome in the control group falls on the 14th day, since the periods of resorption and rejection of necrotic tissues in the wound, damage to the vessels of the microvasculature (microangiopathy) are prolonged, and edema is observed for a long time. The wound process against the background of DM is characterized by late formation of angiogenesis, slowing down and impaired maturation of granulation tissue, marginal epithelialization. In the experimental group, in rats, along with the local traditional method of wound treatment, the drug Reomannisol was used intraperitoneally, as a result, wound healing was recorded on the 10th day from the moment the wound was

applied to the foot of the rats. The use of local treatment and Reomannisol can enhance angiogenesis in the early stages of the experiment and restore disturbed microcirculation (neoplasms of blood vessels), increase macrophage response, fibroblast proliferation, maturation and remodeling of granulation tissue and its epithelization, reduce the inflammatory response, which leads to more effective and early healing wound area.

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