Platelet function activity in black-motley calves during the dairy phase

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Abstract. Platelet activity in cattle can change under the influence of many factors. Their assessment in the second phase of their early ontogenesis is of great interest, taking into account the breed of calves. The purpose of the work is to evaluate platelet activity in black-and-white breed dairy calves. The study was conducted on 41 calf of black-motley breed, which was obtained from healthy cows as a result of 2–3 pregnancies. The calves were examined on the 11th, 15th, 20th, 25th and 30th day of ontogenesis. The study used biochemical, hematological and statistical methods. In animals, an increase in platelet aggregation with all tested inductors was detected during the milk feeding phase. The number of discoid platelets in the blood of calves observed during the second phase of early ontogenesis decreased by 10.5%. Moreover, the total number of active platelets increased by 24.0%. The levels of small, as well as medium and large aggregates of platelets present in the blood also increased during the milk feeding phase by 28.6 and 27.3%, respectively. This was achieved in the observed calves by an increase of 9.6% in the synthesis of thromboxane in platelets due to an increase in the activity of cyclooxygenase in them by 9.4% and thromboxane synthetase by 9.3%. This was also influenced by the increase in the platelet content of adenosine phosphates and the increase in their secretion. The levels of actin and myosin in inactive calf platelets increased during the milk feeding phase by 9.7 and 13.2%, respectively. This provides them with a high degree of preservation of blood volume in case of damage to blood vessels. The growth of intravascular platelet activity in these calves also contributes to the creation of the necessary conditions to minimize blood loss and ensure homeostasis.

1 Introduction

As the functionality of the hemostatic system seriously identifies hemocirculation [1, 2]. Great value in this belongs to platelets. Their hemostatic properties strongly govern the operation of microcirculation [3, 4] in all organisms [5, 6]. Previously shown that platelet activity changes during ontogeny [7] and in aging [8, 9], the development of dysfunctions [10, 11], the development of pathology [12], the formation of parasopati [13, 14] and on the background of therapeutic effects [15–17]. However, many aspects of the functioning of platelets in cattle so far been studied very poorly. There are some works devoted to the platelets of these productive animals in the individual age groups [18]. These individual details do not allow having a holistic view on the issue and dictating the need to perform extensive research. The nee for these works is due to the great importance of the functions of platelets for the flow of blood flow in the capillaries. It is clear that microcirculation affects the growth and development of animals is strong, thus realizing their productive potential [19]. Due to genetic differences between individual breeds of cattle and high physiological values of platelets activity in the realization of the potential productivity, it is necessary to evaluate the properties of thrombocytes in calves belonging to highly productive on the milk yield of black-motley breed in the second phase of their early ontogeny.

The goal is to evaluate the activity of platelets in calves’ milk of black-motley breed.

2 Materials and methods

The research was conducted in strict accordance with ethical principles established by the European Convent on protection of the vertebrata used for experimental and other scientific purposes (adopted in Strasbourg March 18, 1986, and confirmed in Strasbourg June 15, 2006) and approved by the local ethic committee of All-
Russian II of Physiology, Biochemistry and Animals' feeding (Record №11, dated December 4, 2015). The study was conducted on 41 calf black-and-white breed, which were obtained from healthy cows 2–3 of pregnancy. All the calves were examined for the phase of milk feeding 5 times: 11, 15, 20, 25 and 30 day of life.

The determination of the intensity of the formation of thromboxane in platelets and evaluation of the activity of cyclooxygenase and thromboxane synthetase was performed in three sample transfer using the photoelectrocolorimeter [20]. In platelets was assessed by the levels of adenosine triphosphate (ATP) and adenosine diphosphate (ADP), the intensity of their secretion when introduced into the environment of collagen and number of actin and myosin in the protein composition of the cytoskeleton of intact and exposed to platelet activation in response to ADP [20].

The development period of platelet aggregation (AP) was found by visual micro-method [21], using as inductor ADP (0.5x10⁻⁴ M), collagen (dilution 1:2 primary suspension), thrombin (0.125 unit/ml), epinephrine (5.0x10⁻⁶ M) and ristomycin (0.8 mg/ml) in plasma, which were standardized level of platelets 200x10⁹ platelets. Intravascular platelet activity was assessed using phase-contrast microscopy [22]. Statistical processing of received information was made with the help of a program packet "Statistics for Windows v. 6.0", "Microsoft Excel". Differences in data were considered reliable in case р<0.05.

3 Results

The number of platelet-disc cells in the blood of calves taken into the study during the phase of milk feeding decreased to 70.1±0.24 %. During the observation of them, the total number of active platelets in their blood increased by 24.0 %. The number of platelet aggregates of any size in their blood between 10 and 30 days of life increased by 28.6 and 27.3 %, respectively.

In dairy calves of black and white breed, an increase in the hemostatic properties of platelets was noted. Thus, in the examined calves, on the 11th day of life, AP occurred with collagen for 31.8±0.10 s, then accelerating by 30 days of ontogenesis to 28.6±0.15 s. A similar acceleration of antibodies was detected with ADP and ristomycin up to 36.5±0.09 s and 43.6±0.17 s, respectively. For antibodies with thrombin and adrenaline, the time was also reduced to 48.7±0.11 s and 91.1±0.21 s.

An important mechanism ensuring the acceleration of antibodies in black-motley breed milk calves is the enhancement of thromboxane synthesis in their platelets. This was indicated by an increase of 9.6 % AP in a simple transfer sample. The basis of this in these calves was an increase in the activity of platelet cyclooxygenase and thromboxane synthetase. This was indicated by an increase in the recovery of antibodies in a collagen-aspirin sample, which indirectly estimates the state of cyclooxygenase (by the end of the observation, 86.6±0.14 %). The level of AP recovery in the collagen-imidazole test, which indirectly measures the activity of thromboxane synthetase in calf platelets, also increased and amounted to 44.8±0.16 % on day 30.

Initially, a low number of ATP and ADP of calves in platelets increased, reaching 5.96±0.012 and 3.61±0.009 μmol/10⁹ platelets by 30 days of life. Under these conditions, their secretion from platelets between 10 and 30 days of life increased to 33.9±0.14 and 44.8±0.13 %.

The amount of actin and myosin on day 11 in inactive calf platelets was 27.8±0.10 and 13.6±0.13 % of the total protein in the platelet. By the end of the observation, it amounted to 30.5±0.14 and 15.4±0.16 % of the total protein in the platelet. During the observation, the calves also had an increase in platelet aggregation of actin formation by 11.1 % and myosin by 9.8 %.

4 Discussion

Recent studies have led to an understanding of the great biological significance of the dynamics of hematological parameters in humans and animals due to the fact that they reveal various aspects of the functioning of homeostasis [23, 24]. Given the great importance for the work of the body of calves of high-milk breeds, platelet activity in them is studied very poorly. This was the motivation for performing this study on calves of the black-motley breed of dairy food.

In the assessment of calves AP with collagen and ristomycin, a gradual increase in the adhesion of their platelets during the milk feeding phase was found. Obviously, this happened according to two mechanisms [25]. The first mechanism was revealed by the found acceleration of their aggregation with collagen. This should be associated with an increase in calf platelet membranes during the observation of the number of glycoproteins IIa-IIa and VI, which are collagen receptors.

The second mechanism for enhancing platelet adhesion in black-motley breed milk calves is an increase in the number of von Willebrand factor receptors (GPIb) on their surface with an increase in their blood concentration. This was indicated by the acceleration of AP found in the examined calves in response to ristomycin.

The accelerated platelet aggregation detected in calves of dairy black-motley breed undoubtedly gives a high degree of protection for their body from blood loss. Some acceleration of their antibodies to strong inducers (collagen and thrombin) is provided by a slight increase in the number of receptors for them on platelet membranes in combination with activation of phospholipase C and phosphoinositol pathway against the background of increased phosphorylation of proteins of their contractile system.

The activation of formation of inositol trisphosphate calves in platelets is apparently ensured at this age by enhancing the release of Ca²⁺ from dense granules, which is a mechanism for stimulating self-assembly and reducing actomyosin [26].
n increase in blood plates of dairy calves. cycl
the synthesis of thromboxane A₂, also stimulated
activity of cyclooxygenase and thromboxane synthetase,
synthesis of thromboxane A₂ from arachidonic acid membranes provided enhanced
an increase in the activity of phospholipase A₂, which accel-
red with age. This was undoubtedly associated with an increase in the
expression of fibrinogen receptors (GPIIb-IIIa), and an increase in the activity of phospholipase A₂ in them. The intensification of the release of phospholipids from arachidonic acid membranes provided enhanced synthesis of thromboxane A₂ [27]. The increase in the activity of cyclooxygenase and thromboxane synthetase, found in black and white breed calves, also stimulated the synthesis of thromboxane A₂. This was confirmed by transfer tests, which showed an increase in the activity of cyclooxygenase and thromboxane synthetase in the blood plates of dairy calves. Another important mechanism for accelerating AP in dairy calves is the increase in actinogenesis and myosin formation under conditions of the appearance of an aggregation inducer in the plasma and increased secretion of ATP and ADP calf platelet granules from the platelets.

To estimate the start of platelet activation in calves milk supply black-and-white was clarified by intravascular platelet activity during vasovasostomy microscopy. These calves were found the increase in the number of active platelets in the field. It talked about the increase in the sensitivity of platelets to aggregation of stimulants. The increase of intravascular platelet activity, also spoke about increasing availability to the blood clots of the blood vessels, including because of the rise in their blood svobodnozhivushchikh trombotsitarnyh units. It also pointed to the increase in

Table 1. Platelet characteristics of black-motley calves of dairy nutrition

| Parameter                                | 11 day                  | 15 day                  | 20 day                  | 25 day                  | 30 day                  |
|------------------------------------------|-------------------------|-------------------------|-------------------------|-------------------------|-------------------------|
| The number of ATP in platelets before secretion, µmol /10^9 platelets | 5.4±0.010               | 5.5±0.016               | 5.6±0.018               | 5.7±0.015               | 5.9±0.012               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The number of ADP in platelets before secretion, µmol /10^9 platelets | 3.27±0.009              | 3.37±0.011              | 3.45±0.007              | 3.52±0.012              | 3.61±0.009              |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The severity of secretion ATP, %         | 30.3±0.10               | 31.0±0.13               | 31.6±0.06               | 32.8±0.10               | 33.9±0.14               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The severity of ADP secretion, %         | 40.4±0.10               | 41.0±0.12               | 42.1±0.15               | 43.0±0.17               | 44.8±0.13               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| AP recovery level when conducting collagen-aspirin test, % | 79.2±0.08               | 80.1±0.10               | 80.8±0.08               | 82.1±0.11               | 86.6±0.14               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| AP recovery level when conducting collagen-_imidazole test, % | 41.0±0.09               | 41.8±0.14               | 42.6±0.12               | 43.1±0.09               | 44.8±0.16               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| AP level in a simple transfer sample, %  | 30.3±0.08               | 30.9±0.05               | 31.7±0.06               | 32.6±0.08               | 33.2±0.07               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The amount of actin in inactive platelets, % of the total protein in platelets | 27.8±0.10               | 28.1±0.07               | 28.8±0.05               | 29.4±0.10               | 30.5±0.14               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The amount of actin in platelets with ADP-aggregation, % of total protein in platelets | 38.6±0.15               | 39.0±0.17               | 39.8±0.13               | 41.6±0.15               | 42.9±0.13               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The amount of myosin in inactive platelets, % of the total protein in platelets | 13.6±0.13               | 14.2±0.10               | 14.8±0.09               | 15.2±0.12               | 15.4±0.16               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The amount of myosin in platelets during ADP-aggregation, % of the total protein in platelets | 28.5±0.11               | 28.9±0.10               | 29.6±0.14               | 30.0±0.12               | 31.3±0.13               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| AP development time with ADP, s          | 39.8±0.18               | 39.1±0.16               | 38.4±0.10               | 37.7±0.14               | 36.5±0.09               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The development time of AP with collagen, s | 31.8±0.10               | 31.2±0.16               | 30.6±0.12               | 29.4±0.13               | 28.6±0.15               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The development time of AP with thrombin, s | 52.7±0.12               | 51.6±0.14               | 50.5±0.09               | 49.3±0.16               | 48.7±0.11               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The development time of AP with ristomycin, s | 47.6±0.14               | 46.8±0.10               | 46.0±0.18               | 45.2±0.12               | 43.5±0.17               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The development time of AP with adrenaline, s | 97.2±0.20               | 96.2±0.17               | 95.5±0.23               | 94.0±0.19               | 91.1±0.21               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| Platelet-discoyte level, %               | 77.5±0.14               | 76.3±0.12               | 74.1±0.16               | 72.3±0.15               | 70.1±0.24               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| Sum of activated platelet forms, %       | 22.5±0.10               | 23.7±0.16               | 25.9±0.12               | 26.7±0.14               | 27.9±0.18               |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The number of small platelet aggregates, per 100 free platelets | 3.5±0.11                | 3.6±0.09                | 3.8±0.06                | 4.1±0.08                | 4.5±0.07                |
| p<0.05                                  |                         |                         |                         |                         |                         |
| The number of medium and large platelet aggregates per 100 free platelets | 0.11±0.018              | 0.11±0.015              | 0.12±0.016              | 0.13±0.023              | 0.14±0.020              |
| p<0.05                                  |                         |                         |                         |                         |                         |

Note: p – is the reliability of the dynamics of indicators in relation to 11 days of age.
the blood of the calves-dairy producers of black-motley breed of other inducers of platelet aggregation (ADP, thrombin, epinephrine) [28]. Gain from the observed calves of platelet aggregation leads to increased level of active forms and their units of any size in the blood. This is an important mechanism eliminating the risk of bleeding in the absence of blockade functionally required number of microvascular platelet aggregates and maintains the optimum level of physiological processes of platelet-vascular interactions. Found intravascular aggregation of blood platelets in calves milk of black-motley breed indicates high activity adequate adhesion and aggregation properties of blood platelets and suggests that they are physiologically sufficient level of disaggregation, apparently due to the high density of receptors antiplatelet agents on the membranes of platelets.

5 Conclusion

For calves’ milk of black-motley breed is characterized by the increased activity of platelets. It maintains homeostasis by ensuring that conditions to minimize their bleeding, while maintaining optimal conditions for microcirculation in the tissues. This ensures they gain in the physiologically acceptable limits of the activity of the mechanisms of platelet adhesion, aggregation and secretion. The increased intravascular activity of platelets in calves of black-motley breed during the phase of milk feeding significantly ensures the preservation of homeostasis under normal condition of blood flow in the vessels and optimum metabolism in their muscles and internal organs, which creates conditions for the normal growth and development of calves of this breed.

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