**RéSUMÉ**

Rhabdomyolyse accompagnant la septicémie staphyloccique

Introduction. La rhabdomyolyse est un syndrome clinique et de laboratoire résultant de lésions musculaires squelettiques avec libération d’un grand nombre de substances intracellulaires de myocytes dans la circulation systémique et endotoxémie. L’entrée des produits de destruction musculaire dans la circulation systémique entraîne le développement de manifestations systémiques, de troubles graves de l’homéostasie, du syndrome de défaillance multiviscérale, y compris une insuffisance rénale aiguë, menaçant souvent la vie du patient.

Cas rapporté. Un cas rare de rhabdomyolyse en présence de septicémie staphyloccique est décrit. La présence de la rhabdomyolyse est confirmée par les résultats d’études pathologiques et anatomiques. Les causes possibles de la rhabdomyolyse dans la septicémie staphyloccique sont discutées ici.

Conclusions. Seuls quelques cas de rhabdomyolyse à fond de septicémie staphyloccique ont été trouvés.

**ABSTRACT**

Introduction. Rhabdomyolysis (RM) is a clinical and laboratory syndrome resulting from skeletal muscle damage, with the release of a big number of intracellular substances of myocytes into the systemic circulation and endotoxemia. The entry of the muscle destruction products into the systemic circulation leads to the development of systemic manifestations, serious disorders in homeostasis, multiple organ failure syndrome, including acute renal failure, often threatening the patient’s life.

Case report. A rare case of RM in a patient with staphyloccocal sepsis (SS) and development of multiple organ failure is described. The diagnosis of RM was confirmed by the results of pathological and anatomical studies. Possible causes of RM in staphyloccocal sepsis are discussed here.

Conclusions. Only a few cases of RM upon SS were found in the literature. The development of RM in combination with the systemic inflammatory reaction syndrome in SS exacerbates the course of the disease and worsens its prognosis. Therefore, the timely diagnosis of RM and appropriate treatment are extremely important. The mechanism of RM development upon...
SS is complex. After staphylococcus was isolated from the affected muscles in a patient with SS, one of the first descriptions of RM upon SS suggested the role of direct invasion of the infectious agent into the muscles, followed by the development of "pyomyositis." Another mechanism for RM occurrence of PM may be the presence of toxin upon the staphylococcal toxic shock syndrome in patients with SS.

**Keywords:** rhabdomyolysis, staphylococcal sepsis, infective endocarditis, acute renal failure.

**List of abbreviations:**
- RM: rhabdomyolysis
- SS: staphylococcal sepsis
- ARN: acute renal failure
- IE: infectious endocarditis
- CPK: creatine phosphokinase
- ALT: alanine aminotransferase
- AST: aspartate aminotransferase

**Introduction**

Rhabdomyolysis (RM) is a clinical and laboratory syndrome resulting from skeletal muscle damage, with the release of a large number of intracellular substances of myocytes into the systemic circulation (myoglobin, lysosomal and mitochondrial enzymes, histamine, serotonin, oligo- and polypeptides, etc.) and endotoxemia. The entry of the muscle destruction products into the systemic circulation leads to the development of systemic manifestations, serious disorders in homeostasis, multiple organ failure syndrome, including acute renal failure (ARN), often threatening the patient's life.

The various causes of RM (traumatic injuries, metabolic disorders, epilepsy, excessive muscle stress, drugs, etc.) specifically include various infections\(^1\)\(^-\)\(^3\). A search in the Medline system for the publication of RM cases with infections over 30 years (1966-1996) returned information on viral and bacterial infections as the cause of RM in 59 and 60 cases, respectively\(^4\),\(^5\). Recent publications of the last century cover two cases of RM upon SS: in one patient with acquired immunodeficiency syndrome\(^6\), and in another – with staphylococcal pneumonia\(^7\). It is noteworthy that in almost 30 years after the first descriptions of RS upon SS, there were no publications on this topic, and only in the second millennium did individual information reappear. Thus, a report was published on the development of RM in a 12-year-old child with staphylococcal IE, manifested by myalgia, fever, and secretion of dark urine\(^8\). In adults, two cases of RM upon IE caused by methicillin-sensitive staphylococci have been described. One patient previously operated for a congenital malformation developed IE complicated by severe RM with high CPK indices (up to 49068 IU), successfully treated with antibiotics followed by surgical intervention\(^9\). In another patient with IE, RM was diagnosed in combination with purpura, purulent pericarditis and intracerebral hematoma. Despite intensive therapy (antibiotics, drainage of the pericardial cavity and cerebral hematoma), a fatal outcome occurred\(^10\).
In this article, we present a patient with severe RM that developed against the background of SS.

**CASE PRESENTATION**

A 46-year-old man was admitted to the neurological department of the State Clinical Hospital „S.S. Yudin“ (Moscow, Russian Federation) on June 28, 2019, with complaints of pain in the lower spine, and weakness in the legs. The medical history included type 2 diabetes mellitus, for which he received insulin preparations of short and prolonged action, nephrectomy for urolithiasis, polytrauma (fracture of ribs, cranio-cerebral trauma, fractures in the vertebra, contusion of the liver and spleen) at the age of 20 years.

The condition at admission is satisfactory. The consciousness is clear. Forced position on the back. Vesicular breathing, without wheezing, breath rate 16/min. The heart sounds are rhythmic, heart rate 80/min, blood pressure 120/80 mm Hg. The abdomen is soft and painless on palpation. The liver and spleen are not enlarged. No acute neurological and surgical pathology, or foci of infection were found.

We noted the changes of laboratory tests in dynamics. Peripheral blood showed leukocytosis (32-35 x 10^9/L) with a sharp shift to the left (17-37% of stab neutrophils) to the appearance of young forms in the blood (myelocytes, metamyelocytes). Hemoglobin and platelet counts were within the normal range. Biochemical blood tests revealed an increase in the level of C-reactive protein (CRP) (302-440 mg/L), procalcitonin (5.3-17.02 ng/mL), AST and ALT activity (118 IU/L and 101 IU/L, respectively), creatine phosphokinase (2200-2700 IU/L), urea (16-23 mmol/L), creatinine (153-234 mmol/L), potassium (5.2-5.5 mmol/L), glucose (20 mmol/L). During his stay in the neurological department, the patient’s condition deteriorated, with sharp pain in the muscles of the limbs, pain on palpation of the muscles, disorders of the central nervous system (lethargy, confusion), a decrease in the blood pressure to 90/70 mm Hg. Myoglobin (14350 ng/mL) was detected in the blood. Brown urine secretion was marked. The patient was transferred to the intensive care unit on July 01, 2019, where intubation of the trachea, and catheterization of the internal jugular vein were performed, followed by artificial ventilation, hemodiafiltration, and no-radrenaline infusion.

Ultrasound of the soft tissues of the medial surface of the upper third of the thigh revealed an increase in echogenicity, without clear delineations, with single an-echogenic inclusions (edema?, infiltrate?). No fluid accumulation detected in soft tissues. Electrocardiogram showed episodes of atrial flutter, arrested by cordarone. The values of urea (45.3 mmol/L), creatinine (332 µmol/L), CPK (4809 U/L), and lactate dehydrogenase (729 U/L) were increasing. The microbiological examination of the blood and urine revealed Staphylococcus aureus sensitive to methicillin. Antibacterial therapy (oxacillin, linezolid) was prescribed. Despite intensive therapy, the condition deteriorated progressively and the patient died on July 11, 2019.


The pathological diagnosis after necropsy was: Sepsis. Septic shock: shocked kidneys, liquid blood in the cavities of the heart and large vessels. Bilateral focal-confluent pneumonia in 6-9 segments. Multiple organ failure syndrome: parenchymal dystrophy of the liver, myocardium, kidneys. Myocardial infarction in the posterior wall of the left ventricle, with dimensions of 2.5x3.0 cm about one day old. Focal necrosis of the ileum. Acute erosion of the stomach and duodenum. Pulmonary edema. Cerebral edema. Attention was drawn to the change in the appearance of striated muscles, especially the femoral and iliac muscles, which were mottled, with uneven blood supply and flabby (“boiled meat” appearance) (Fig. 1 a,b).

**DISCUSSION**

The presence of a systemic inflammatory reaction syndrome (marked leukocytosis with a shift to myelocytes, a high-level of CRP), the results of the microbiological study (Staphylococcus aureus in the blood and urine), and a high-level of procalcitonin indicated the development of SS, although the primary site of infection could not be defined. Peculiarities of SS progression in the form of myalgia syndrome in combination with elevated CPK, ALT, AST, the presence of myoglobin in the blood, brown urine, and ultrasound data on soft tissues indicated the diagnosis of RM, as a rare complication of SS. This clinical and laboratory syndrome raised the suspicion of polymyositis, which had to be differentiated from RM in various pathological processes. Similar difficulties in the differential diagnosis were reported in one of the RM case descriptions upon SS in a patient with IE, successfully treated with antibiotics, followed by the disappearance of RM signs^{14}. Muscle lesion was confirmed by autopsy and subsequent histological examination.

Only a few cases of RM upon SS were found in the literature, and in the second millennium, only three observations were published^{12-14}. The development of RM in combination with the systemic
inflammatory reaction syndrome in SS exacerbates the course of the disease and worsens its prognosis. Therefore, the timely diagnosis of RM and appropriate treatment are extremely important.

The mechanism of RM development upon SS is complex. After staphylococcus was isolated from the affected muscles in a patient with SS, one of the first descriptions of RM upon SS suggested the role of direct invasion of the infectious agent into the muscles, followed by the development of „pyomyositis.“ Another mechanism may be the presence of toxins upon the staphylococcal toxic shock syndrome in patients with SS15.

Among other toxic mechanisms not related to the toxic shock syndrome, are the staphylococcal enterotoxins associated with the development of RM in a patient with SS, who was found to have the genes encoding staphylococcal enterotoxins C, G and I. One of the mechanisms of myocyte damage upon infections may be the release of interleukin-1 from cells, which is, moreover, a pyrogen during fever development16. The role of the increased content of intracellular calcium entering the cells under the action of prostaglandins and leading to cell destruction activated by neutral proteases is not excluded17. Various additional risk factors may contribute to the development of RM upon SS (statins, alcohol consumption, etc)18. In our patient, one additional factor may be considered the presence of diabetes mellitus.

CONCLUSIONS

One of the RM complications is renal damage, with the development of ARF19. The incidence of ARF reaches 40% among all cases of RM20. RM is the cause of 15% of ARF cases21. The main pathogenetic mechanism of renal damage upon RM is the appearance of myoglobin, having a nephrotoxic effect, in the blood. Furthermore, the fluid redistribution into the muscles results in hypovolemia, leading to renal hypoperfusion. Attempts to predict the risk of developing ARF in patients with RM based on the values of CPK, creatinine, potassium, calcium, and the content of myoglobin in urine were inconclusive20. An additional factor for the development of ARF in our patient could be the nephrectomy in the past.

The treatment of SS patients complicated by RM includes, along with adequate antibacterial therapy, the correction of hypovolemia, electrolyte imbalance (acidosis, hyperkalemia, hypercalcemia), and the hemostatic system, alkalization of urine in order to prevent the nephrotoxic effect of myoglobin, as well

Fig. 1 a,b. Iliac muscles of the mottled flabby appearance with uneven blood supply. 
as the methods of replacement therapy for ARF (hemodialysis)².

Author Contributions:
L.I.D., E.V.S., and S.S.B. were responsible for the diagnostic procedures, clinical diagnosis, and treatment decisions. G.O.Z. made the histopathological diagnosis. L.I.D. and M.A.K. analyzed the literature and wrote the manuscript. All authors have read the text and reached an agreement for the manuscript text.

Compliance with Ethics Requirements:
"The authors declare no conflict of interest regarding this article"

"The authors declare that all the procedures and experiments of this study respect the ethical standards in the Helsinki Declaration of 1975, as revised in 2008(5), as well as the national law. Informed consent was obtained from the patient included in the study"

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References