

1 **AGGREGATION OF BASIC REGULAR BLOOD ELEMENTS IN**
2 **CALVES DURING THE PHASE OF MILK FEEDING**

3 **ABSTRACT**

4 **Aim.** The aim was to examine aggregation activity of basic regular blood
5 elements of calves during the phase of milk feeding.

6 **Study design.** The study used 39 calves of black and white breed which were
7 taken into the research on the 11th day of life. They were examined on the 11th,
8 15th, 20th, 25th and 30th days of life.

9 **Place and duration of the study.** The study was conducted on “Kolos” farm of
10 Fatezh district in Kursk region, Russia, in spring, 2014.

11 **Methodology.** We used biochemical, hematological and statistical methods of
12 investigation. We estimated the intensity of lipids' peroxidation in plasma,
13 aggregation of erythrocytes, platelets and neutrophils.

14 **Results.** The calves were noted to have an upward trend of erythrocytes'
15 spontaneous aggregation during the phase of milk feeding.. It could be judged
16 by a light upward trend of erythrocytes' summary quantity in an aggregate,
17 quantity rise of aggregates themselves and number lowering of disaggregated
18 erythrocytes. All the calves were noted to have a trend to strengthening of
19 platelets' aggregation during the phase of milk feeding. So, on the 11th day of
20 life their period of platelets' aggregation development under collagen impact
21 was equal to $30.7 \pm 0.12s$. It decreased to some extent during the research.
22 Similar state of healthy animals' platelets' aggregation was noted for adenosine
23 diphosphate (to the end of the phase - $38.1 \pm 0.15s$) and ristomicin (to the end of
24 the phase - $46.2 \pm 0.17s$). In later period platelets' aggregation which was
25 developed with thrombin and adrenaline, also had a trend to light acceleration
26 during the research and to its end was equal to $51.3 \pm 0.18s$ and $98.0 \pm 0.34s$,
27 respectively. The calves were also noted to have a little trend to strengthening of

28 neutrophils' aggregation during the phase of milk feeding. So, their neutrophils'
29 aggregation during the research rose with lectin on 4.6%, with concanavalin A -
30 on 6.4%, with phytohemagglutinin - on 3.2%.

31 **Conclusion.** During the phase of milk feeding the calves showed low activity of
32 lipid peroxidation in plasma, which had a slight up-ward tendency. The calves
33 of the age between 11 to 30 days of life were found to have little strengthening
34 of regular blood elements' aggregation.

35 **Key words:** phase of milk feeding, calves, aggregation, erythrocytes, platelets,
36 white blood cells.

37 **1. INTRODUCTION**

38 Blood consists of regular elements and plasma. It continuously circulates along
39 vessels in a living body [1]. It provides gas metabolism and delivery of nutrients
40 and biologically active substances to tissues [2,3]. It also provides removal of
41 metabolic waste products out of them [4,5]. The efficiency of hemocirculation,
42 especially in microcirculation system, mostly depends on regular blood
43 elements' aggregation [6,7]. It's evident under constant control from the side of
44 a vascular wall [8,9]. It was noted that surplus aggregation of erythrocytes,
45 platelets and leucocytes could inhibit metabolic processes in a body [10,11]. In
46 this connection, we are sure that estimation of the degree of regular blood
47 elements' aggregation in calves at the beginning of their ontogenesis - in the
48 phase of milk feeding - is very urgent [12]. Given researches are important for
49 both fundamental science and practice as abnormalities in the processes of
50 aggregation and disaggregation in blood play essential role in pathogenesis of

51 many diseases [13,14]. Both physiology of animals and veterinary science need
52 precisely adjusted normative indices of basic regular blood elements'
53 aggregation [15]. These norms are necessary for estimation of dynamics of
54 cattle state, including milk fed calves, in case of application of various impacts
55 on their bodies [16].

56

57 The following aim was set in our research - to examine aggregation activity of
58 regular blood elements in calves during the phase of milk feeding.

59 **2. MATERIALS AND METHODS**

60 The research was conducted in strict accordance with ethical principles
61 established by the European Convent on protection of the vertebrata used for
62 experimental and other scientific purposes (adopted in Strasbourg in March,
63 18th, 1986, and confirmed in Strasbourg in June, 15th, 2006) and approved by
64 the local Ethics Committee of Kursk Institute of Social Education, a branch of
65 Russian State Social University (record №12, dated December, 3rd, 2015) and
66 the local Ethics Committee of All-Russian Scientific Research Institute of
67 Physiology, Biochemistry and Animals' Feeding (record №11, dated December,
68 4th, 2015).

69

70 The study used 39 calves of black and white breed, taken into the research on
71 the 11th day of life. All the calves were received in autumn. The animals were
72 kept in Kursk region (Central Russia) in calf-sheds of the farm “Kolos” without

73 special heating. They drank whole milk in the amount of 6-7 liters a day from
74 the teaspoon drinking bowls, which amounted to approximately 12-14% of their
75 body weight. They were examined five times during the phase of milk feeding -
76 on the 11th, 15th, 20th, 25th and 30th days of life.

77

78 The activity of the processes of lipids' peroxidation (LPO) in plasma was
79 estimated according to the content of thiobarbituric acid (TBA)-active products
80 with the help of a set "Agat-Med" and acyl hydroperoxides (AHP). Antioxidant
81 potential of liquid part of blood was determined according to its antioxidant
82 activity (AOA) [17].

83

84 The evidence of erythrocytes' aggregation was determined with the help of a
85 light microscope in Gorjaev's box. We registered the quantity of erythrocytes'
86 aggregates, the number of aggregated and disaggregated erythrocytes [18].

87

88 Platelets' aggregation (AP) was estimated with the help of visual micromethod
89 of AP estimation [19] with the usage of adenosine diphosphate (ADP) (0.5×10^{-4}
90 M), collagen (dilution 1:2 of basic suspension), thrombin (0.125 un/ml),
91 ristomicin (0.8 mg/ml) and adrenaline (5.0×10^{-6} M) in rich in platelets plasma
92 with standardized platelets' quantity 200×10^9 tr. Activity of neutrophils'
93 aggregation was estimated with the help of a photoelectrocolorimeter. We used

94 lectin of wheat foetus in a dose of 32 mkg/ml, concanavalin A - 32 mkg/ml and
95 phytohemagglutinin - 32 mkg/ml as inductors.

96

97 Statistical processing of received data was made with the help of a program
98 package "Statistics for Windows v. 6.0", "Microsoft Excel". A single-factor
99 analysis of variance was used with application of the F-reliability criterion of
100 Fisher. Differences in data were considered reliable in case of $p < 0.05$.

101

102 **3. RESULTS AND DISCUSSION**

103 Examined calves were noted to have little LPO activity of plasma with a slight
104 trend to increase during the period of the research. The content of AHP in it rose
105 from 1.44 ± 0.17 D₂₃₃/1ml to 1.47 ± 0.25 D₂₃₃/1ml, TBA-active products - from
106 3.59 ± 0.15 umol/l to 3.64 ± 0.28 umol/l. It was accompanied by a trend to some
107 weakening of plasma AOA from $33.5 \pm 0.38\%$ on the 11th day of life to
108 $33.0 \pm 0.34\%$ on the 30th day of life (table 1).

109

110 During the phase of milk feeding the calves were noted to have unexpressed
111 upward trend of spontaneous erythrocytes' aggregation. It could be judged by a
112 slight upward trend of summary erythrocytes' quantity in an aggregate (on
113 1.9%), quantity rise of aggregates themselves (on 2.4%) and number lowering
114 of disaggregated erythrocytes (on 2.2%) (table 1).

115

116 All the milk fed calves were noted to have a trend to strengthening of platelets'
117 aggregation. So, on the 11th day of life their period of AP development under
118 the impact of collagen was equal to 30.7 ± 0.12 s. It decreased to some extent
119 during the research. Similar AP state of healthy animals was noted for ADP (to
120 the end of the phase - 38.1 ± 0.15 s) and ristomicin (to the end of the phase -
121 46.2 ± 0.17 s). In later period AP which was developed with thrombin and
122 adrenaline, also had a trend to slight acceleration during the research and to its
123 end was equal to 51.3 ± 0.18 s and 98.0 ± 0.34 s, respectively (table 1).

124 During the phase of milk feeding the calves were also noted to have a little
125 trend to strengthening of neutrophils' aggregation. So, during the research their
126 neutrophils' aggregation rose with lectin on 4.6%, with concanavalinA - on
127 6.4%, with phytohemagglutinin - on 3.2% (table 1).

128 The consumption of milk and beef by the population of the planet increases. It
129 dictates the necessity of constant development of this agricultural branch. It can
130 be achieved in the result of continuation of active scientific researches in the
131 field of cattle physiology [15,20]. In this connection, special significance is
132 given to researches of calves' blood physiology at the beginning of ontogenesis
133 [21,22]. Much attention is paid to studying of calves which prepare to switch to
134 the consumption of vegetable feeding. In our work it was found that calves at
135 the age between 11 to 30 days of life had stable plasma AOA. It was
136 accompanied by a stable level of LPO products in plasma. Found facts were
137 supported by the results of earlier researches [23]. It is known that intensity of

138 freely-radical processes in plasma influences significantly the morpho-
139 functional state of erythrocytes, platelets and leucocytes [24,25]. It can explain
140 the slight ability of milk fed calves in aggregation of basic regular blood
141 elements.

142

143 In our work special attention was paid to aggregation of basic regular blood
144 elements. Intra vascular formation of units and success of microcirculation in
145 many respects depend on its level. In this regard, processes of metabolism and
146 intensity of animals' growth depend on the activity of regular blood elements'
147 aggregation.

148

149 It is obvious that a large number of electronegative proteins on erythrocytes'
150 surface [26,27] largely provides low activity of erythrocyte aggregation in
151 calves during the phase of milk feeding. High control over generation of oxygen
152 active forms in calves provides minimization of oxidative damages of
153 membrane erythrocyte proteins and globular plasma proteins which participate
154 in aggregation [28,29]. In this connection, we can come to the conclusion that
155 the phase of milk feeding of calves is characterized by optimum of metabolic
156 and receptor processes in erythrocytes. Received estimation results of
157 erythrocytes' aggregation are confirmed by the single work. It contains
158 information about the trend to its strengthening in calves of the given age [30].
159 We should compare received results with literature data with great caution. In

160 previous researches the groups were mixed, as far as breed was concerned, but
161 calves of Simmental breed prevailed. Besides, they were received in autumn. It
162 also makes comparison of results difficult.

163

164 Noted in calves trend to strengthening of platelets' aggregative activity during
165 the phase of milk feeding was connected with activity increase of their receptors
166 and postreceptor mechanisms of aggregation [31]. Concentration of von
167 Willebrand Factor - cofactor of platelets' adhesion - gradually rose in calves'
168 blood at the age of 11-30 days. It was accompanied by little number increase of
169 receptors to it - (GPIb) on platelets' surface. It was pointed by a downward trend
170 of AP period in calves in response to ristomicin. Found AP dynamics in
171 response to strong and weak agonists of aggregation could be explained by
172 physiologically approved activity changes of platelet phospholipase A₂ and C.
173 They provided functioning of thromboxane and phosphoinositol ways of
174 platelets' activation [32,33]. In literature there is rather poor information about
175 platelets' activity in milk fed calves [34]. Famous sources confirm that calves
176 have a trend to strengthening of platelets' aggregation during the phase of milk
177 feeding. But comparison of these results with received ones should be done with
178 great caution. It's connected with the fact that experimental calves in previous
179 researches were kept in Central Russia in calf-sheds with special heating, and
180 they received substitutes of whole milk and fodder concentrated products.

181

182 It is known that activity of neutrophils' aggregation in mammals is provided by
183 locuses' quantity in their glycoprotein receptors' composition. These receptors
184 can connect lectins [35]. It is firmly established that phytohemagglutinin can
185 mostly interact with parts of bD-galactose of glycoproteins, lectin of wheat
186 foetus - with N-acetyl-D-glycosamin и N-acetyl-neuraminic (sialic) acid, and
187 concanavalin A – with N-glycans containing mannose [11]. That's why, the
188 state of lectin stimulated neutrophils' aggregation of calves is determined by the
189 expression level of receptors' adhesion. These receptors have such parts in their
190 composition. Taking it into consideration, we can come to the conclusion that
191 found growth trend of neutrophils' aggregation at calves' age of 11-30 days
192 was, evidently, connected with the rise of sensitivity and density of leucocytes'
193 glycoprotein receptors. It happened simultaneously with changing of their
194 composition. Gradual strengthening of lectin - and concanavalin A - induced
195 neutrophils' aggregation in experimental calves was provided by expression
196 increase of adhesion receptors on their surface and by some growth of areas
197 containing N-acetyl-D-glucosamine, N-acetyl-neuraminic acid and mannose.
198 Strengthening increase of aggregation, induced by phytohemagglutinin in calves
199 between 11to 30 days of life, was provided by an upward trend of areas of
200 glycoproteins, containing bD-galactose [11], in their neutrophils' receptors.
201 Neutrophils' aggregation was not studied earlier in productive animals and,
202 moreover, in calves. With the help of available literature sources, containing
203 information about researches aimed at human beings, it becomes clear that the

204 role of receptor mechanisms in its realization is great, and that it can be quickly
205 damaged in case of unfavorable environmental and metabolic conditions
206 [11,32].

207

208 Noted strengthening of aggregative activity of erythrocytes, platelets and
209 neutrophils in calves during the phase of milk feeding was mostly caused by
210 processes of growth and strengthening of environmental impacts against their
211 background [36]. Sufficient activity of adaptive mechanisms keeps the balance
212 of aggregation and disaggregation in calves' blood in these conditions on the
213 level which is necessary for optimum of internals' blood supply [37].

214

215 **4. CONCLUSION**

216 The phase of milk feeding is an important stage in the development of
217 hematological indicators in cattle. During the phase of milk feeding, the calves
218 showed stability of lipids' peroxidation in plasma. It was found that calves at
219 the age of 11-30 days had a weak upward trend in aggregation of the basic
220 blood elements. This situation is, in many respects, the basis for the optimal
221 bloodstream along small vessels in milk fed calves and the processes of their
222 growth.

223 **REFERENCES**

- 224 1. Medvedev IN, Zavalishina SYu. Platelet Activity in Patients With Third
225 Degree Arterial Hypertension and Metabolic Syndrome. *Kardiologiia*.
226 2016;56(1):48.
- 227 2. Medvedev IN, Gromnatskii NI, Golikov BM, Al'- Zuraiki EM, Li VI.
228 Effects of lisinopril on platelet aggregation in patients with arterial
229 hypertension with metabolic syndrome). *Kardiologiia*. 2004;44(10):57-59.
- 230 3. Medvedev IN, Gromnatskii NI, Mokhamed A.-ZE. Comparative
231 Assessment of Effects of Qadropiril and Enalapril on Intravascular Activity
232 of Platelets in Hypertensive Patients with Metabolic Syndrome.
233 *Kardiologiia*. 2004;44(12):44-46.
- 234 4. Medvedev IN, Gromnatskii NI, Volobuev IV, Osipova VM, Dement'ev VI,
235 Storozhenko MV. Thrombocytic hemostasis in hypertensive patients with
236 metabolic syndrome and its correction with lovastatin. *Klinicheskaiia*
237 *meditsina*. 2004;82(10):37-41.
- 238 5. Simonenko VB, Medvedev IN, Tolmachev VV. Comparative evaluation of
239 the influence of sulfhydryl and phosphate ACE inhibitors on thrombocyte
240 aggregation in patients suffering from arterial hypertension with metabolic
241 syndrome. *Clinical Medicine*. 2007;85(4):24-27.
- 242 6. Medvedev IN. A comparative analysis of normodipin and spirapril effects
243 on intravascular activity of platelets in patients with metabolic syndrome.
244 *Terapevticheskii Arkhiv*. 2007;79(10):25-27.

- 245 7. Kutafina NV, Medvedev IN. Platelet Aggregation in Clinically Healthy
246 Persons of the Second Coming-of-Age Living in Kursk region. *Advances in*
247 *Gerontology*.2015;5(4):267-270.
- 248 8. Medvedev IN, Skoryatina IA. Aggregation properties of blood cells and
249 vascular control over them in patients with arterial hypertension and
250 dyslipidemia. *Russian Journal of Cardiology*. 2015;4(120):18-22.
- 251 9. Simonenko VB, Medvedev IN, Mezentseva NI, Tolmachev VV. The
252 antiaggregation activity of the vascular wall in patients suffering from
253 arterial hypertension with metabolic syndrome. *Klinicheskaja meditsina*.
254 2007;85(7):28-30.
- 255 10. Medvedev IN, Gamolina OV. Lisinopril effects on platelet activity in
256 patients with arterial hypertension and impaired glucose tolerance. *Russian*
257 *Journal of Cardiology*. 2008;3:45-48.
- 258 11. Medvedev IN, Skoryatina IA. The aggregation capacity of neutrophils in
259 patients with arterial hypertension and dyslipidemia treated with fluvastatin.
260 *Klinicheskaja meditsina*. 2015;93(1):66-70.
- 261 12. Kutafina NV. Platelet Parameters of Holstein Newborn Calves. *Annual*
262 *Research & Review in Biology*. 2017; 15(2): 1-8. doi:
263 10.9734/ARRB/2017/35214
- 264 13. Medvedev IN, Skoriatina IA. Dynamics of microrheologic properties of
265 erythrocytes in patients with arterial hypertension and dyslipidemia treated
266 with atorvastatin. *Klinicheskaja meditsina*. 2012;90(6):42-45.

- 267 14. Medvedev IN, Lapshina EV, Zavalishina SYu. Experimental methods for
268 clinical practice: Activity of platelet hemostasis in children with spinal
269 deformities. Bulletin of Experimental Biology and Medicine.
270 2010;149(5):645-646.
- 271 15. Medvedev IN. Vascular-platelet interaction in pregnant cows. Bulg. J.
272 Agric. Sci.2017;23(2):310-314.
- 273 16. Zaitsev SY, Maksimov VI, Bardyukova TV. Supramolecular enzymatic
274 systems of the dog blood: Clinical diagnostic implications. Moscow
275 University Chemistry Bulletin. 2008;63(2):99-102.
- 276 17. Zaytsev GS, Bikbulatova AA, Egorova NA, Mozdykov AV, Kalashkova
277 DO. Liminal aspects of dreams. Man In India. 2016; 96(12) : 5719-5734.
- 278 18. Medvedev IN, Maksimov VI, Parakhnevich AV, Zavalishina SYu, Kutafina
279 NV. Rapid assessment of aggregation abilities and surface properties of
280 platelets and red blood cells. International Journal of Pharma and Bio
281 Sciences. 2016 April;7(2):(B)793-797.
- 282 19. Medvedev IN, Savchenko AP, Zavalishina SYu, Krasnova EG, Kumova
283 TA, Gamolina OV, Skoryatina IA, Fadeeva TS. Methodology of blood
284 rheology assessment in various clinical situations. Russian Journal of
285 Cardiology. 2009;5:42-45.
- 286 20. Kutafina NV, Medvedev IN. Platelet aggregation in clinically healthy
287 persons of the second coming of age living in Kursk region. Advances in
288 gerontology. 2015;28(2):321-325.

- 289 21. Medvedev IN. Microrheology of erythrocytes in arterial hypertension and
290 dyslipidemia with a complex hypolipidemic treatment. Russian Journal of
291 Cardiology. 2017;4(144):13-17.
- 292 22. Gromnatskii NI, Medvedev IN. Non-pharmacological correction of
293 impaired platelet hemostasis in hypertensive patients with metabolic
294 syndrome. Klinicheskaiia meditsina. 2003;81(4):31-34.
- 295 23. Zavalishina SYu. Physiological Features of Hemostasis in Newborn Calves
296 Receiving Ferroglukin, Fosprenil and Hamavit, for Iron Deficiency. Annual
297 Research & Review in Biology. 2017;14(2):1-8. doi:
298 10.9734/ARRB/2017/33617
- 299 24. Medvedev IN. Dynamics of violations of intravascular platelet activity in
300 rats during the formation of metabolic syndrome using fructose models.
301 Problems of nutrition. 2016;85(1):42-46.
- 302 25. Simonenko VB, Medvedev IN, Tolmachev VV. Effect of irbesartan of the
303 function of hemocoagulative component of hemostasis in patients with
304 arterial hypertension during metabolic syndrome. Klinicheskaiia meditsina.
305 2010;88(6):27-30.
- 306 26. Medvedev IN, Skoryatina IA. Erythrocyte aggregation in patients with
307 arterial hypertension and dyslipidemia treated with pravastatin.
308 Klinicheskaiia meditsina. 2014;92(11):34-38.
- 309 27. Medvedev IN, Skoriatina IA. Effect of lovastatin on adhesive and
310 aggregation function of platelets in patients with arterial hypertension and
311 dyslipidemia. Klinicheskaiia meditsina. 2010;88(2):38-40.

- 312 28. Simonenko VB, Medvedev IN, Kumova TA. Pathogenetic aspects of
313 hypertension in case of metabolic syndrome. *Voenno-meditsinskii zhurnal*.
314 2010;331(9):41-44.
- 315 29. Medvedev IN, Plotnikov AV, Kumova TA. Rapid normalization of platelet
316 hemostasis in patients with arterial hypertension and metabolic syndrome.
317 *Russian Journal of Cardiology*. 2008;2:43-46.
- 318 30. Glagoleva TI, Zavalishina SYu. Aggregative Activity of Basic Regular
319 Blood Elements and Vascular Disaggregating Control over It in Calves of
320 Milk-vegetable Nutrition. *Annual Research & Review in Biology*. 2017;
321 12(6): 1-7. doi: 10.9734/ARRB/2017/33767
- 322 31. Medvedev IN, Gromnatskii NI. Correction of thrombocyte hemostasis and
323 biological age reduction in metabolic syndrome. *Klinicheskaiia meditsina*.
324 2005;83(8):54-57.
- 325 32. Medvedev IN, Skoryatina IA. Fluvastatin effects on blood cell aggregation
326 in patients with arterial hypertension and dyslipidemia. *Cardiovascular*
327 *Therapy and Prevention*. 2013;12(2):18-24.
- 328 33. Medvedev IN, Kumova TA, Gamolina OV. Renin-angiotensis system role
329 in arterial hypertension development. *Russian Journal of Cardiology*.
330 2009;4:82-84.
- 331 34. Zavalishina SYu, Kutafina NV, Vatnikov YuA, Makurina ON, Kulikov EV.
332 Platelet-Activity Dependence on the Age of Rats with Experimental

333 Dyslipidemia. Biol Med (Aligarh). 2016; 8: 326. doi: 10.4172/0974-
 334 8369.1000326.

335 35. Medvedev IN, Skoryatina IA. Pravastatin in correction of vessel wall
 336 antiplatelet control over the blood cells in patients with arterial hypertension
 337 and dyslipidemia. Cardiovascular therapy and prevention. 2014;13(6):18-22.

338 36. Simonenko VB, Medvedev IN, Briukhovetskii AG. Effect of therapy with
 339 diuretics on the functional activity of platelets in patients with arterial
 340 hypertension and abdominal obesity. Klinicheskaia meditsina. 2012;
 341 90(11):54-56.

342 37. Simonenko VB, Medvedev IN, Gamolina OV. Primary hemostasis activity
 343 in patients with arterial hypertension and impaired glucose tolerance treated
 344 with trandolapril. Klinicheskaia meditsina. 2011; 89(2):29-31.

345
 346
 347
 348

Table 1. The activity of the processes of lipids' peroxidation in plasma and aggregation of blood elements in milk fed calves

Registered parameters	Age of calves (n=39, M±m)				
	11 days	15 days	20 days	25 days	30 days
acyl hydroperoxides, D ₂₃₃ /1ml	1.44±0.17	1.46±0.12 F= 0.357 (p≤0.425)	1.47±0.20 F= 1.102 (p≤0.282)	1.47±0.15 F= 1.124 (p≤0.271)	1.49±0.25 F= 1.348 (p≤0.249)
TBA-active products, umol/l	3.59±0.15	3.63±0.22 F= 0.218 (p≤0.615)	3.60±0.26 F= 0.416 (p≤0.431)	3.62±0.19 F= 1.320 (p≤0.232)	3.64±0.28 F= 2.264 (p≤0.096)
AOA, %	33.5±0.38	33.3±0.36 F= 1.220 (p≤0.252)	33.1±0.34 F= 1.758 (p≤0.189)	32.9±0.29 F= 1.974 (p≤0.192)	32.4±0.32 F= 2.126 (p≤0.174)
sum of all the erythrocytes in an aggregate	40.1±0.19	40.2±0.24 F= 0.123 (p≤0.726)	40.4±0.29 F= 1.117 (p≤0.294)	40.6±0.25 F= 1.112 (p≤0.295)	40.9±0.32 F= 1.344 (p≤0.250)
quantity of aggregates	8.2±0.12	8.2±0.10 F= 0.017	8.3±0.16 F= 0.019	8.4±0.19 F= 1.286	8.4±0.11 F= 2.912

		(p≤0.896)	(p≤0.890)	(p≤0.260)	(p≤0.092)
quantity of free erythrocytes	245.7±2.19	244.2±2.25 F= 3.122 (p≤0.081)	241.8±2.01 F= 2.284 (p≤0.135)	242.0±1.90 F= 1.529 (p≤0.220)	240.4±2.46 F= 1.032 (p≤0.313)
AP with ADP, s	39.2±0.16	39.0±0.12 F= 0.645 (p≤0.424)	38.7±0.13 F= 1.779 (p≤0.186)	38.4±0.10 F= 3.110 (p≤0.081)	38.1±0.15 F= 3.189 (p≤0.078)
AP with collagen, s	30.7±0.12	30.5±0.10 F= 0.025 (p≤0.876)	30.3±0.09 F= 0.295 (p≤0.588)	30.1±0.11 F= 0.724 (p≤0.397)	29.7±0.14 F= 1.704 (p≤0.196)
AP with thrombin, s	52.7±0.15	52.6±0.10 F= 0.238 (p≤0.627)	52.2±0.16 F= 1.207 (p≤0.275)	51.7±0.10 F= 2.505 (p≤0.117)	51.3±0.18 F= 3.039 (p≤0.085)
AP with ristomicin, s	47.5±0.12	47.2±0.16 F= 0.771 (p≤0.383)	46.9±0.22 F=0.877 (p≤0.352)	46.6±0.26 F= 2.505 (p≤0.117)	46.2±0.17 F= 3.057 (p≤0.084)
AP with epinephrine, s	97.8±0.42	97.4±0.36 F= 0.504 (p≤0.479)	97.1±0.32 F= 0.798 (p≤0.374)	98.5±0.45 F= 1.008 (p≤0.318)	98.0±0.34 F= 1.167 (p≤0.283)
Aggregation of neutrophils with lectin, %	14.5±0.16	14.5±0.17 F= 0.716 (p≤0.399)	14.7±0.15 F= 1.010 (p≤0.318)	14.9±0.26 F= 1.467 (p≤0.229)	15.2±0.22 F= 1.781 (p≤0.186)
Aggregation of neutrophils with concanavalin A, %	14.5±0.10	14.6±0.12 F= 0.529 (p≤0.469)	14.9±0.16 F=1.037 (p≤0.312)	15.1±0.11 F= 1.349 (p≤0.249)	15.5±0.13 F= 1.982 (p≤0.163)
Aggregation of neutrophils with phytohemagglutinin, %	27.1±0.19	27.2±0.23 F= 0.693 (p≤0.408)	27.4±0.14 F=0.877 (p≤0.352)	27.8±0.26 F= 1.104 (p≤0.297)	28.0±0.21 F=2.683 (p≤0.106)

349

350 Note:

351 F – the value of Fisher test when the indicators are compared with their values

352 at the age of 11 days throughout the entire observation,

353 p – possibility of unmistakable prognosis.