

ISSN 2518-1629 (Online),  
ISSN 2224-5308 (Print)

ҚАЗАҚСТАН РЕСПУБЛИКАСЫ  
ҰЛТТЫҚ ҒЫЛЫМ АКАДЕМИЯСЫНЫҢ  
Өсімдіктердің биологиясы және биотехнологиясы институтының

# Х А Б А Р Л А Р Ы

---

---

## ИЗВЕСТИЯ

НАЦИОНАЛЬНОЙ АКАДЕМИИ НАУК  
РЕСПУБЛИКИ КАЗАХСТАН  
Института биологии и биотехнологии растений

## NEWS

OF THE NATIONAL ACADEMY OF SCIENCES  
OF THE REPUBLIC OF KAZAKHSTAN  
of the Institute of Plant Biology and Biotechnology

**БИОЛОГИЯ ЖӘНЕ МЕДИЦИНА  
СЕРИЯСЫ**



**СЕРИЯ**

**БИОЛОГИЧЕСКАЯ И МЕДИЦИНСКАЯ**



**SERIES**

**OF BIOLOGICAL AND MEDICAL**

**2 (320)**

**НАУРЫЗ – СӘУІР 2017 ж.**

**МАРТ – АПРЕЛЬ 2017 г.**

**MARCH – APRIL 2017**

1963 ЖЫЛДЫҢ ҚАҢТАР АЙЫНАН ШЫҒА БАСТАҒАН  
ИЗДАЕТСЯ С ЯНВАРЯ 1963 ГОДА  
PUBLISHED SINCE JANUARY 1963

ЖЫЛЫНА 6 РЕТ ШЫҒАДЫ  
ВЫХОДИТ 6 РАЗ В ГОД  
PUBLISHED 6 TIMES A YEAR

АЛМАТЫ, ҚР ҰҒА  
АЛМАТЫ, НАН РК  
ALMATY, NAS RK

Б а с р е д а к т о р

ҚР ҰҒА академигі, м. ғ. д., проф.

**Ж. А. Арзықұлов**

**Абжанов Архат** проф. (Бостон, АҚШ),  
**Абелев С.К.** проф. (Мәскеу, Ресей),  
**Айтқожина Н.А.** проф., академик (Қазақстан)  
**Акшулаков С.К.** проф., корр.-мүшесі (Қазақстан)  
**Алшынбаев М.К.** проф., корр.-мүшесі (Қазақстан)  
**Березин В.Э.**, проф., корр.-мүшесі (Қазақстан)  
**Бисенбаев А.К.** проф., корр.-мүшесі (Қазақстан)  
**Бишимбаева Н.К.** проф., корр.-мүшесі (Қазақстан)  
**Ботабекова Т.К.** проф., корр.-мүшесі (Қазақстан)  
**Ellenbogen Adrian** prof. (Tel-Aviv, Israel),  
**Жамбакин К.Ж.** проф., корр.-мүшесі (Қазақстан), бас ред. орынбасары  
**Ishchenko Alexander**, prof. (Villejuif, France)  
**Қайдарова Д.Р.** проф., корр.-мүшесі (Қазақстан)  
**Күзденбаева Р.С.** проф., академик (Қазақстан)  
**Лось Д.А.** prof. (Мәскеу, Ресей)  
**Lunefeld Bruno** prof. (Израиль)  
**Миербеков Е.М.** проф. (Қазақстан)  
**Муминов Т.А.** проф., академик (Қазақстан)  
**Purton Saul** prof. (London, UK)  
**Рахыпбеков Т.К.** проф., корр.-мүшесі (Қазақстан)  
**Сапарбаев Мұрат** проф. (Париж, Франция)  
**Сарбассов Дос** проф. (Хьюстон, АҚШ)

«ҚР ҰҒА Хабарлары. Биология және медициналық сериясы».

**ISSN 2518-1629 (Online),**

**ISSN 2224-5308 (Print)**

Меншіктенуші: «Қазақстан Республикасының Ұлттық ғылым академиясы» РҚБ (Алматы қ.)

Қазақстан республикасының Мәдениет пен ақпарат министрлігінің Ақпарат және мұрағат комитетінде  
01.06.2006 ж. берілген №5546-Ж мерзімдік басылым тіркеуіне қойылу туралы куәлік

Мерзімділігі: жылына 6 рет.

Тиражы: 300 дана.

Редакцияның мекенжайы: 050010, Алматы қ., Шевченко көш., 28, 219 бөл., 220, тел.: 272-13-19, 272-13-18,  
[www.nauka-nanrk.kz](http://www.nauka-nanrk.kz) / [biological-medical.kz](http://biological-medical.kz)

---

© Қазақстан Республикасының Ұлттық ғылым академиясы, 2017

Типографияның мекенжайы: «Аруна» ЖК, Алматы қ., Муратбаева көш., 75.

Г л а в н ы й р е д а к т о р  
академик НАН РК, д.м.н., проф.

**Ж. А. Арзыкулов**

**Абжанов Архат** проф. (Бостон, США),  
**Абелев С.К.** проф. (Москва, Россия),  
**Айтхожина Н.А.** проф., академик (Казахстан)  
**Акшулаков С.К.** проф., чл.-корр. (Казахстан)  
**Алчинбаев М.К.** проф., чл.-корр. (Казахстан)  
**Березин В.Э.**, проф., чл.-корр. (Казахстан)  
**Бисенбаев А.К.** проф., чл.-корр. (Казахстан)  
**Бишимбаева Н.К.** проф., чл.-корр. (Казахстан)  
**Ботабекова Т.К.** проф., чл.-корр. (Казахстан)  
**Ellenbogen Adrian** prof. (Tel-Aviv, Israel),  
**Жамбакин К.Ж.** проф., чл.-корр. (Казахстан), зам. гл. ред.  
**Ishchenko Alexander** prof. (Villejuif, France)  
**Кайдарова Д.Р.** проф., чл.-корр. (Казахстан)  
**Кузденбаева Р.С.** проф., академик (Казахстан)  
**Лось Д.А.** prof. (Москва, Россия)  
**Lunenfeld Bruno** prof. (Израиль)  
**Миербеков Е.М.** проф. (Казахстан)  
**Муминов Т.А.** проф., академик (Казахстан)  
**Purton Saul** prof. (London, UK)  
**Рахыпбеков Т.К.** проф., чл.-корр. (Казахстан)  
**Сапарбаев Мурат** проф. (Париж, Франция)  
**Сарбассов Дос** проф. (Хьюстон, США)

«Известия НАН РК. Серия биологическая и медицинская».

**ISSN 2518-1629 (Online),**

**ISSN 2224-5308 (Print)**

Собственник: РОО «Национальная академия наук Республики Казахстан» (г. Алматы)

Свидетельство о постановке на учет периодического печатного издания в Комитете информации и архивов  
Министерства культуры и информации Республики Казахстан №5546-Ж, выданное 01.06.2006 г.

Периодичность: 6 раз в год

Тираж: 300 экземпляров

Адрес редакции: 050010, г. Алматы, ул. Шевченко, 28, ком. 219, 220, тел. 272-13-19, 272-13-18,

[www.nauka-nanrk.kz/biological-medical.kz](http://www.nauka-nanrk.kz/biological-medical.kz)

---

© Национальная академия наук Республики Казахстан, 2017

Адрес типографии: ИП «Аруна», г. Алматы, ул. Муратбаева, 75

Editor in chief

academician of NAS RK, doctor of medical science, professor

**Zh. A. Arzykulov**

**Abzhanov Arkhat** prof. (Boston, USA),  
**Abelev S.K.** prof. (Moscow, Russia),  
**Aitkhozhina N.A.** prof., academician (Kazakhstan)  
**Akshulakov S.K.** prof., corr. member. (Kazakhstan)  
**Alchinbayev M.K.** prof., corr. member. (Kazakhstan)  
**Berezin V.Ye.**, prof., corr. member. (Kazakhstan)  
**Bisenbayev A.K.** prof., corr. member. (Kazakhstan)  
**Bishimbayeva N.K.** prof., corr. member. (Kazakhstan)  
**Botabekova T.K.** prof., corr. member. (Kazakhstan)  
**Ellenbogen Adrian** prof. (Tel-Aviv, Israel),  
**Zhambakin K.Zh.** prof., corr. member. (Kazakhstan), deputy editor in chief  
**Ishchenko Alexander**, prof. (Villejuif, France)  
**Kaydarova D.R.** prof., corr. member. (Kazakhstan)  
**Kuzdenbayeva R.S.** prof., academician (Kazakhstan)  
**Los D.A.** prof. (Moscow, Russia)  
**Lunefeld Bruno** prof. (Israel)  
**Miyerbekov Ye.M.** prof. (Kazakhstan)  
**Muminov T.A.** prof., academician (Kazakhstan)  
**Purton Saul** prof. (London, UK)  
**Rakhypbekov T.K.** prof., corr. member. (Kazakhstan)  
**Saparbayev Murat** prof. (Paris, France)  
**Sarbassov Dos**, prof. (Houston, USA)

**News of the National Academy of Sciences of the Republic of Kazakhstan. Series of biology and medicine.**

**ISSN 2518-1629 (Online),**

**ISSN 2224-5308 (Print)**

Owner: RPA "National Academy of Sciences of the Republic of Kazakhstan" (Almaty)

The certificate of registration of a periodic printed publication in the Committee of information and archives of the Ministry of culture and information of the Republic of Kazakhstan N 5546-Ж, issued 01.06.2006

Periodicity: 6 times a year

Circulation: 300 copies

Editorial address: 28, Shevchenko str., of. 219, 220, Almaty, 050010, tel. 272-13-19, 272-13-18,

<http://nauka-nanrk.kz/biological-medical.kz>

---

© National Academy of Sciences of the Republic of Kazakhstan, 2017

Address of printing house: ST "Aruna", 75, Muratbayev str, Almaty

## NEWS

OF THE NATIONAL ACADEMY OF SCIENCES OF THE REPUBLIC OF KAZAKHSTAN

## SERIES OF BIOLOGICAL AND MEDICAL

ISSN 2224-5308

Volume 2, Number 320 (2017), 27 – 33

UDK581.4:581.1

**Dzhangalina Erika, Zhumabayeva Beibitgul, Aitasheva Zaure, Lebedeva Lina**

Al-Farabi Kazakh National University,  
SRI of Biology and Biotechnology Problems MES RK, Almaty, Kazakhstan.  
E-mail: Erika.Dzhangalina@kaznu.kz

**BIOLOGICAL ACTIVITY OF LECTINS EXTRACTED  
FROM LEGUMES' CALLUSES**

**Abstract.** Nowadays cell and tissue cultures *in vitro* are widely used in many biological experiments, particularly in production of natural biologically active food supplements and protein substances for different purposes. Legumes are characterized by high level of proteins and protein compounds such as lectins. Studying different patterns of hormonal regulation of lectins *in vitro* makes possible understand the physiological mechanisms of growth and development of legumes. In this regard selecting the source of explants and optimizing concentration of minerals and hormones in culture media allowed us to obtain morphogenic and non-morphogenic calluses. Usage of 2,4-D helped to increase frequency of morphogenic calluses more than 5 times in contrast with NAA. Biological activity of lectins, extracted from different types of calluses, was determined visually by intensity of hemagglutination and their titer. It is proved, that lectin activity of legume specimens depends on genotype and type of calluses, which is determined by ingredients of culture media. The level of lectin is controlled and regulated by concentration of auxins and cytokinins in culture media. All morphogenic samples of calluses demonstrated higher activity than non-morphogenic ones. And maximum dose of lectins was observed at concentration of 1 mg/l 2,4-D and 0.5 mg/l of kinetin. The highest lectin activity was marked for two representatives of morphogenic type of calluses – "Aktatti" and "Jura-vushka". Real experiments demonstrated that determination of agglutinative activity in plants and the level of lectins in calluses allows to see fluctuations in lectin activity in cells and tissues. Also this data might be used to study the hormonal processes of differentiation, proliferation and early development of legumes and to improve current methods of lectin extraction.

**Key words:** *Phaseolus vulgaris*, lectins, callus culture, activity, hormonal dependence.

**Introduction.** One of the main functions of lectins is participation in processes of cell division, stretching and differentiation and, consequently, regulation of morphological and physiological processes in plants.

Some hormones such as ABA, IAA, BAP, GA regulate synthesis of lectins. Biological role of these substances is associated with the seeds formation, the stage of rest and the stage of awakening [1, 2]. Mechanism of action of plant growth regulators is controlled by changing in activity and synthesis of different proteins. There is information that lectins have an ability to attach molecules of phytohormones. So, lectin in wheat germ agglutinin has a high affinity to many phytohormones such as auxins, cytokinins and gibberellic acid [3]. It is supposed, that complex of lectins and phytohormones takes participation in storing of hormones and regulation of plants growth [4, 5]. Recently it is not doubt that there is a specific lectin-dependent system to regulate some functions inside animal cells. Probably the similar system might be founded in plants too [6]. In scientific articles there are proves that lectins with phytohormones may be involved in regulation of growth processes in vegetating plants. This suggestion is based on the facts that various lectins can interact with hormones. Independence of kinetin in different culture media at the level of lectin in calluses usually varies [7].

Also it was revealed, that cooperation between K on A with IAA and its derivatives and lectins in lima bean with cytokinins is possible because of hydrophobic binding sites in lectins [8].

It is shown that PBA can result in a rapid changing in hormonal system of beans, demonstrating an ability of PBA complex and phytohormones to regulate proliferation in roots. Obtained results prove that various phytolectins are involved in regulation such an important process as a plant growth [9].

Thus, one of the most prospective ways to study physiological mechanisms of growth regulation of legumes is a chance to observe different patterns of hormonal regulation of quantitative content in legumes *in vitro*.

The aim of the research is to determine biological activity and lectin concentration in beans calluses, having differences in morphogenetic capabilities, to find sources of lectins and study the process of lectin regulation.

**Materials and methods.** To get calluses specimens «Aktatti», «Juravushka», «Kamelia», «Red Goya» were used. Primary callus cultures were cultivated on a special media Murasige-Skuga, containing 2.4-D (2-8 mg/l) and kinetin (0.25 mg/l). The role of explants was played by epikotyls of 7-14-days aseptic seedlings. Calluses were grown at 23-25 °C in the light, during 16-hours photoperiod with light intensity equal to 10 000 lux. Passage occurred once per 28 days. Lectins were extracted from calluses according to the method invented by us in the previous researches [10]. The content of lectins was determined right after extraction and dialysis. Extracted mass was weighted and calculated in mg per 100g of wet weight. The lectin activity was defined by reaction of hemagglutination with rabbit blood.

The lectin activity was determined by its titer and measured in units of inverse  $[\text{mg/ml}]^{-1}$ . It means the minimal concentration of protein leads to hemagglutination. For this purpose lectins were diluted five times [11]. To determine the impact of phytohormones on lectins accumulation, calluses were grown on a culture media, containing 1 mg/l of 2.4-D and 0-10 mg/l of kinetin. The content of lectins was fixed after 21 days. To exclude mistakes and errors this experiment was repeated three times. Microphotographies were given by Motic DM 143 SERIES, a microscope.

**Results and discussion.** In many scientific articles it was proved that formation of one or another morphological type of callus tissue might be regulated by different phytohormones, by changing compounds in a media, conditions of cultivation, and also by using species and specimens with a high potential *in vitro*. The last factor is very important for the process of cell and tissue cultivation in some legumes which have different traits appearing at the level of morphogenetic processes [12-14]. In the previous researches it was shown, that for selected specimens of legumes the most intensive process of callus formation occurred on a modified Murasige-Skuga media [15]. Formation of morphogenic and non-morphogenic type of calluses depended on the type and concentration of phytohormones. The presence of 2 mg/l of 2.4-D and IAA in a media resulted in formation of 87% and 15 % of morphogenic calluses respectively. Increasing concentration of 2.4-D provoked a degradation of morphogenic calluses and in presence of 8 mg/l of 2.4-D necrosis was observed. It was supposed that high concentrations of 2.4-D increased the speed of ethylene formation, contrariwise it decreased the speed of cell stretching. Probably high concentrations of auxins influence on suppression of growth of dicotyledonous plants and associate with the synthesis of ethylene [16]. Analysis of extracts showed their differences in doses of lectin. The concentration of lectin in morphogenic calluses was significantly higher than in non-morphogenic ones and varied from 37.3 mg/100g of raw mass in "Juravushka" to 26,0 mg/100g in "Kamelia" (figure 1).

Non-morphogenic calluses are characterized by low concentration of lectins (about 18.4-25.2 mg/100 g of wet weight) in all studied samples. It was assumed that these differences in lectin concentration between morphogenic and non-morphogenic calluses might be related to hormones in the media, because of morphogenic type was formed on culture media with IAA and low concentrations of 2.4-D. According to literature sources it is known that synthesis of lectins is regulated by abscisic acid and high concentrations of 2.4-D decrease the content of ABA [17-19].

The next stage of this research was to determine lectin activity in morphogenic and non-morphogenic calluses of different legumes. In the first serial of experiments the intensity of hemagglutination was assessed visually using bands from 1 to 10. It was found that agglutinating activity in all specimens depended both on genotype and type of callus tissue. In contrast with non-morphogenic type morphogenic calluses demonstrated higher activity which varied from 10 to 13 bands. Two specimens "Aktatti" and "Juravushka" had 13 bands, "Kamelia" had 11 bands, "Red Goya" had 8 bands, but non-morphogenic calluses got not more than 5 bands. Also in "Aktatti" hemagglutination was observed during the first minute after start, in "Juravushka" the same indicator was observed in two minutes. It is necessary to mention

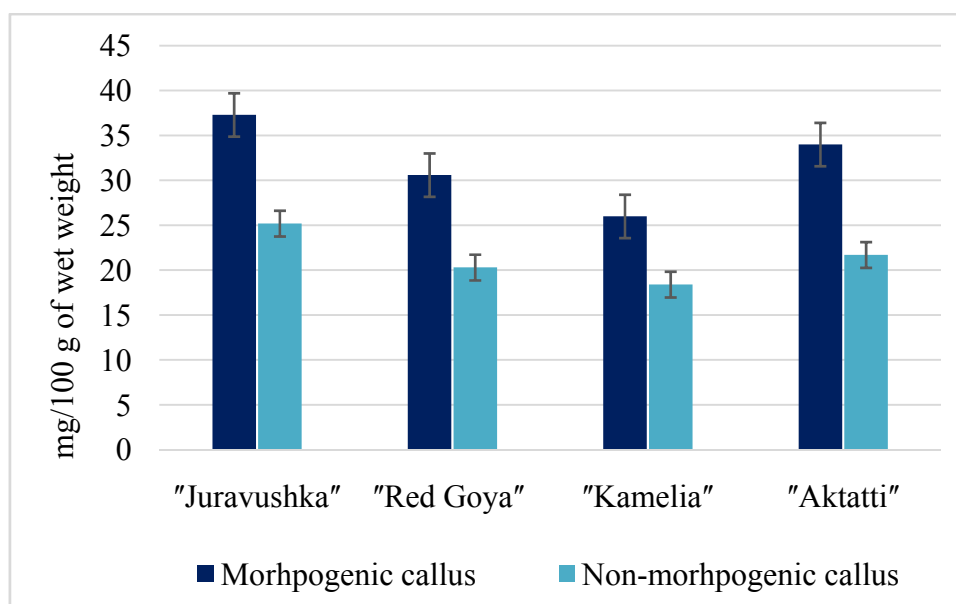


Figure 1 – The presence of lectins in callus tissues of various legume samples

that in this case the effect of hemagglutination was very strong and could be observed even after 5 dilutions. Shaking samples did not provoke destruction of red blood cells, liquid also remained clear and transparent. All samples were differed from the timing of hemagglutination. In "Kamelia" (morphogenic type of calluses) this reaction started later than in other specimens, but it was very strong and could be observed after several dilutions. "Red Goya" (a non-morphogenic one) demonstrated very low lectin activity, as the reaction of agglutination started 55 minutes later and was observed after the first dilution only.

15 minutes later after the reaction of hemagglutination in "Aktatti", "Juravushka" and "Kamelia" was more perceptible. Red blood cells formed stable structures and liquid was perfectly transparent. In 20 minutes in all dilutions reaction was still perceptible, but not sustained. Agglutinates did not break down into pieces and liquid was still clear. In 30 minutes after shaking it was observed that agglutinates disintegrated, liquid was muddy. That was the first sign that hemagglutination was very weak (figures 2 and 3).

In other cases reaction was moderate and agglutinates accumulated at the edges of holes. In one hour there was not any visible changing.

Further experiments of studying hemmagglutinative activity were performed by measurement of lectin titer, i.e. maximum dilution of minimum concentration in its solution in which red blood cells still can make agglutinates. As a result of these experiments it was proved that morphogenic calluses have high titer of lectin ( $27.7-30.8 \text{ [mg/ml]}^{-1}$ ). The highest activity was detected for "Aktatti" and "Juravushka". "RedGoya" and "Kamelia" demonstrated a little bit smaller activity than the previous ones. Non-morphogenic calluses had the smallest titer (not more than  $10-11 \text{ [mg/ml]}^{-1}$ ). Probably these distinguishes can depend on types and concentration of phytohormones, used for callus growth induction. Obtained data does not contradict with the results of similar experiments with wheat [20, 21].

As mentioned earlier, lectins take participation in different processes such as cell growth and differentiation. Therefore, regulation of lectin concentration in callus cultures might be under impact of various phytohormones. Concerning, the influence of auxins and cytokinins on induction of lectin formation in morphogenic legume "Juravushka" was studied. Asauxinsitwasused 2.4-D (1 mg/l), the role of cytokinins played kinetin (0-10 mg/l). As a result it was stated that the highest concentration of lectins is observed on a culture media containing 1 mg/l of 2.4-D and 0,5mg/lof kinetin (figure 4).

During the cell growth cycle (7 weeks) the concentration of lectins was determined by passaging 1.5 g of callus on a culture media containing 1 mg/l of 2.4 D and 0.1 mg/l of kinetin. This procedure was repeated every 10 days. The highest concentration was marked in its initial stage of growth. This can be explained by more intensive cell division, which is common for the phase (figure 5).

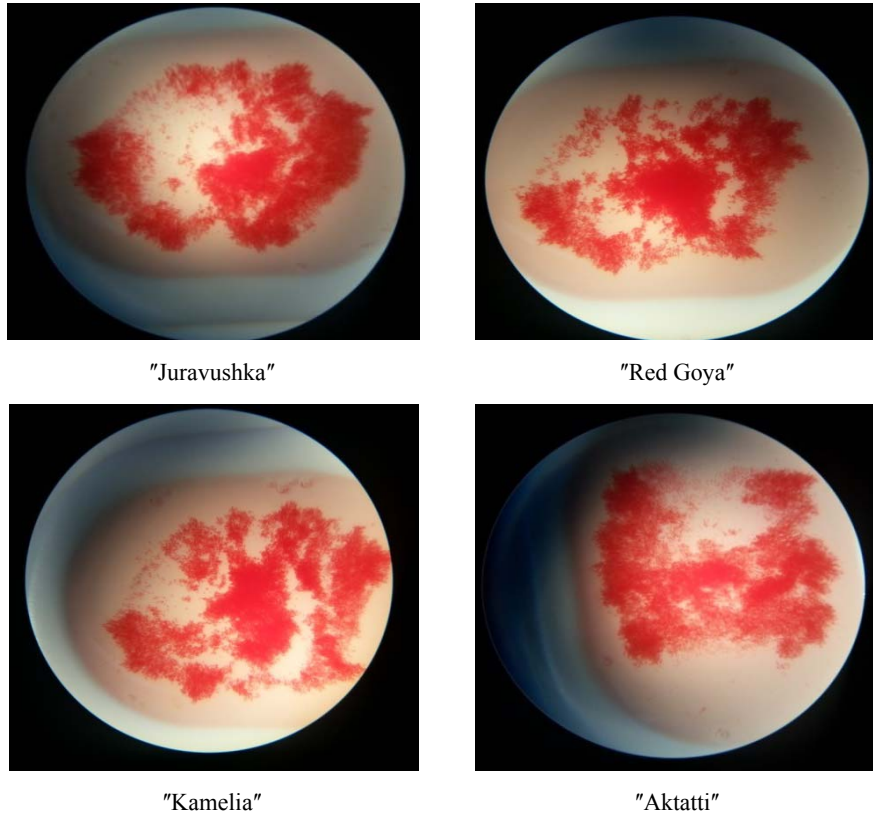


Figure 2 – Microphotograph of hemagglutination of morphogenic calluses (x20)

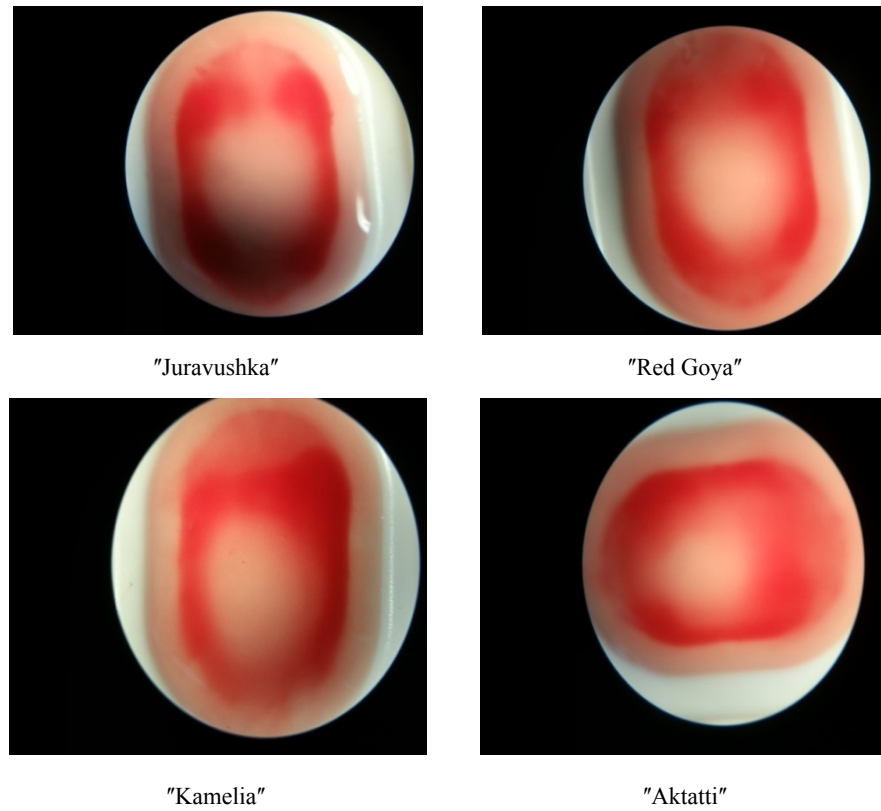


Figure 3 – Microphotograph of hemagglutination of non-morphogenic calluses (x20)



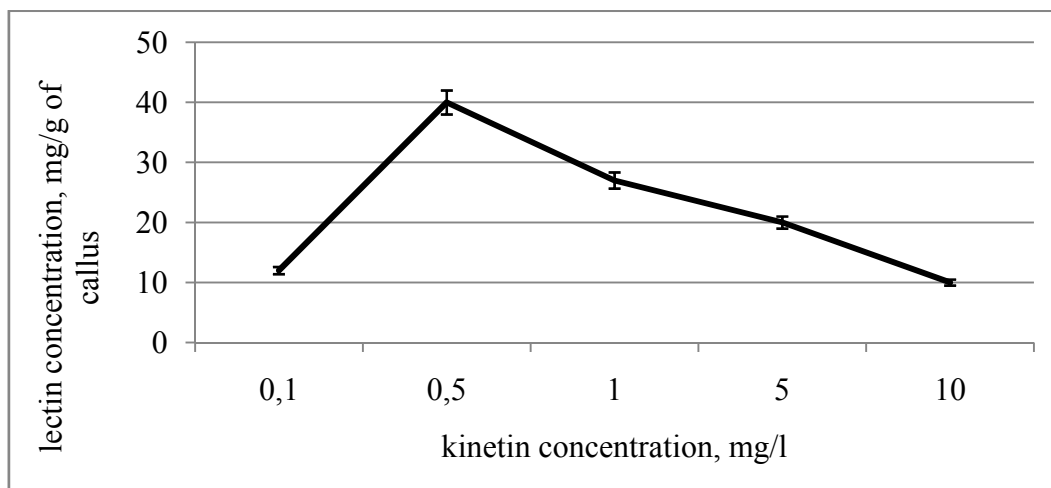


Figure 4 – Lectin concentration in callus tissue in dependence on dose of kinetin

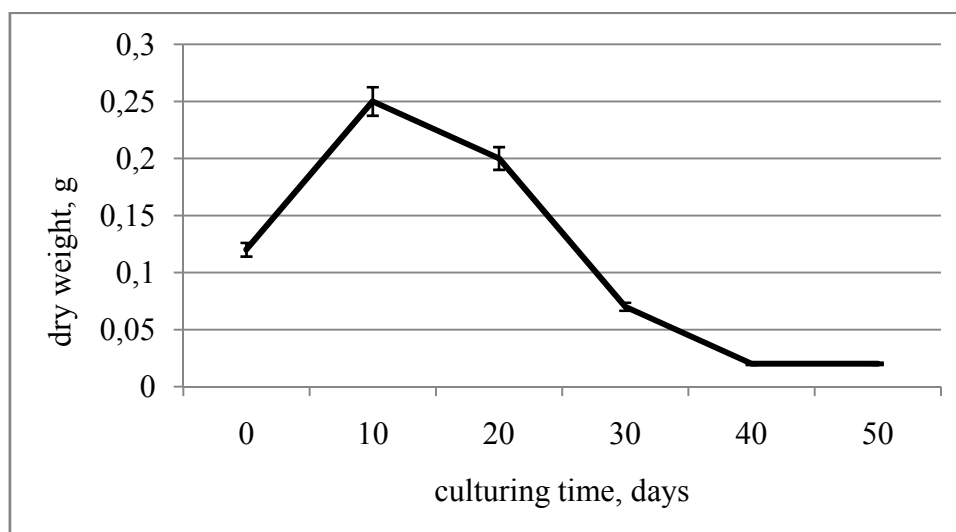


Figure 5 – Dependence between lectin concentration and growth cycle of calluses

Thus, this research on studying biological activity of lectins showed that accumulation, concentration and activity of lectins strictly depend on hormones in culture media. So, in the future calluses may be used as a biotechnological source to extract lectins for different purposes.

*This work has been carried out as part of the RK MES Project: 0825 / GF3: Study of the biological activity of the protein components of promising bean specimens and producing cell cultures with a high lectin content. (GR №: 0113RK00269).*

#### REFERENCES

- [1] Koval'chuk NV, Musatenko LI (2000) *Lektinipridozrivanninasinnjakvasoli 7:169-173 (In Ukrainian)*
- [2] Ed PJ Davies (2010) *Plant hormones: biosynthesis, signal transduction, action*. Springer, Netherlands. ISBN 978-1-4020-2686-7
- [3] Bogoeva VP, Radeva MA, Atanasova LY, Stoitsova SR, Boteva RN. (2004) Fluorescence analysis of hormone binding activities of wheat germ agglutinin, *BiochimBiophysActa*.6;1698(2):213-218DOI:10.1016/j.bbapap.2003.12.002
- [4] Oliveria BN, Stiirmer JC, Filcho JDS. (2008) Plant growth regulation activity of steviol and derivatives, *Phytochemistry*, 69:1528-1533DOI: 10.1016/j.phytochem.2008.01.015
- [5] Manoj K Rai, Shekhawat N.S, HarishAmit K, Gupta M Phulwaria, Kheta Ram and U. Jaiswal.(2011) The role of abscisic acid in plant tissue culture, *Plant cell, tissue and organ culture*, 106:2:179-190.DOI 10.1007/s11240-011-9923-9
- [6] Holf PLD, Brill LM, Hirsch AM. (2009) Plant lectins: the sites that bind in root symbiosis and plant defense, *Mol. Genet. Genomics*, 282:1-15DOI: 10.1007/s00438-009-0460-8

- [7] Borrebaeck CA, Linsefors L.(1985) Hormonal Regulation of the Lectin Biosynthesis in Callus Culture of the *Phaseolus vulgaris* plant, *Plant Physiol.* 79(3):659-662
- [8] Esteban R, Dopico B, Munoz FJ, Romo S, Labrador E.(2002) A seedling specific vegetative lectin gene is related to development in *Cicer arietinum*, *Physiol.Plant.* 114:619-626 DOI: 10.1034/j.1399-3054.2002.1140416.x
- [9] Zhao H. (2001) Interaction of alpha-agglutinin and a-agglutinin, *Saccharomyces cerevisiae* sexual cell adhesion molecules, *Bacteriol.* 183(9):2874-2880 DOI:10.1128/JB.183.9.2874-2880.2001
- [10] Zhumabaeva BA, Ajtasheva ZG, Dzhangalina ED, Zulpuhar Zh. (2016) Optimizacija uslovijvy delenija lektinov iz kallusnyh kul'tur fasoli // *Izvestija NANRK*, 1:29 – 34
- [11] Koval'chuk N.V. *Dinamika aktivnosti lektina pri prorastani isemjan fasoli // Ukr. biohim. zhurn.* - 2006. - T. 78 № 1. - S. 130-134
- [12] Bibikova A.V., Gorpenchenko T.Ju., Zhuravljov Ju.N. *Rastenija kak objekt biotekhnologii // Komarovskie chtenija.* - 2007. - Vyp. 55. - S. 184-211
- [13] Gatica AM, Jenny MV, Pilar RF, Marta VM. (2010) *In vitro* plant regeneration system for common bean (*Phaseolus vulgaris*): effect of N6-benzylaminopurine and adenine sulphate, *Electronic Journal of Biotechnology.* 13:1:2-8. DOI: 10.2225/vol13-issue1-fulltext-7
- [14] Veltcheva MR, Svetleva DL. (2005) *In vitro* regeneration of *Phaseolus vulgaris* L. via organogenesis from petioles explants, *Journal Central European Agriculture*, 6:1:53-58 DOI: <http://dx.doi.org/10.5513/jcea.v6i1.246>
- [15] Zhumabaeva B.A., Dzhangalina E.D., Ajtasheva Z.G., Zhigitbekova A.D. *Morfogeneticheskie osobennosti kallusnyh kul'tur fasoli obyknovnoy // Vestnik KazNU. Seriya Jekologicheskaja.* – 2012. - № 3(35). – S.113-117
- [16] Pavlovskaja N.E. *Belkovyj kompleks zernobobovyh kul'tur iputipovysheniya ego kachestva - Orel: OGAU, 2003.* – 216 s
- [17] Kol'man Ja., Rem G. (2000) *Nagljadnaja biohimija*, Mir, Russia. ISBN: 978-5-9963-0126-3, 978-5-03-003811-7.
- [18] Zadorin A.D. *Selekcija, semenovodstvo zernobobovyh ikrupjanyh kul'tur: sostojanie, problemy, perspektivy // Vestnik Semenovodstva.* – 2001. – №4. – S. 24–31;
- [19] Shajahmetov I. F. *Rol' lektin apshenicyabscizovoj kisloty v regeneracii rastenij // Uspehi sovremennoj biologii, 2004, tom 124, № 6, s. 602-611*
- [20] Wasilewska A., Vlad F., Sirichandra C. et al. (2008). An update on abscisic acid signaling in plants, *Molecular Plant*, 1:2:198-217 DOI: 10.1093/mp/ssp022.
- [21] Singh PS, Bhaglal P, Bhullar SS. (2000) Wheat germ agglutinin (WGA) gene expression and ABA accumulation in the developing embryos of wheat (*Triticum aestivum*) in response to drought, *Plant Growth Reg.* 30:145-150 DOI:10.1023/A:1006317302405

Э. Д. Джангалина, Б. А. Жумабаева, З. Г. Айташева, Л. П. Лебедева

Әл-Фараби атындағы Қазақ ұлттық университеті, Алматы, Қазақстан

#### ҮРМЕ БҮРШАҚ КАЛЛУСТЫҚ КУЛЬТУРАСЫНАН БӨЛІНІП АЛЫНҒАН ЛЕКТИНДІК БЕЛОҚТАРДЫҢ БИОЛОГИЯЛЫҚ АКТИВТІЛІГІ

**Аннотация.** Қазіргі уақытта *invitro* клетка культуралары мен ұлпалары биологиялық зерттеулер үшін кең қолданылуда. Көп зерттеулер табиғи биологиялық активті заттар мен түрлі спектрде әрекет ететін белоктық компоненттер алу мақсатында жүргізілуде. Биохимиялық құрамы бойынша бұршақ тұқымдастары басқаларға қарағанда белок және белоктық компоненттердің көп болуымен ерекшеленеді. Солардың бірі - лектиндік белоктар. *In vitro* клетка культураларында лектиндердің сандық құрамының гормональдық реттелуі сияқты әр түрлі жолдармен зерттеу арқылы - өсу процесін реттеу, бұршақ тұқымдастардың дамуы сияқты физиологиялық механизмдерді дәлелдеуге болады. Осындай жолмен эксплантты іріктеп алу үшін, қоректік ортаның минералды және гормональдық құрамын оңтайландыру мақсатында үрмебұршақтың морфогенді және морфогенді емес каллус ұлпалары алынды. 2,4-Д қолдану кезіндегі морфогендік каллустардың пайда болу жиілігі НСК қолданғандағымен салыстырғанда бес есе жоғары болды. Әр түрлі каллустардан бөлініп алынған лектиндердің биологиялық активтілігі гемагглютинация реакциясының қарқындылығын сырт көзбен бақылау және лектиндерді титрлеп өлшеу жолдарымен анықталды. Бізге белгілі үрмебұршақтың әрбір сорт үлгілеріндегі лектиндік активтілік қасиеті каллус ұлпасының түрі мен генотипіне байланысты. Ал ол, өз кезегінде қоректік ортаның гормональды құрамымен анықталды. Лектиндік белоктардың құрамы қоректік ортадағы ауксин мен цитокининнің концентрациясымен реттеліп отырады. Морфогендік түрлі каллустардың барлық үлгілерінің активтілігі морфогендік емес каллустармен салыстырғанда айтарлықтай жоғары болды. Ал лектиндердің максималды (ең көп мөлшері) жинақталуы - 2,4-Д – 1 мг/л, кинетин – 0,5 мг/л концентрацияда байқалды. Ең жоғары лектиндік активтілік «Актатти» және «Журавушка» атты морфогенді типті каллус үлгілерінде анықталған.

Жүргізілген зерттеулер нәтижелері көрсеткендей, белгілі фитогемаглютеиндеуші активтілікті және өсімдік каллус культураларындағы лектиндердің құрамын анықтау жұмыстары лектиндік активтіліктің түрленгіштік деңгейін клеткалық және ұлпалық деңгейде анықтап, табуға мүмкіндік береді және дифференциация

ция процесіндегі гормональдық реттелуді жете зерттеу үшін келешегі бар тәсіл болып табылады. Сондай ақ, бұл жұмыстар бұршақ тұқымдастардың өсіп өнуі мен дамуын реттеуге және әр түрлі биотехнологиялық жолмен лектиндерді алу тәсілдерін жетілдіруге мүмкіндіктер береді.

**Түйін сөздер:** *Phaseolus vulgaris*, лектиндер, каллустық культура, белсенділік, гормонға тәуелділік.

Э. Д. Джангалина, Б. А. Жумабаева, З. Г. Айташева, Л. П. Лебедева

Казахский национальный университет им. аль-Фараби,  
Институт проблем биологии и биотехнологии МОН РК, Алматы, Казахстан

### **БИОЛОГИЧЕСКАЯ АКТИВНОСТЬ ЛЕКТИНОВ, ВЫДЕЛЕННЫХ ИЗ КАЛЛУСНЫХ КУЛЬТУР ФАСОЛИ**

**Аннотация.** Культура клеток и тканей *in vitro* в настоящее время находит применение в широком диапазоне биологических исследований, в частности получения природных БАВ и белковых компонентов различного спектра действия и применения. По биохимическому составу бобовые культуры отличаются высоким содержанием белков и белковых компонентов, к которым относятся, например, лектины. Путем изучения различных путей гормональной регуляции количественного содержания лектинов в культуре *in vitro* возможно установление физиологических механизмов регуляции роста и развития бобовых культур. В связи с этим путем подбора источника экспланта, оптимизации минерального и гормонального состава питательных сред, получены морфогенные и неморфогенные каллусные ткани фасоли. Частота образования морфогенных каллусов при использовании 2,4-Д была в пять раз выше, по сравнению с НУК. Биологическая активность лектинов, выделенных из различных типов каллусов определялась визуально по интенсивности реакции гемагглютинации и путем измерения титра лектинов. Установлено, что лектиновая активность сортообразцов фасоли зависит от генотипа и типа каллусной ткани, который определяется гормональным составом питательной среды. Содержание лектинов регулируется концентрацией ауксинов и цитокининов в питательной среде. У всех образцов активность морфогенного типа каллусов была значительно выше по сравнению с неморфогенным, а максимальное накопление лектинов наблюдалось при концентрации 2,4-Д – 1 мг/л, кинетин – 0,5 мг/л. Наибольшая лектиновая активность отмечена для морфогенного типа каллуса сортообразцов «Актатти» и «Журавушка».

Проведенные исследования показывают, что определение фитогемагглютинирующей активности и содержания лектинов в каллусных культурах растений позволяет установить варибельность уровня лектиновой активности на клеточном и тканевом уровнях и может служить перспективным подходом для изучения гормональной регуляции процессов дифференциации, роста и развития бобовых, а также усовершенствовать биотехнологические подходы получения лектинов.

**Ключевые слова:** *Phaseolus vulgaris*, лектины, каллусные культуры, активность, гормональная зависимость.

## **Publication Ethics and Publication Malpractice in the journals of the National Academy of Sciences of the Republic of Kazakhstan**

For information on Ethics in publishing and Ethical guidelines for journal publication see <http://www.elsevier.com/publishingethics> and <http://www.elsevier.com/journal-authors/ethics>.

Submission of an article to the National Academy of Sciences of the Republic of Kazakhstan implies that the described work has not been published previously (except in the form of an abstract or as part of a published lecture or academic thesis or as an electronic preprint, see <http://www.elsevier.com/postingpolicy>), that it is not under consideration for publication elsewhere, that its publication is approved by all authors and tacitly or explicitly by the responsible authorities where the work was carried out, and that, if accepted, it will not be published elsewhere in the same form, in English or in any other language, including electronically without the written consent of the copyright-holder. In particular, translations into English of papers already published in another language are not accepted.

No other forms of scientific misconduct are allowed, such as plagiarism, falsification, fraudulent data, incorrect interpretation of other works, incorrect citations, etc. The National Academy of Sciences of the Republic of Kazakhstan follows the Code of Conduct of the Committee on Publication Ethics (COPE), and follows the COPE Flowcharts for Resolving Cases of Suspected Misconduct ([http://publicationethics.org/files/u2/New\\_Code.pdf](http://publicationethics.org/files/u2/New_Code.pdf)). To verify originality, your article may be checked by the Cross Check originality detection service <http://www.elsevier.com/editors/plagdetect>.

The authors are obliged to participate in peer review process and be ready to provide corrections, clarifications, retractions and apologies when needed. All authors of a paper should have significantly contributed to the research.

The reviewers should provide objective judgments and should point out relevant published works which are not yet cited. Reviewed articles should be treated confidentially. The reviewers will be chosen in such a way that there is no conflict of interests with respect to the research, the authors and/or the research funders.

The editors have complete responsibility and authority to reject or accept a paper, and they will only accept a paper when reasonably certain. They will preserve anonymity of reviewers and promote publication of corrections, clarifications, retractions and apologies when needed. The acceptance of a paper automatically implies the copyright transfer to the National Academy of Sciences of the Republic of Kazakhstan.

The Editorial Board of the National Academy of Sciences of the Republic of Kazakhstan will monitor and safeguard publishing ethics.

Правила оформления статьи для публикации в журнале смотреть на сайте:

[www.nauka-nanrk.kz](http://www.nauka-nanrk.kz)

**ISSN 2518-1629 (Online), ISSN 2224-5308 (Print)**

<http://www.biological-medical.kz/index.php/ru/>

Редактор *М. С. Ахметова, Д. С. Аленов, Т. М. Апендиев*  
Верстка на компьютере *Д. Н. Калкабековой*

Подписано в печать 06.04.2017.  
Формат 60x881/8. Бумага офсетная. Печать – ризограф.  
16,4 п.л. Тираж 300. Заказ 2.