

# About 4-day Rhythm of Proliferative Activity of Fibroblast-like Cell Cultures Depends on the Environmental Factor

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## Research Article

**Keywords:** cell culture, fibroblast-like cells, L929, freezing, ifradian rhythm, proliferative activity

**Posted Date:** November 19th, 2021

**DOI:** <https://doi.org/10.21203/rs.3.rs-1046150/v1>

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# Abstract

A study of the 4-day rhythm of the proliferative activity of the embryonic fibroblast-like cells in the logarithmic growth phase was carried out. It was shown that in cell cultures obtained on different days from embryos of different ages, the phase of the 4-day rhythm coincides. *In vitro* the maxima of the proliferative activity were consistent with the minima of the motor activity of mice. Freezing the culture for 2 or 6 days does not cause a shift in the phase of the 4-day rhythm of cell proliferative activity compare with the unfreezing culture. That indicates the existence of an external synchronizer, which determines the 4-day infradian rhythm of the proliferative activity of embryonic cells.

Then we daily thawed samples of single L-929 culture of mice fibroblast-like cells for 22 and 17 days and researched the dynamics of its proliferative activity. We also showed 4-day rhythm of the simultaneous increase in the number of cells for all thawed samples. Taking into account that deep freezing of a culture leads to the cessation of all life processes, the fact we obtained indicates an exogenous mechanism of the formation of about a 4-day rhythm of the proliferative activity of cell culture.

# Introduction

Infradian about 4-day fluctuations were found in many biological parameters: concentrations of melatonin, glucocorticoid and sex hormones [1-6], motor activity [7], immune system [8, 9], mitotic activity of cells [10, 11]. Based on the facts that 1) 4-day rhythms are preserved during the removal both of the testes and adrenal glands [12], and 2) hibernating hedgehogs with reduced hormonal activity had the stable rhythms of spontaneous short-term arousals [13], it can be argued that the formation of this infradian rhythm is not associated with hormonal regulation. It is interesting that the removal of the pineal gland, which is involved in the formation of circadian rhythms [14], also does not affect the period and phase of the 4-day rhythm of mitotic activity of the rat esophagus epithelium [7]. Probably, the 4-day rhythm is determined by the activation of the sympathetic nervous system, which is indicated by the presence of this rhythm in the dynamics of epinephrine excretion in the urine [2], as well as the whole complex of signs associated with an increase in motor activity, the release of glucocorticoid hormones, suppression of immune reactions, a decrease in proliferative cell activity.

The mitotic activity of epithelial cells is the convenient model of the stable manifestation of 4-day rhythm [11]. However, the formation mechanisms of this rhythm have not been established yet. *In vivo* studies do not allow to exclude the influence on the mitotic activity of cells of humoral factors of the body internal environment, that have a 4-day rhythm determined by the activation of the sympathetic nervous system. Based on this hypothesis, the 4-day rhythm of mitotic activity in an isolated cell culture should be absent. However, studying the daily dynamics of the proliferative activity of the culture of embryonic fibroblast-like cells in the logarithmic growth phase, we revealed a 4-day rhythm that persists for at least three weeks of cultivation [15].

There are three possible variants of this rhythm's origin: 1) completely endogenous, self-sustaining on the basis of molecular genetic interactions, and, therefore, such a rhythm will not be synchronous in different cell cultures; 2) completely exogenous, caused by the influence of an external environmental factor; 3) endogenous, but having an external synchronizer, and, therefore, these two rhythms will be synchronous.

To determine the possible origin of the 4-day rhythm of the proliferative activity of fibroblast-like cells, it is necessary to establish:

- 1) are the manifestations of the 4-day rhythm of proliferative activity in cultures obtained on different days from embryos of the same age synchronous?
- 2) does the phase of the rhythm in cell culture comparable with that *in vivo*? Under *in vivo* conditions, the phase of the infradian rhythm of the motor activity of animals is observed in antiphase to the 4-day rhythm of mitotic activity (mitotic index) of the esophageal epithelium [7].
- 3) will freezing of cells for a period not multiple of 4 days cause a phase shift of the investigated infradian rhythm relative to the intact culture?

However, immediately after thawing, embryonic fibroblast-like cells do not begin to divide intensively. It is a considerable time period between the isolation of the culture and the beginning of the experiment, during which the cultures can synchronize their endogenous rhythms. Therefore, at the final stage of our research, we studied the changes in the proliferative activity of the fibroblast-like cells mice culture L-929, which has a high proliferative potential.

The aim of the 4th point of the study was: to establish the exogenous/endogenous nature of about 4 days of the rhythm of proliferative activity of fibroblast-like L-929 cells by daily thawing of same frozen samples of the same culture. If, after defrosting, the rhythm is immediately observed in the phase, as if it were not stopped, then we can suggest its exogenous origin, but if the restoration of the rhythm phase is observed gradually, then this is an endogenous rhythm that has an external synchronizer.

## Results

### Infradian rhythm of increase of embryonic cells' number

The highest rate of the number of embryonic fibroblast-like cells increase was observed on the second and third days after sowing, when the cultures were in the phase of logarithmic growth. Following gradual decrease in growth rates was recorded due to contact inhibition of cells (Fig. 1). As an indicator of proliferative activity, the daily increase in the number of cells on the second and third days of cultivation was calculated.

In order to establish the synchronicity of the 4-day rhythm, the proliferative activity of fibroblast-like cells obtained from embryos conceived on different dates - November 4 and 6, 2019, and November 17 and 19,

2019 was investigated. The difference in the date of conception of the studied embryos was 2 days, which corresponds to half period of the 4-day rhythm.

Figure 2A shows daily cell growth rates in two cultures, one of which was isolated on November 16 from embryos conceived on November 4, and the other – on November 18 from embryos conceived on November 6. The correlation coefficient between the indicators of these cultures was  $r = 0.67$  ( $p = 0.008$ ), therefore, their dynamics can be considered synchronous. The maximum indicators of the increase in the cells number in both cultures were noted on November 24, 28, December 2-3, 5-6 and 9-10. The minimum indicators of the increase in the cells number were registered on November 26 and 30, December 4, 8 and 12. So that we observed about 4-day rhythm.

In the first 12 days, the growth of the culture was high. We found out a statistical difference of the daily cells number increase between the acrophase (November 24, 28 and December 2) and bathyphase (November 26, 30 and December 4) of the 4-day rhythm. At the acrophase, the cells number increased 140 (120; 220) thousand cells and at the bathyphase, the increase was only up on 60 (40; 120) thousand cells ( $p = 0.006$ ). The existence of a 4-day rhythm was confirmed by autocorrelation analysis. As the cell growth rate gradually decreased, its deviations from the approximating trend line were calculated. The autocorrelation coefficients are presented in the Table 1. The correlation coefficient between the original series and the series shifted by 2 days is  $r = -0.36$  ( $p = 0.11$ ), and by 4 days -  $r = 0.49$  ( $p = 0.04$ ).

Table 1  
The calculated autocorrelation coefficients for the dynamics of the mean of proliferative activity of the culture cells #1 and #2, received on November, 16 and 18, 2019.

lag, day	r	p
1	-0.07	0.73
2	-0.36	0.11
3	0.002	0.99
4	0.49	0.04

Figure 2B shows the daily growth of two other cells of cultures. The first culture was isolated on November 29 from embryos who had been conceived on November 17, 2019. The second one was isolated on December 1 from embryos who had been conceived on November 19. We also revealed the synchronicity in the dynamics of these cultures and coefficient of correlation ( $r$ ) was 0.68 ( $p = 0.007$ ). The maximum indicators were registered on 2-3, 5-6 and 9-10, 14 and 18 December, and the minimum - on 4, 8, 12, 16 and 19-20 December. The indices of daily growth in the acrophase of the 4-day rhythm were 160 (130; 280) thousand cells, and in the bathyphase were 100 (70; 135) thousand cells. The daily growths in

the acrophases and the bathyphases statistically significantly differed among themselves ( $p = 0.01$ ). The autocorrelation coefficient between the original series and the series shifted by 2 days is statistically significant and the coefficient correlation ( $r$ ) is  $-0.69$  ( $p = 0.002$ ), which also confirms the stability of the 4-day rhythm in the dynamics of proliferative activity of culture cells.

In all four studied cultures, the maximum proliferative activity got in the same calendar dates; therefore, the phase of the 4-day rhythm of these cultures coincided.

### *Matching of infradian rhythms between proliferative activity of embryonic cell culture and motor activity of mice*

We investigated the daily motor activity of male mice in the period from November 27 to December 21, 2019 to identify the synchronicity of the 4-day rhythm of proliferative activity *in vitro* and the motor activity of mice *in vivo* (Fig. 3). The maxima of the motor activity of mice were at the minima of the proliferative activity of the culture cells. A negative correlation coefficient was revealed between these parameters:  $r = -0.47$  ( $p = 0.016$ ).

## **Freezing and thawing of embryonic cell cultures**

To test the hypothesis about the endogeneity of the 4-day rhythm, we froze the cells for a period not multiple of 4 days. In the case of the endogenous nature of this rhythm, such a procedure should have led to a shift the phase of the proliferative activity of the frozen cell culture in relation to the intact one. Figure 4 shows the daily increase in the cells number of one culture, a part of that were frozen for 2 and 6 days, and the other part were cultured as usual. We detected a 4-day rhythm of the dynamics of proliferative activity of all three cultures. The maximum indicators were noted on January 2-3, 6-7 and 10-11, and the minimum on January 4-5, 8-9 and 12-13. The correlation coefficient between frozen and intact cultures was  $r = 0.78$  ( $p < 0.001$ ) and  $r = 0.72$  ( $p = 0.008$ ). Consequently, freezing the culture for 2 and 6 days did not lead to a change in the phase of the 4-day rhythm of proliferative activity relative to the intact one.

## **Freezing and thawing of L929 cells**

The next task was to study the dynamics of proliferative activity of single L-929 cell culture samples, which were thawed daily and immediately plated into Petri dishes (Table 4, first series). Figure 5 shows the dynamics of the daily increase in the cell number of the studied cultures after the 2nd, 3rd, 4th and 5th days of cultivation. It can be noted that synchronous changes are observed in the dynamics of proliferative activity. Indicators of the daily increase in the number of cells have maximums on January 18-19, 22-23, 27, 30-31 and February 4. The minimum increase in the cell number was noted on January 16, 21-21, 24, 28-29, February 1 and 5, as well as an additional minimum on February, 3. The correlation coefficient of the increase in the cell number is maximum between 2 and 3 days ( $r = 0.54$   $p = 0.024$ ).

To approve the discovered rhythm of L929 cell proliferative activity for the period from January 16 to February 1, the autocorrelation coefficient of the average index of proliferative activity for the 2nd - 5th

days of cultivation was calculated (Fig. 6). Autocorrelation analysis confirms the statistical significance of the 4-day rhythm.

Thus, in the dynamics of the L-929 cell proliferative activity over a long period from January 16 to February 1, a 4-day rhythm was revealed. During the acrophase the indicator of daily growth after the 2nd and 3rd days of cultivation (January, 18-19, 22-23, 27, 30-31) was 1.97 (1.69–2.12) and during the bathiphase (January, 16, 20-21, 24, 28-29) was equal to 1.40 (1.30-1.61). There was a statistically significant difference between the values in acrophase and bathiphase ( $p = 0.0015$ ).

Based on the fact that the maximum correlation coefficient is observed between the 3 and 4 days of an increase in the L929 cells number, it can be assumed that it is during this period that the culture is most sensitive to external influences that affect the proliferative activity of cells. The indicators of 2 days after the start of cultivation have great instability, probably caused by the procedure of thawing and sowing the cells. The indicator of proliferative activity after 5 days of cultivation has a small value, which is probably affected by the exhaustion of the nutrient medium and the high density of the monolayer formed by this time. Therefore, in the next series of experiments, we analyzed the proliferative activity after 3 days of cultivation.

In the second series of experiments on daily thawing of cryoprobes in the period February 15 - March 3, 2021 (Table 5), a similar dynamic was revealed. The maximum daily increases in the cell number were observed on February 20, 23, 27 and March 3 (Fig. 7). The minimum values of proliferative activity were established on February 21, 24-26, March 1 and 5. It should be noted that a statistically significant correlation is observed between experimental samples, that were daily thawing and control samples that were parallel cultivated and not subjected to freezing ( $r = 0.65$ ,  $p = 0.004$ ).

## Discussion

In cell cultures obtained from embryos of the same age, but conceived on different calendar dates, the phase of the 4-day rhythm of proliferative activity coincides. Therefore, there is an external effect that synchronizes the proliferative activity of fibroblast-like cells *in vitro*. A similar pattern was previously established by us for *in vivo* conditions: the 4-day rhythm of glucocorticoid hormones and the mitotic activity of epithelial cells of the esophagus coincides in phase not only between individuals of the same species, but also between species belonging to the class of mammals and birds [16]. Consequently, the phase of the 4-day rhythm of the proliferative activity of the primary cell culture *in vitro* continues to be synchronized with the analogous rhythm of mitotic activity *in vivo*.

Freezing the culture for 2 or 6 days (a period not multiple of 4 days) does not cause a shift in the phase of the 4-day rhythm of the proliferative activity of embryonic fibroblast-like cells relative to the intact culture. Thus, the studied rhythm is, most likely, either a consequence of the direct action of an external factor on the cells, i.e. exogenous, or endogenous, but having an external synchronizer as after thawing the culture, before the start of the experiment, it was necessary to carry out one passage for 5 days,

perhaps, the cells managed to synchronize their endogenous rhythms with an external factor during this short period.

A long-term study of the L-929 cell culture proliferative activity during to simultaneously freezing of single samples and then thawing daily for 20 days, showed existence of a 4-day rhythm. We suggest that as after deep freezing all vital processes stop in the cells, the presence of a 4-day rhythm in the daily increase in L-929 cell number is caused an exogenous mechanism of the formation of this rhythm. The hypothesis that the studied rhythm is endogenous, but it is synchronized by an external factor, isn't consistent, as we observed the rhythm synchronicity already two days after defrosting. If the synchronization by a not constantly acting external environmental factor was being, synchronous rhythmic changes in proliferative activity would be observed later after defrosting.

Thus, our results point out an exogenous mechanism of a 4-day rhythm formation of the daily increase in cells number both in embryonic and L-929 cultures. The direction for further research is the search for an external environmental factor that determines the rhythmic changes in the proliferative activity of cell culture.

Oscillations of the geomagnetic field can be a possible external synchronizer. It was found that the screening of the Earth's magnetic field causes a decrease in the proliferative activity of the Fibrosarcoma HT1080 tumor cells and colorectal HCT116 cancer cells [17]. In [18] it was shown that 24-hour exposure of the endothelial cell culture in a weak constant magnetic field leads to an increase in their proliferation.

Currently, there are reasons to consider natural electromagnetic fields as a possible synchronizer of infradian and ultradian rhythms [19, 20]. In addition to the established mechanism for the perception of these oscillations at the whole organism [21-23], there are few data that the reception of electromagnetic oscillations is possible within the one cell. It was shown that irradiation of a suspension of T- and B-lymphocytes with low-intensity electromagnetic radiation (42.2 GHz, 1  $\mu\text{W}/\text{cm}^2$  amplitude modulation 10 Hz) suppressed their blast transformation [24]. Effecting of periodic weak outside pulsed magnetic fields to the activated neutrophils, attached to a glass substrate, increase by them the production of reactive oxygen species and NO [25]. To manifest this effect, the frequency of the external magnetic effect must coincide with the natural frequency of concentration fluctuations NAD(P)H in a cell, which is about 20 s, and the presence of calcium ions in the medium is necessary for the perception of an external magnetic (or electric) field, which indicates the participation of calcium channels in magneto reception [25].

Revealing the nature and parameters of an external factor regulating proliferative activity of isolated cells will make it possible to develop a method for modulating regeneration processes.

Based on the exogenous mechanism of the about 4-day rhythm formation of the cell culture proliferative activity and indirect evidence of the coincidence of the phase of the 4-day rhythm of the proliferative activity of the cell culture *in vitro* with the similar rhythm of mitotic activity *in vivo*, it can be suggested that the infradian rhythms of other biological parameters *in vivo* are caused by a constantly or almost

constantly manifested rhythmic factor of the environment. However, this statement requires further experimental verification.

## **Material And Methods**

### **Animals**

3 male and 10 female C57Bl/6 mice were used for obtained embryonic fibroblast-like cells. 7 male C57Bl/6 mice were used for estimation of motor activity. All animals were purchased from the “Stolbovaya” branch of the Federal State Budgetary Institution of Science and Technology of the Federal Medical and Biological Agency of Russia. The study received permission from the Bioethics Committee of the Research Institute of Human Morphology (Protocol No. 24, September 2, 2020). The study is done in accordance with ARRIVE guidelines (<https://arriveguidelines.org>) and all methods were carried out in accordance with relevant guidelines and regulations. All manipulations with animals were carried out according to the European convention for the protection of vertebrate animals used for experimental and other scientific purposes (ets no. 123), Strasbourg, 2006, and all efforts were made to minimize the suffering and distress of animals. 5 female mice per cages (40x14.5x24 cm) were housed in temperature-regulated room at 12:12 h light–dark cycle, relative humidity, between 55 and 65%; and unlimited access to water and food (“Char”, JSC “Range-Agro”, Russia). Male mice (n=10) were kept in individual plastic cages (40x14.5x24 cm).

### **Obtaining of embryonic fibroblast-like cells**

Fibroblast-like cells were isolated from 12-13th day embryos of pregnant C57Bl/6 female mice. In this period of the development an embryo contains a high percentage of undifferentiated mesenchyme, which is the main source of fibroblasts [26]. In order to establish a dated gestation date, female mice were housed with males for only one night. Accordingly, after 12 days, the females were scarified by dislocation of the cervical vertebrae. For the following study embryos were taken without internal organs and head. In total, 5 cultures were obtained in the work from 5 pregnant female mice respectively (Table 2). To obtain a culture, embryos were taken from one female. The fibroblast-like cells obtained from different embryos of one female were mixed with each other. The number of embryos taken depended on the volume of cell culture required for the experiment.

Table 2  
Calendar date of experimental manipulations with embryonic cells culture from November, 2019 to January, 2020

Number of culture	Conception date	Date of separation of the primary culture	Freezing date	Defrost date	Initiating date of culture daily sowing on Petri dishes	Initiating date of daily cell counting	Finished date
1	Nov, 4	Nov, 16	-	-	Nov, 20	Nov, 21	Dec, 13
2	Nov, 6	Nov, 18	-	-	Nov, 22	Nov, 23	Dec, 7
3	Nov, 17	Nov, 29	-	-	Dec, 3	Dec, 4	Dec, 21
4	Nov, 19	Dec, 1	-	-	Dec, 5	Dec, 6	Dec, 20
5	Dec, 3	Dec, 16	Dec, 20	Dec, 26	Dec, 31	Jan, 1	Jan, 15
			Dec, 22	Dec, 26	Dec, 31	Jan, 1	Jan, 15
			Dec, 26	Dec, 28	Jan, 2	Jan, 3	Jan, 15

## Culturing of embryonic fibroblast-like cells

We used the method of enzymatic disaggregation to isolate cells: the embryos were placed in a Petri dish (diameter 6 cm) with 2 ml of 0.25% trypsin (PanEco, Russia), crushed into pieces 1-0.5 mm in size. Then 3 ml of trypsin was added and incubated in a thermostat at 37 ° C for 15-30 minutes, pipetted every 5 minutes. Then it was centrifuged for 7 min at 100g, the pellet was resuspended in a medium with 10% fetal bovine serum [26]. Cells in the amount of 1-2 million in a volume of 10 ml of complete culture medium were placed in mattresses of 25 cm<sup>2</sup> for 4 hours, after that the medium was changed on the new one. 3-5 days after sowing, the culture was subcultured into a new flask or used in the experiment.

The cells were cultured in DMEM/F12 medium with L-glutamine (PanEco, Russia) 10% FBS (Biosera France) and 0.004% gentamicin (Borisov Medicines Plant, Belarus) at 37°C in a humid atmosphere containing 5% CO<sub>2</sub> [26].

After subculturing, cells in the amount of 300 000 were plated on 12 Petri dishes with a diameter of 6 cm in a volume of 5 ml of medium. To standardize the experiment and the constant presence of cells in the logarithmic growth phase, the cells were seeded in Petri dishes daily during the study period. The experimental scheme is shown in Table 3.

Table 3  
Date of cell counting in embryonic cells culture

		Date of cell counting in Petri dishes																							
		Nov 21	Nov 22	Nov 23	Nov 24	Nov 25	Nov 26	Nov 27	Nov 28	Nov 29	Nov 30	Dec 1	Dec 2	Dec 3	Dec 4	Dec 5	Dec 6	Dec 7	Dec 8	Dec 9	Dec 10	Dec 11	Dec 12	Dec 13	
Date of culture daily sowing on Petri dishes	Nov,20																								
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\* Black block – is a date of cell counting of embryonic fibroblast-like cell culture

Each subsequent day, for the purpose of counting, the cells were removed from 2 Petri dishes by trypsinization: 2 ml of 0.02% chymotrypsin diluted in Versene solution was added and incubated for 4 min at 37°C. Cells were counted with using an automatic cell counter TC-20 (Bio-Rad Laboratories, USA).

## Freezing and thawing of embryonic fibroblast-like cells

To change the phase of the infradian rhythm of the proliferative activity of embryonic fibroblast-like cells, they were exposed to short-term freezing. Culture samples from the embryos of one female were frozen on December 20, 22 and 26. Procedure of freezing included the next steps:  $2-3 \times 10^6$  cells were added to 0.5 ml of cryopreservation medium (90% FBS and 10% DMSO), the cell suspension was poured into ampoules for freezing. To ensure a cooling rate of 1°C/min, the ampoules were placed on cotton wool in a polystyrene foam box with walls 15 mm thick and put them into a -70°C fridge. After 2 hours, when the ampoules reached -70°C, they were immersed in a liquid nitrogen freezer.

In 2, 4 and 6 days after freezing, two cells cultures were defrosted in a thermostat at 37° C, respectively, on December 26 and 29, 2019. After defrosting, a small amount of growth medium was slowly added to the ampoule, then the contents of the ampoule were transferred to a centrifuge tube and the medium was slowly added to the end volume of 10 ml. It was centrifuged for 3 min at 100g. The cells were resuspended in fresh growth medium and then transferred to culture flacons. Then cells were cultured in a mattress for 5 days, and then they were plated on Petri dishes. The experimental and cell counting scheme did not differ from that presented above for all other cultures (Table 3).

## Culturing of L929 cells

The L-929 culture of transformed mouse fibroblasts was obtained from the collection of vertebrate cell cultures of the Institute of Cytology, Russian Academy of Sciences (Russia, St. Petersburg). The L-929 cells were cultured in growth medium DMEM/F12 with L-glutamine (Capricorn Scientific, Germany), 10% FBS (Biosera France), 50 U/ml penicillin and 50 µg/ml streptomycin (PanEco, Russia).

To obtain a series of identical samples, cells after several passages were simultaneously frozen. DMSO (dimethyl sulfoxide) was used as a cryoprotectant for freezing the L-929 culture. The cell suspension of L-929 was frozen in 10% DMSO and 90% of cattle serum, for this,  $2.5 \times 10^6$  cells were diluted in 1 ml of freezing medium and placed in cryotubes. To ensure cooling of cryoprobes at a rate of 1°C/ min, the cryotubes were placed on cotton wool in a polystyrene foam box with 15 mm wall thickness, then put into a low-temperature freezer at -70 ° C.

The culture was thawed by placing it in a thermostat for 5 min at 37°C, then centrifuged at 200g for 4 min, resuspended in a growth medium and centrifuged again. After centrifugation, it was diluted in 2 ml of growth medium. Viable cells were counted on an automatic cell counter TC-20 (Bio-Rad Laboratories, USA). Viable cells of one cryoprobe with  $150 \times 10^3$  cells in 3 ml of growth medium were plated on Petri dishes 35 mm in diameter.

## Freezing and thawing of L929 cells

We carried out two series of experiments with freezing/thawing of the L-929 culture. To assess the proliferative activity of cells in the logarithmic growth phase, every day from January 14 to February 3 and from February 15 to March 3, 2021 we thawed one cryoprobe (Table 4). In the first series, the indicators of the increase in the cells number after the 2nd, 3rd, 4th and 5th days of cultivation were analyzed, while the cells were counted daily from one of the Petri dishes (Tables 5 and 6). In the second series, the indicators were obtained only after the 2nd and 3rd days of cultivation, three Petri dishes from each cryoprobe were analyzed daily (Table 5 and 6).

Table 4  
Calendar date of experimental manipulations with L-929 cells

Nº series	Freezing date	Defrost date of the first cryosample	Defrost date of the last cryosample	Number of repetitions	Number of days after seeding, when the determination of the number of cells in the Petri dish was carried out	The duration of the obtained data series of the daily increase in the number of cells
1	06.01.21	14.01.21	03.02.21	1	1, 2, 3, 4, 5,	22 days
2	10.02.21	15.02.21	03.03.21	3	2, 3	17 days

Table 5.

Date of cell counting in L-929 cells culture from January, 15 to February, 6, 2021

		Date of cell counting in Petri dishes																							
		Jan, 15	Jan, 16	Jan, 17	Jan, 18	Jan, 19	Jan, 20	Jan, 21	Jan, 22	Jan, 23	Jan, 24	Jan, 25	Jan, 26	Jan, 27	Jan, 28	Jan, 29	Jan, 30	Jan, 31	Feb, 1	Feb, 2	Feb, 3	Feb, 4	Feb, 5	Feb, 6	
Date of culture daily sowing on Petri dishes	Jan,14																								
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\* Black block – is a date of cell counting of L-929 culture

Table 6

Date of cell counting in L-929 cells culture from February, 17 to March, 6, 2021

		Date of cell counting in Petri dishes																					
		Feb, 17	Feb, 18	Feb, 19	Feb, 20	Feb, 21	Feb, 22	Feb, 23	Feb, 24	Feb, 25	Feb, 26	Feb, 27	Feb, 28	Mar, 1	Mar, 2	Mar, 3	Mar, 4	Mar, 5	Mar, 6				
Date of culture daily sowing on Petri dishes	Feb, 15																						
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\* Black block – is a date of cell counting of L-929 culture

# Motor activity of mice

To compare the phases of the 4-day infradian rhythm in the fibroblast-like cell culture and in the organism we determined the rhythm of motor activity of mice. The motor activity of the animals was determined with intraperitoneally implanted DTN4-28/TL4-28 sensors (EMBI RESERCH, Novosibirsk). Implantation of sensors into the abdominal cavity had been carried out under anesthesia by Zoletil (Virbac Sante Animale, France) at the dose of 5-7 mg/kg. Measurements were carried out at a frequency of 1 time per minute. The total daily activity was determined as the sum of the minute readings of the accelerometer built into the sensor.

## **Statistical analyzes**

Statistical analyzes was performed using the Statistica 7.0 software package. The obtained data were expressed as the median and interquartile range Me (Q1-Q3). To identify the periods of infradian rhythms, the autocorrelation coefficient was calculated between the original series and the series shifted by 1, 2, 3, and 4 days. In order to identify the synchronicity of changes in the dynamics of the studied parameters, the Spearman correlation coefficient was calculated. Differences were considered statistically significant at  $p < 0.05$ .

## **Declarations**

## **Acknowledgments**

The work was performed within the State Task of the Research Institute of Human Morphology AAAA-A19-119021490067-4.

## **Author contributions**

Marina A. Diatroptova, Anna M. Kosyreva & Mikhail E. Diatroptov contributed equally.

## **Contributions**

M.A.D. and M.E.D. designed the project, analysed and discussed the data. A.M.K. and M.A.D. performed the experiment, prepared samples and analysed data. M.A.D., A.M.K. and M.E.D. prepared final manuscript

## **Conflict of interest statement**

The authors have no potential conflicts of interest with respect to the research, authorship and/or publication of this article.

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## Figures

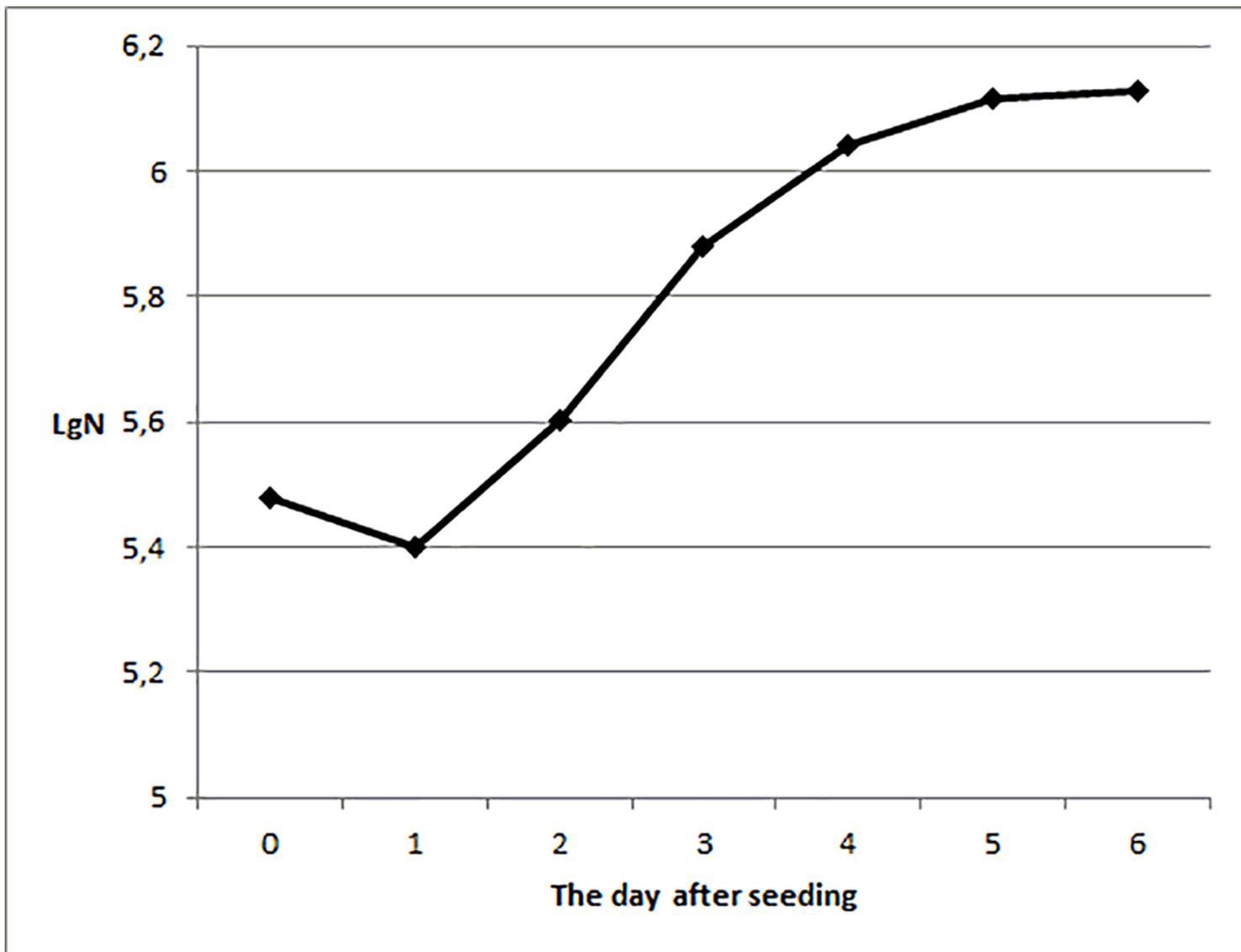
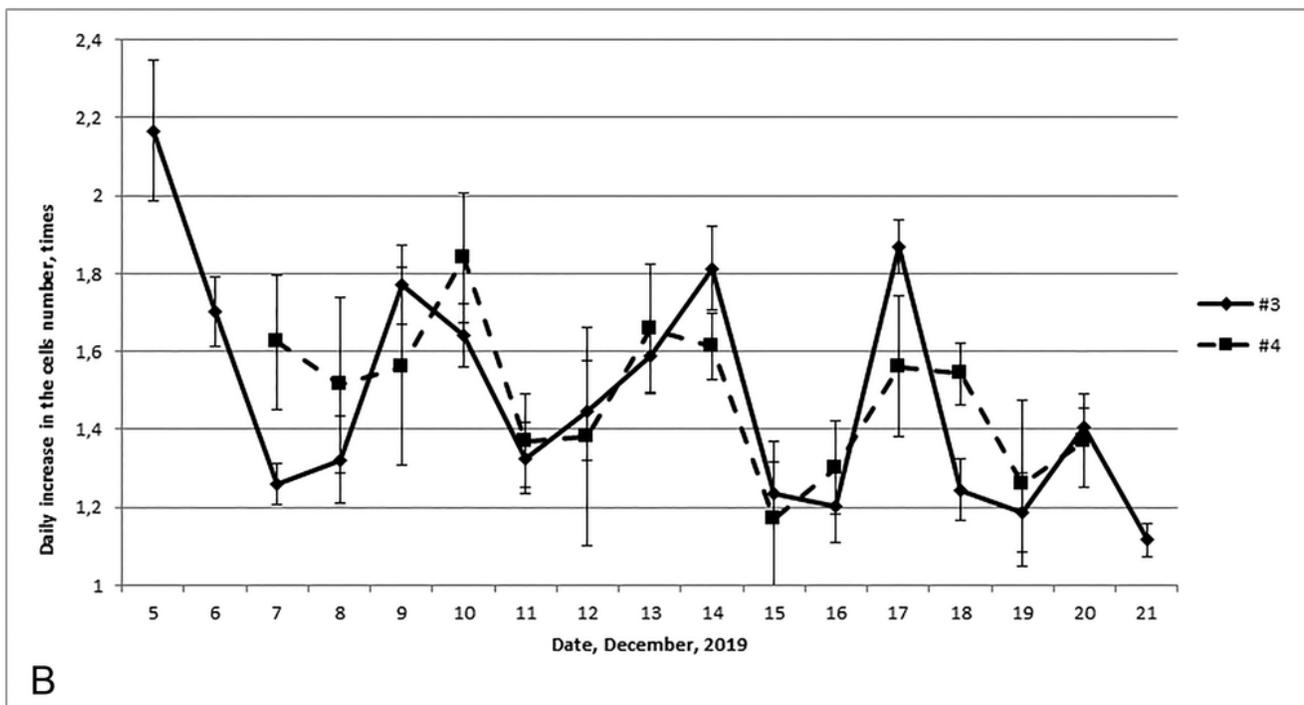
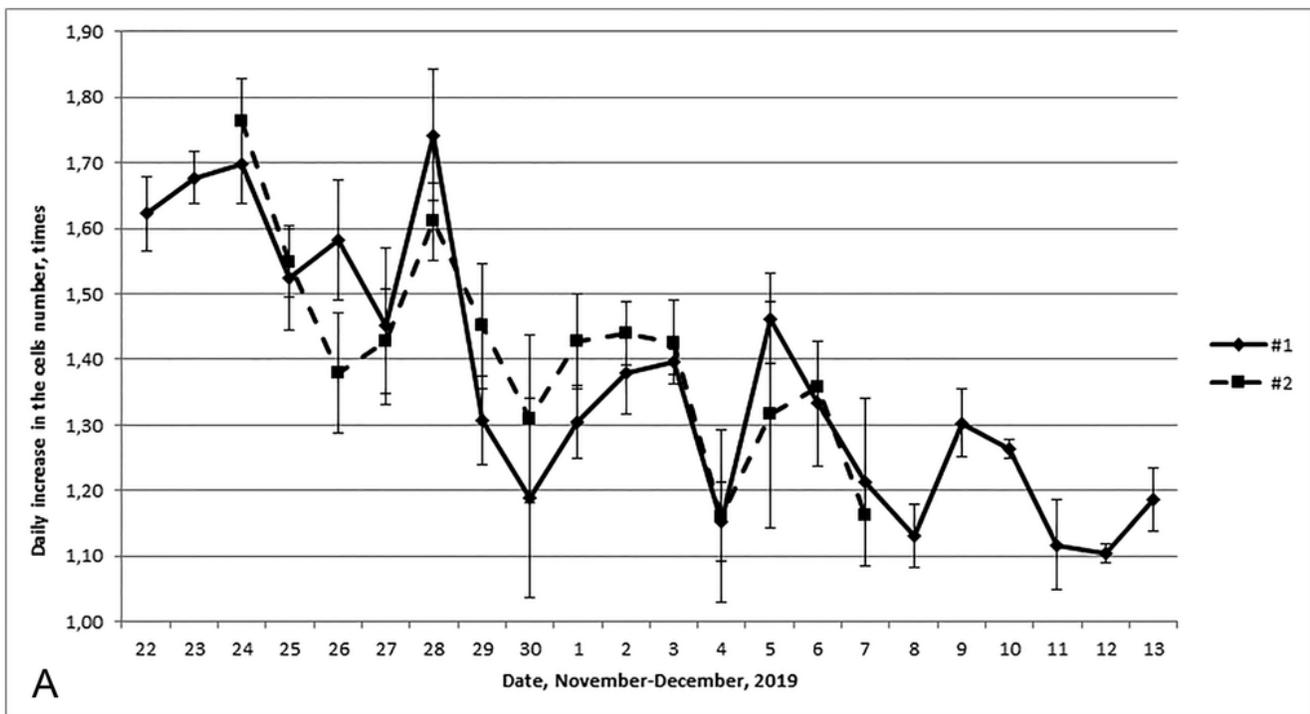


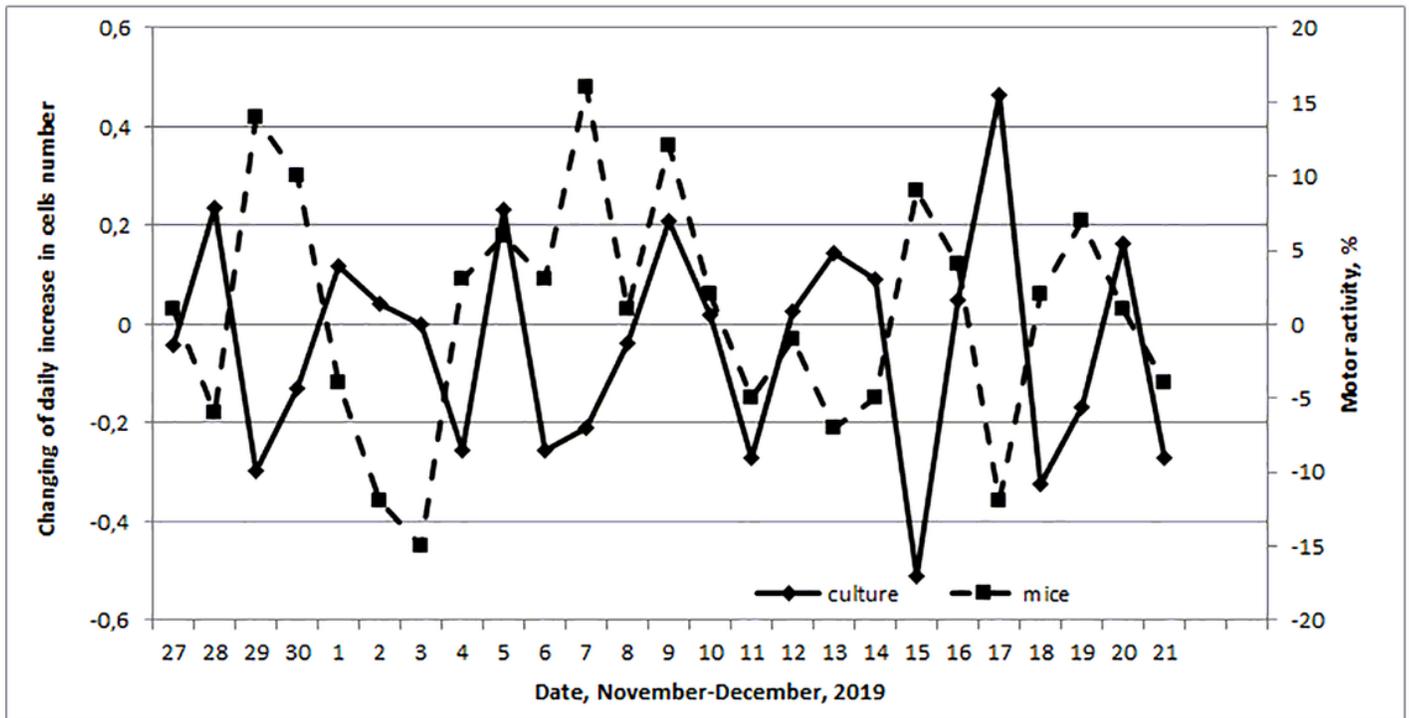
Figure 1

Typical growth curve of the cell culture of embryonic fibroblast-like cells.



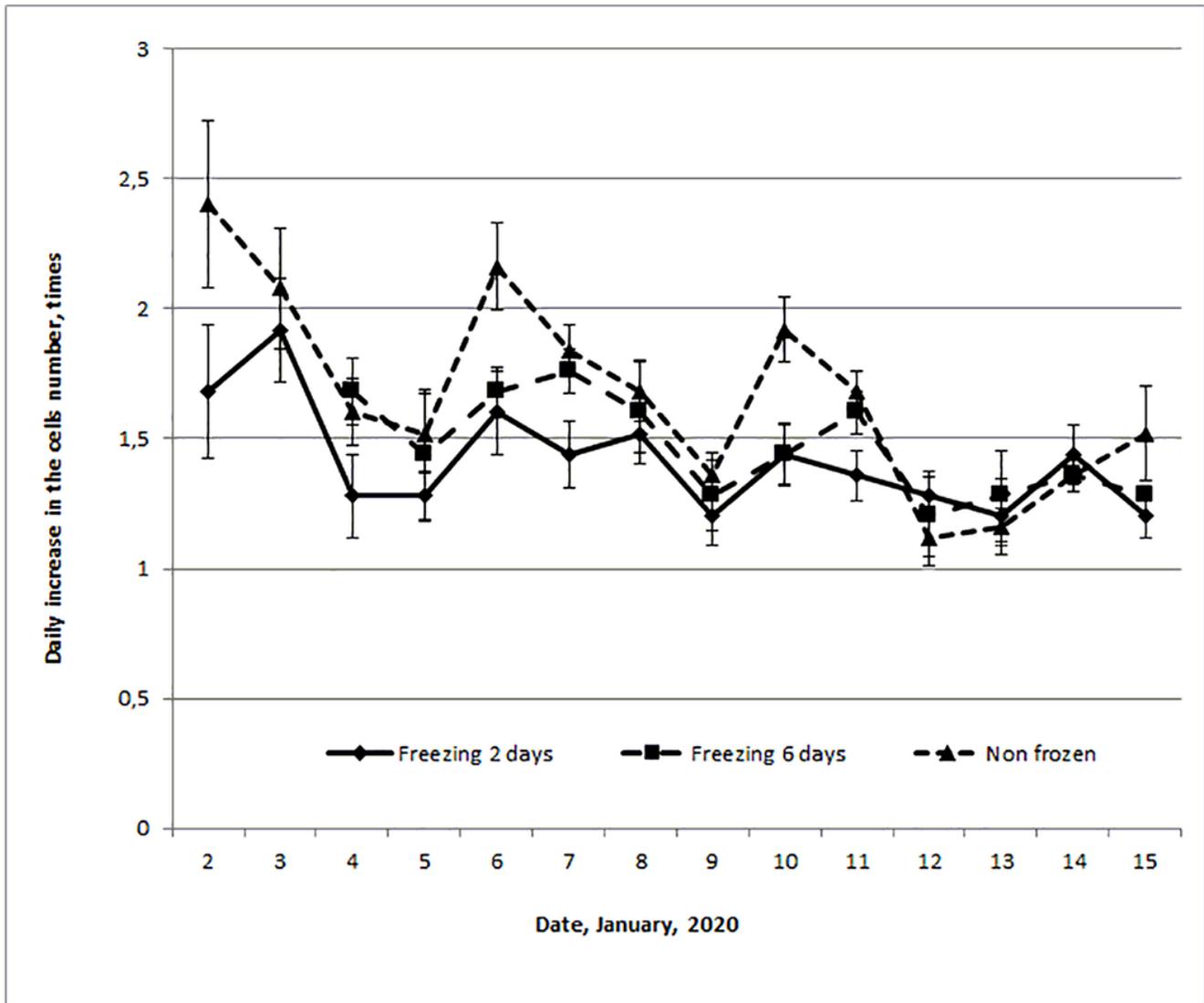
**Figure 2**

Dynamics of the daily increase in fibroblast-like cells number during the period of the logarithmic growth state obtained in different phases of the 4-day rhythm: (A) row 1 - November 16, 2019; row 2 - November 18, 2019. (B) row 1 - November 29, 2019; row 2 - December 1, 2019.



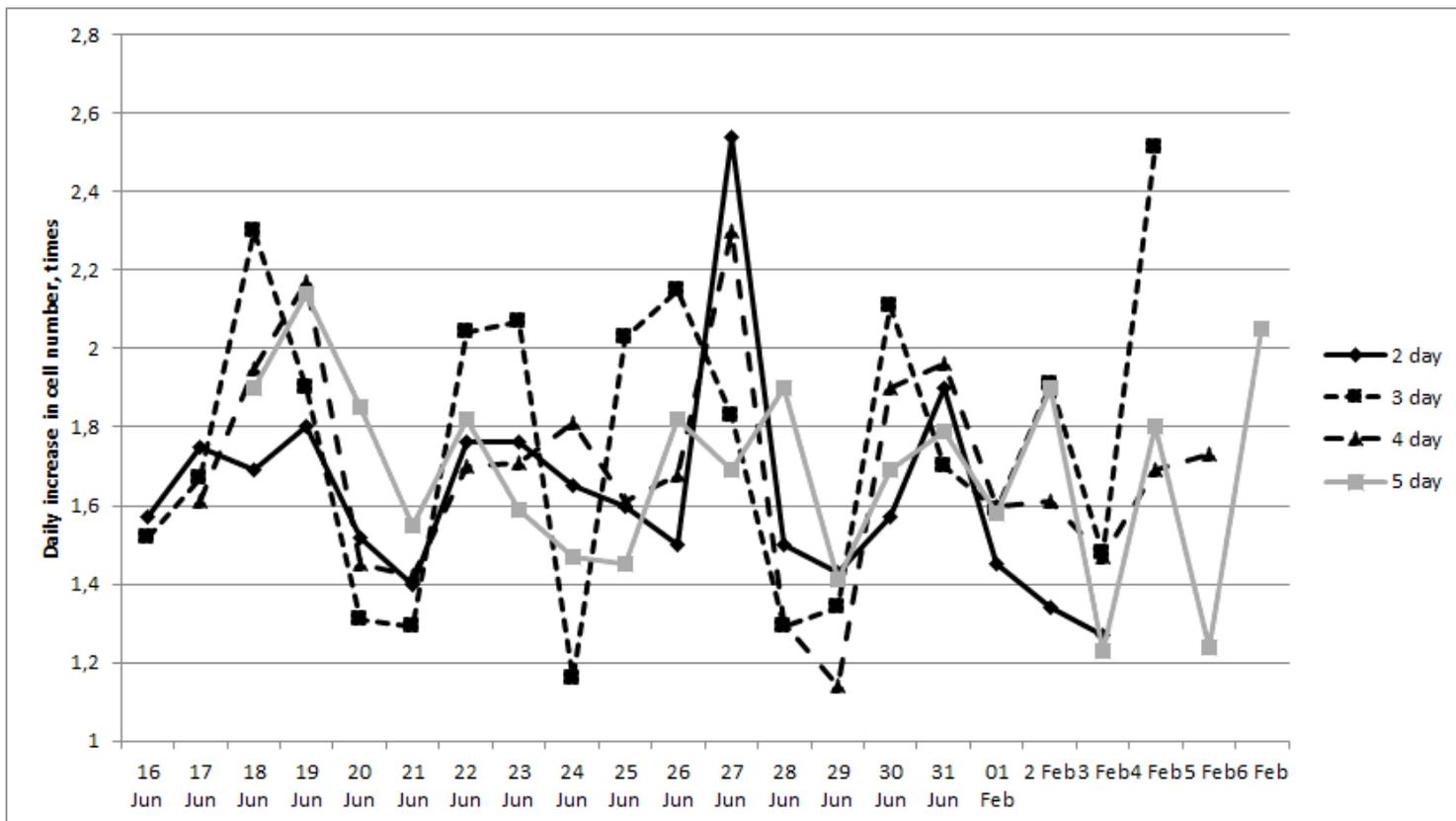
**Figure 3**

Dynamics of the deviation of the daily increase in the fibroblast-like cells number from the trend line (linear approximation) in comparison with the total motor activity of male mice (n = 7); the percentage deviations from the individual mean level are presented.



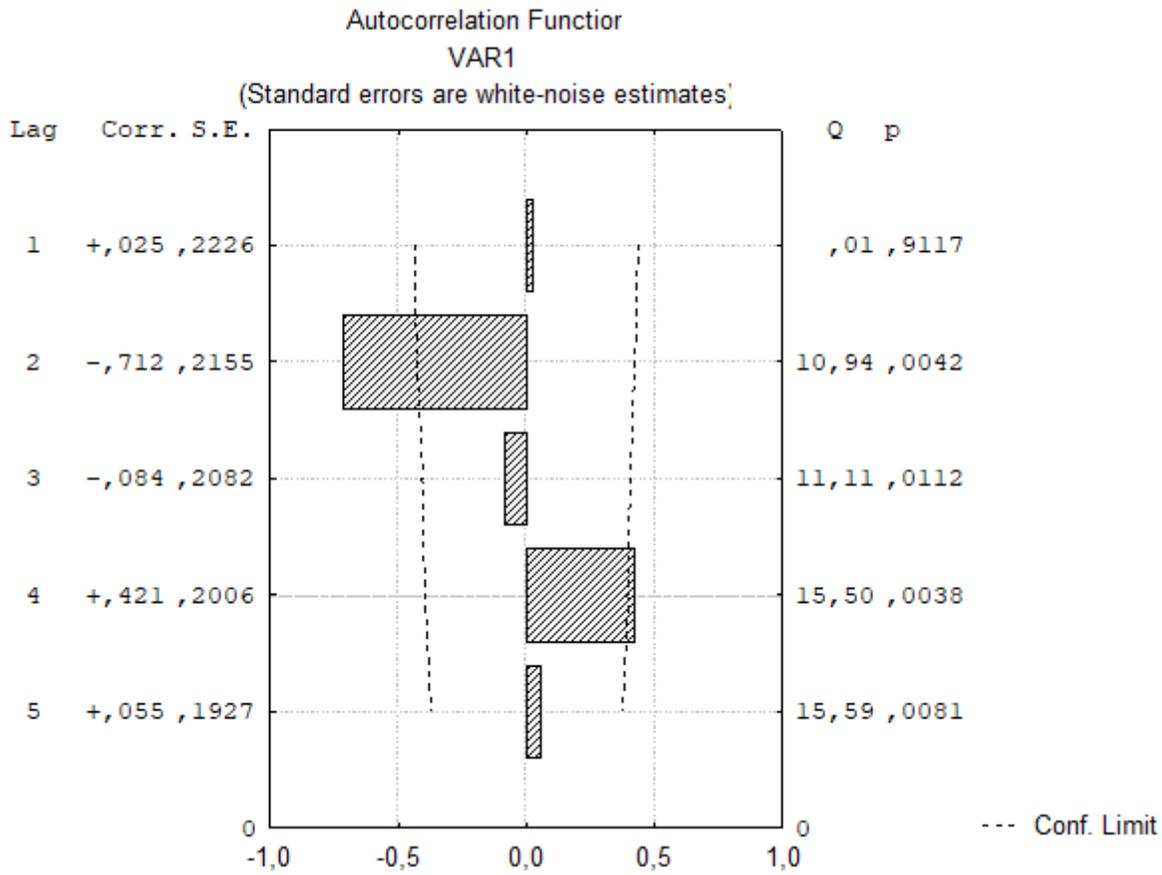
**Figure 4**

Dynamics of the daily increase in the fibroblast-like cells number of single culture, some of them were frozen for 2 and 6 days (rows 1 and 2), and some were cultured as usual (row 3).



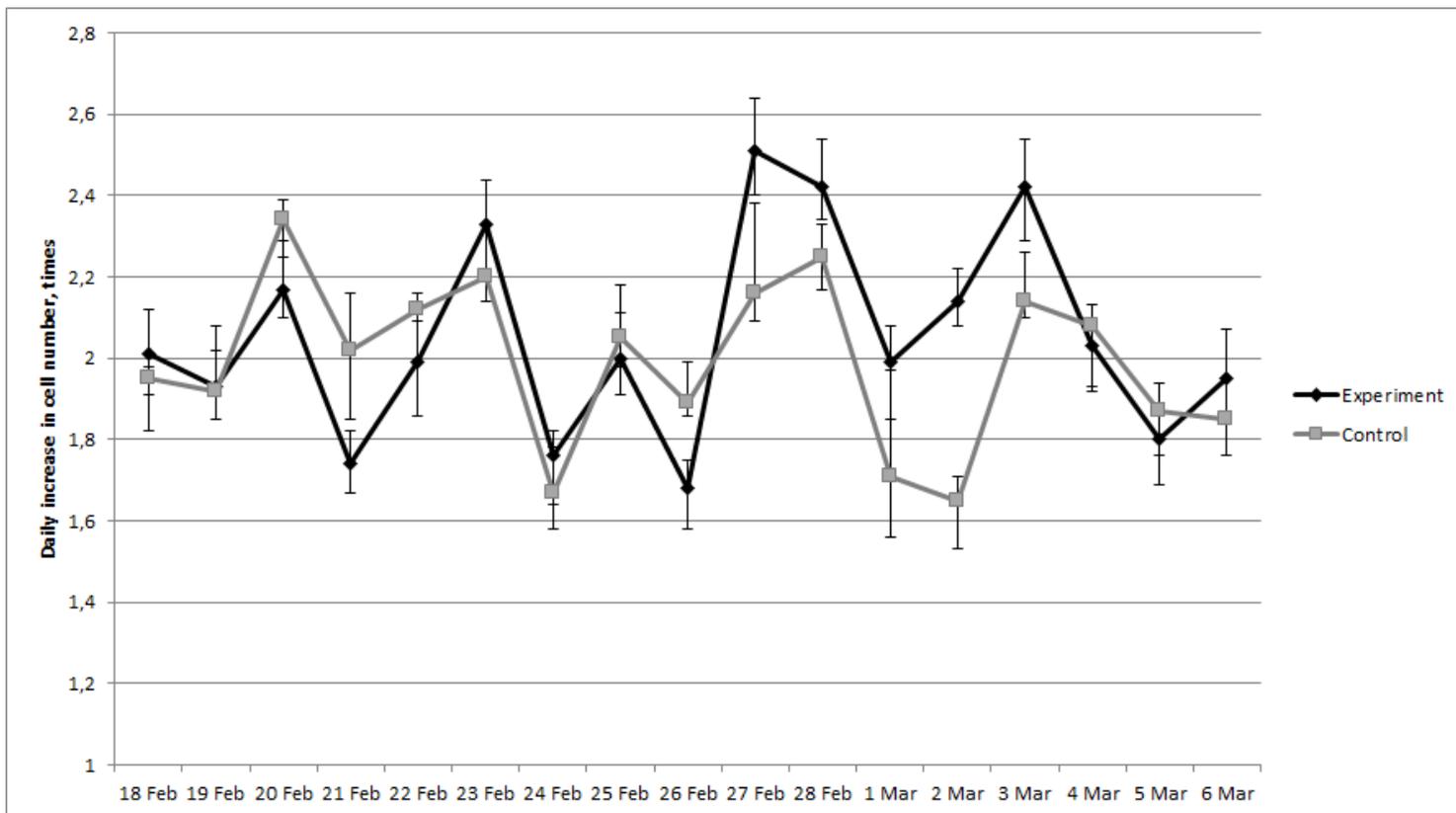
**Figure 5**

Dynamics of the daily increase in the L-929 cells number in the samples of the culture, analyzed in 2, 3, 4 and 5 days after thawing.



**Figure 6**

Autocorrelation function of the dynamics of the daily increase in the number of L929 cells from January 16, 2021 to February 1, 2021



**Figure 7**

Dynamics of the daily increase in the L-929 cells number after thawing (experiment) and in not frozen culture (control).