Introduction

Skeletal muscle and peripheral nerve damage after compression injury of the limb is a common complication. Long-lasting compression results in direct trauma and necrosis of muscle tissue, followed by ischemic reperfusion injury. The mechanisms underlying ischemia-reperfusion are complex and distinguish two types of interdependent changes – structural and metabolic. Acute circulatory hypoxia of limb tissues remains less studied than structural changes, which are more isolated and understandable.

Muscle damage occurs not only as a result of direct injury to the limb but can occur as a consequence of compartment syndrome or on the background of limb’s blood vessel disorders [2]. Irreversible changes in the skeletal muscles of the extremities begin after 2 hours of ischemia, and after 4 hours, necrosis can reach 40% of muscle tissue [3]. Restoration of blood flow during reperfusion causes secondary tissue damage, as the accumulated cytotoxic metabolites penetrate the tissues, including through structurally damaged vessels of the limb [4]. As a result, edema, biochemical changes, and tissue necrosis progress.

It has been suggested that it is possible to correct the condition of ischemic injured limb muscles by injecting platelet plasma, bone marrow aspirate cells,
or adipose tissue. Research results indicate the potential impact of these suspensions in delaying muscle damage and dystrophy after trauma or ischemia [5, 6, 7, 8, 9]. Therefore, studying the effects of cell technologies on ischemic muscles is relevant.

Materials and Methods

The developed experimental model consists of a reproduction of the combined mechanical and ischemic damage of an extremity’s tissues reached by imposing on a hind leg of a rabbit Chinchilla (weighing 4, 2-4, and 5 kg) a medical elastic plait. The number of tourniquets is 8 per limb, from the middle third of the thigh to the ankle joint. It achieved sufficient pressure and mechanical trauma to the extremity. The duration of limb immobilization was 6 hours. During this time, severe mechanical damage to the superficial skeletal muscle group occurs in the limb. When measuring subfascial pressure using a device “Stryker Intra-Compartmental Pressure Monitor” (USA), it was confirmed an increase in pressure in over 30 mmHg, which was evaluated as a compartment syndrome in the bone-fascial sheaths of the lower leg. After confirmation of the syndrome, autologous platelet plasma, bone marrow aspirate, or SVF was injected into the injured leg. The protocol of receipt, preparation, and injection are described in detail in the article [10].

On days 5, 15, and 30, after removal of the tourniquets, changes in skeletal muscle and sciatic nerve in the injured limbs were histologically examined. The materials were fixed in 10% formalin solution, sealed in paraffin (Leica Surgipath Paraplast Regular, Formula ‘R’, Germany) through isopropanol-chloroform. Micropreparations 6 μm thick, stained with hematoxylin and eosin, were examined under a microscope Olympus BX 51 (Japan).

Directly, before animals were removed from the experiment, an ultrasound examination was performed to detect and assess the volume of necrotized tissue in the limb. To do this, ultrasound recognized δ-entropy in areas of ischemic tissue and muscle in the contralateral limb. In a separate series of experiments, animals were injected with platelet-rich plasma, bone marrow aspirate, or stromal vascular adipose tissue after knowledge of the elastic tourniquet. All manipulations on the animals were carried out by the provisions of bioethics and the protocol of the bioethics commission of the SI Institute of Orthopedics and Traumatology of NAMS of Ukraine; sodium thiopental was used at a dose of 60 mg/kg. All statistical calculations were performed using the nonparametric Kruskal-Wallis test and one-way analysis of variance ANOVA. The differences were considered significant at p<0.05.

Results and Conclusions

According to the results of sonographic studies, a decrease in entropy and after injection of a suspension of bone marrow aspirate concentrate was revealed only on the 5th day (Fig. 1). On days 15 and 30, the entropy level did not differ from the contralateral limb. Because necrosis of muscle tissue has been detected in recent experiments, it has been concluded that ultrasound has value in assessing changes in ischemic damaged limb tissue only in the early stages of injury.

Macroscopic examination revealed significant damage to the muscles of the calf group: edema with a large amount of exudate (up to 2.0 ml), the acute blood supply to blood vessels and hemorrhages, and reduced density and loss of muscle tissue color (from pink to yellow-white) (Fig. 2). Such characteristic damage to the leg muscles was observed on days 5 and 15. On day 30, free exudate was almost absent, but areas of hemorrhage and muscle necrosis were still recorded. After the introduction of platelet plasma or bone marrow aspirate, no visual differences were detected, while the injection of SVF caused encapsulation of adipose tissue and severe angiogenesis.

Analysis of microstructural changes in the leg muscles revealed the dynamics of skeletal muscle disorders. Initially, necrosis of muscle groups accompanied by edema, necrosis of blood vessels, and hemorrhagic infiltration of the muscle interstitium, then connective tissue replaced the area of necrosis and atrophy of muscle fibers with the formation of a scar. On the 15th day, the density of fibrosis areas significantly increased. On the 30th day, it remained unchanged or tended to increase. Thus, if in the intact muscle the amount of connective tissue was 4.74±0.51%, then after ischemia, it was 53.87±1.61%, 52.68±1.47%, and 69.89±0.85%. There were no differences between the groups with mechanical ischemia and groups injected with platelet-rich plasma, bone marrow aspirate concentrate, and SVF. Although on 30th day, the relative density of collagen sites was slightly lower in the first two groups, which requires further study and possibly is not associated with the direct introduction of these funds. An increase in the number of myocyte nuclei was recorded, which indicates the activation of regenerative processes in the muscle fiber in single damaged muscle fibers.

The topographic dependence of muscle’s damage level was revealed. Necrosis occurred in the superficial groups of muscle fibers. Atrophy was revealed in the middle group of muscles, and deep ones remain almost undamaged. Structurally preserved sciatic and femoral nerves were found between the deep muscle groups. Their fascicular structure remained preserved throughout the experiment; distal branches of the peripheral shin nerves had been damaged and
Fig. 1. The level of entropy in the muscles of the ischemic damaged limb
Note: * – reliable for control (contralateral limb)

Fig. 2. Damage to the limb on the 5th (a) and 15th days (b) after mechanically induced ischemia
degenerative changes were found in them. There was no difference between the density of nerve fibers in the sciatic nerve of the injured limb and contralateral. Degeneration of myelinated nerve fibers was revealed in the fasciculus of the tibial nerve on the 15th and 30th days after the onset of compartment syndrome. These changes were isolated, at the level of individual fascicles that did not affect changes in the density of nerve fibers in the sciatic nerve. It is important to note that on the 15th and 30th days in the areas of atrophy, there were signs of activation of regeneration of individual muscle fibers: the appearance of additional myocyte nuclei and the formation of longitudinal groups of satellite cells. This reaction was observed in almost all samples, but the intergroup comparison was more significant after the injection of bone marrow aspirate concentrate.

According to morphological data, we assume that 6-hour mechanically induced ischemia, which includes two factors (trauma and impaired perfusion in the vessels), causes primary necrosis of muscle tissue in the superficial group of limb muscles. Injuries of deep muscles are associated with ischemia and secondary tissue changes. The revealed changes in the peripheral nerves of the tibia are explained either by the action of the tourniquet or by metabolic changes. The reaction to the injection of adipose tissue was accompanied by activation of fibrosis processes, while the suspension of bone marrow aspirate concentrate did not cause changes (inflammation, osteogenesis, etc.), as well as platelet-rich plasma. We assume that bone marrow aspirate concentrate may promote activation of endogenous regenerative processes in damaged skeletal muscles. It is indicated by the results of biochemical studies [11]. Thus, after the injection of bone marrow aspirate concentrate into ischemic leg muscles, there was a significant increase in the activity of catalase and glutathione peroxidase enzymes for 15 and 30 days and, conversely, decreased NO\textsuperscript{2}. This indicates the activation of the antioxidant defense system and the neutralization of the cytotoxic metabolite NO\textsuperscript{2}. The effect of platelet plasma was less significant in terms of ischemic muscle damage, but also affected the reduction of oxidative modification of proteins. Given the multicomponent nature of bone marrow aspirate concentrate, SVF, and platelet-rich plasma, it is difficult to detect the specific effect of specific drugs. One can only speculatively assume the role of the cytokine system of the bone marrow [8, 12] in the composition of the autologous aspirate in reducing the inflammatory process, delaying malnutrition, and activation of regenerative processes in the muscle tissue of the ischemic damaged limb.

Conflict of interest. The authors declare no conflict of interest. This publication has not been, is not, and will not be the subject of commercial interest in any form.

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Пошкодження нервово-м'язового апарату при механічно індукованій ішемії кінцівки (експериментальна модель)

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Резюме. Актуальність. Травматичне та ішемічне пошкодження кінцівки супроводжується ураженням скелетних м'язів та периферійних нервів кінцівки. Динаміка і наслідки ішемічного ураження залишаються маловивченими і потребують корекції. Мета дослідження. Цількісними морфологічними та сонографічними методами дослідити динаміку пошкодження скелетних м'язів кінцівки після травматичної індукованої ішемії та без введення збагаченої тромбоцитами плазми, аспірату кісткового мозку та фракції жирової тканини. Матеріали і методи. В 3 експериментальних серіях кролям моделювали 6-годинну травматичну ішемію кінцівки з та без введення клітинних суспензій. Результати та висновки. Розроблена модель ішемії кінцівки полягає у 6-годинній іммобілізації кінцівки, на яку накладено медициналасний джгут. Дія джгутів викликає високий підфасціальний тиск у кінцівці і некроз поверхневих груп м'язів нижньої та середньої третини стегна і гомілки. За даними сонографії, ε-ентропія пошкоджених тканин на 5-ту добу знижена щодо неущко-дженого ішемією, як і після введення клітин аспірату кісткового мозку. На 15-ту і 30-ту добу рентгенографія не показала різниці між групами порівняння. Динаміка морфологічних особливостей пошкодження тканин кінцівки полягає у некрозі поверхневих груп м'язів, атрофії у середніх та глибоких груп м'язів. Некроз заміщується рубцовою сполучною тканиною, щільність якої зросла у 11-14 разів і у термінах між 5-ю і 30-ю добою не відрізняється. Введення тромбоцитарної плазми, аспірату кісткового мозку та фракції жирової тканини не вплинуло на зміну динаміки розвитку фіброзних змін у ішемічно пошкодженних груп м'язів. Атрофія м'язів супроводжується активізацією ендогенного відновлення поодиноких м'язових волокон, яка має тенденцію до інтенсифікації після введення аспірату кісткового мозку. Середній нерв пошкодженої кінцівки не зазнав структурного пошкодження з огляду глибокої топографії, водночас у нервах гомілки виникають дегенеративні зміни з 15-ї доби.

Ключові слова: ішемія; травма; скелетні м'язи; клітинна терапія.