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Functional Features Of Platelet Hemostasis In Athletes-Athletes 18-35 Years.

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ABSTRACT

General physical activity can strongly affect all organs and systems of the human body, including the physiological state of the blood cells. This is connected with the fact that in certain diseases it is possible to regulate the degree of activity of platelet functions by physical exertion. This required clarification of the aspects of the influence of long-term regular intense physical exertion in young people on platelet activity in vitro and in vivo and the severity of the functioning of the mechanisms that implement their aggregation function. In the course of the study, it was found that candidates and masters of sports in athletics 18-22 years old have consistently low lipid peroxidation against the background of high activity of the antioxidant platelet system. At the age of 26-35 years old, when switching to irregular workouts, stable normal indicators of antioxidant activity of platelets and a low level of lipid peroxidation in them were noted. The regular training of candidates and masters of sports in athletics, regularly practicing and remaining after 22 years of age, showed a stable low functional activity of platelets. This is largely due to the constancy on their platelets the number of receptors for aggregation inducers.

Keywords: athletics, sport candidates and masters of sport, platelets, lipid peroxidation, physical activity.

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INTRODUCTION

It is known that physical activity can affect all organs and systems of the human body [1,2], including the physiological state of the blood cells [3,4]. There is evidence that in certain diseases it is possible to regulate the activity of platelet functions with physical activity [5,6,7].

At the same time, in the case of long-term regular intense physical exertion in young people, the features of platelet activity in vitro and in vivo and the severity of functioning of the mechanisms that implement their aggregation function are not fully clarified [8,9]. For this reason, the purpose of the study was set: to establish the activity of platelets in healthy candidates and masters of sport in athletics at the age of 18-35 years.

MATERIAL AND METHODS

The research was approved by the Ethics Committee of Russian State Social University (record №5 from 12.05.2014).

The observation group included 125 candidates and masters of sports in track and field athletics 18–22 years old (25 people 18 years old, 26 people 19 years old, 23 people 20 years old, 24 people 21 years old and 27 people 22 years old) and 66 candidates and masters of sport in athletics 26-35 years old, who regularly trained until 22 years old, and now have switched to irregular training, but at least once a week (21 people 26-27 years old, 23 people 30-31 years old, 22 people 34-35 years old).

In all candidates and masters of sports, intra-platelet lipid peroxidation (LPO) was determined by the magnitude of the basal level of malondialdehyde (MDA) in the thiobarbituric acid reduction reaction and by the concentration of acylhydroperoxide (AHP). The functional readiness of intra-platelet catalase and superoxide dismutase was recorded.

The observed levels of platelets in capillary blood were determined using a Goryaev chamber. Platelet aggregation (AP) was recorded by a visual micromethod with a number of inductors of inductors: ADP (0.5×10^{-4} M), collagen (dilution 1: 2 of the main suspension), thrombin (0.125 units / ml), ristomycin (0.8 mg/ml), adrenaline (5×10^{-6} M) and their combinations (ADP and adrenaline, ADP and collagen, adrenaline and collagen) in similar concentrations. The state of intravascular platelet activity was revealed using a phase contrast microscope. Statistical processing of the results obtained by the t-student criterion.

RESEARCH RESULTS

In the sportsmen examined, the physiological and biochemical values taken into account were within the limits of the physiological norm.

The content of AHP in the platelets of candidates and masters of sports 18 years of age regularly practicing was 1.71 ± 0.18 D₂₃₃/10⁹ platelets, not significantly changing until 22 years old (1.66 ± 0.16 D₂₃₃/10⁹ platelets). At the same time, the level of MDA in platelets of 18-year-old athletes was 0.37 ± 0.12 nmol/10⁹ platelets, without experiencing reliable dynamics up to 22 years of age (0.39 ± 0.28 nmol/10⁹ platelets). The number of primary products of LPO-AHP in the platelets of candidates and masters of sports of 26-27 years old, who regularly trained until 22 years, was 1.73 ± 0.16 D₂₃₃/10⁹ platelets, not changing significantly by 34-35 years (1.80 ± 0.24 D₂₃₃/10⁹ platelets). However, the level of MDA in platelets in 26-27 year olds surveyed was 0.41 ± 0.22 nmol/10⁹ platelets, while also remaining unchanged up to 34-35 years of life (0.44 ± 0.30 nmol/10⁹ platelets).

The functional readiness of catalase and superoxide dismutase in platelets in the examined candidates and masters of sports at age 18 amounted to 10500.0 ± 214.5 IU/10⁹ platelets and 1990.0 ± 12.7 IU/10⁹ platelets, respectively. The older athletes of student age did not reveal any significant differences in the activity of these enzymes (at 19 years old 9900.0 ± 271.6 IU/10⁹ platelets, 2100.0 ± 11.9 IU/10⁹ platelets, 20 years old - 9890.0 ± 231.9 IU/10⁹ platelets, 2050.0 ± 21.3 IU/10⁹ platelets, 21 years old - 10600.0 ± 236.4 IU/10⁹ platelets, 1960.0 ± 18.6 IU/10⁹ platelets, 22 years - 10150.0 ± 280.3 IU/10⁹ platelets, 2060.0 ± 12.7 IU/10⁹ platelets, respectively). The state of activity of catalase and superoxide dismutase in the blood plates in older athletes

was not significantly different from 18–22 year olds, not changing from 26–27 years (9920.0 ± 218.6 IU/ 10^9 platelets and 2000.0 ± 20.1 IU/ 10^9 platelets, respectively), up to 34-35 years (9850.0 ± 196.0 IU/ 10^9 platelets, 1920.0 ± 17.5 IU/ 10^9 platelets, respectively).

Candidates and masters of sports in track and field athletics at the age of 18, under the action of collagen, developed at 36.4 ± 0.24 s, being at a comparable level in older athletes. The high duration of AP development in 18 flight observables was noted under the influence of ADP (47.9 ± 0.12 s) and ristomycin (53.2 ± 0.20 s). Later, thrombin and adrenaline AP developed, reaching at 18 years old 59.7 ± 0.18 s and 109.7 ± 0.22 s, respectively, not significantly differing from that of the older patients. Evaluation of the simultaneous effects of several agonists in 18 year old athletes revealed that AP with ADP+adrenaline was 38.5 ± 0.13 s, with ADP + collagen 29.6 ± 0.19 s, with adrenaline + collagen 34.1 ± 0.19 s, staying at this level in all subsequent ages. Thus, in 26–27 years of the observed antibodies, under the influence of collagen, developed over 36.1 ± 0.11 s, being at a similar level in all older subjects. Similar AP activity was noted at this age in young people who stopped regular exercise under the influence of ADP (47.5 ± 0.18 s) and ristomycin (49.1 ± 0.11 s). Later, thrombin and adrenaline antibodies appeared, being at 26-27 years old, 60.3 ± 0.19 s and 105.1 ± 0.23 s, respectively, and did not significantly change in the older observed ones. At 26-27 years old, when combined inductors were used in AP athletes who left training regularly for ADP + adrenaline - 39.2 ± 0.16 s, for ADP + collagen - 30.1 ± 0.22 s, for adrenaline + collagen - 30.4 ± 0.31 s, not significantly different from the duration of AP at the age of 34-35 years.

The discoid platelet blood level of 18-year-old athletes was $88.3 \pm 0.14\%$, not significantly different from the same level in the older ages examined. The number of active forms of platelets, their total number also remained stable in their bloodstream from 18 to 22 years. In the blood of observable athletes of student age, regularly practicing levels of free-circulating small and large platelet aggregates did not experience reliable dynamics, averaging 2.2 ± 0.15 and 0.04 ± 0.016 per 100 free-lying platelets, respectively. The content of platelets involved in the process of aggregation, the observed candidates and masters of sports also did not change between 18 and 22 years old, averaging $5.0 \pm 0.12\%$. The number of discoid platelets in the blood of candidates and masters of sport in athletics who regularly trained to 22 years of age was 26-27 years old $85.1 \pm 0.12\%$, not significantly different from the values in the younger and older age groups included in the study. The content of active forms of platelets also remained unchanged in their bloodstream from 26 to 35 years, which determined the stability of their total number. In the blood of the observed athletes who had previously regularly trained in athletics, the number of free-circulating small and large platelet aggregates did not have a reliable dynamic, reaching 34-35 years old 2.5 ± 0.28 and 0.08 ± 0.039 per 100 free platelets, respectively. The amount of platelet involvement in the process of aggregation in vivo in athletes also remained stable between 26 to 35 years, amounting to 34-35 years $5.9 \pm 0.22\%$.

DISCUSSION

Physical activity is a factor of the environment affecting the human body including all internal organs and blood cells [10,11].

Candidates and masters of sports aged 18-22 years, regularly experiencing intense physical activity revealed low activity of lipid peroxidation increased antioxidant enzymes in platelets [12,13]. At the same time, candidates and masters of sports of athletics 26-35 years to 22 years who trained regularly and remained elevated antioxidant protection of blood platelets when unexpressed in them level of lipid peroxidation [14,15]. Also, regularly and irregularly practicing candidates and masters of sports of 18-35 years revealed consistently small platelet aggregation [16], probably largely associated with constantly low sensitivity of the receptor apparatus platelets [17,18], providing a low level of activity of platelets in vivo [19,20].

The determination of the duration of AP with individual inductors and their combinations in athletes of college age regularly physically training up to 22 years, and subsequently leaving regular training, revealed a low ability of platelets to aggregate at the age of 18-35 years. This AP activity was provided by unexpressed functionalities of phospholipase C [21], which controls the phosphoinositol pathway, phospholirination of actin and myosin, their reduction and the intensity of Ca^{2+} release [22,23] from the depot [24]. Stability in AP athletes with weak inducers was ensured by low expression of fibrinogen receptors (GPIIb-IIIa) [25,26] and unexpressed release of arachidonic acid by phospholipase A_2 from membrane phospholipids [27,28], which goes to synthesis of thromboxane A_2 [29]. The use of several inductors at once revealed their bright mutually

potentiating action [30], confirming the regularities established in the assessment of AP with individual agonists [31,32].

The stability of the intravascular platelet activity of candidates and masters of sport in athletics, who regularly trained at the age of 18-22 years and subsequently switched to irregular classes [33], indirectly indicated that low concentrations of aggregation inducers remained in the bloodstream against the background of a small number of receptors for them platelets [34]. At the same time, a high number of discoid-shaped platelets in the blood of 18–35 years of age is recorded in the blood due to the low activity of their receptors for aggregation inducers and fibrinogen (GP IIb - IIIa) [35-40].

It can be considered that candidates and masters of sports in athletics who regularly exercise physically from 18 to 22 years old, and subsequently who switched to irregular workouts at least until 35 years of age, have a low platelet activity in vitro and in vivo, ensuring their optimal micro-rheological blood properties.

CONCLUSION

The candidates and masters of sport in athletics who regularly train at the age of 18–22, and subsequently who train irregularly for at least 35 years, have consistently low sensitivity of platelets to aggregation inductors and their combinations. Candidates and masters of sports in track and field athletics aged 18-35, who regularly trained until the end of their students, showed low intravascular platelet activity with a low number of free-circulating blood plate aggregates in their blood.

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