

5. Петренко В.А., Татков С.И. Структурно-функциональная организация лейкоцитарного интерферона // Молекулярная биология. – 1990. – 24, Вып.6. – С. 1495-1504.
6. Sambrook J., Russell D. Molecular Cloning: A Laboratory Manual. – Cold Spring Harbour Laboratory Press, 2001. – P.2344.
7. Laemmli U.K. Cleavage of structural proteins during the assembly of the head of bacteriophage – T4. // Nature. – 1970. – 227. – P.680-685.
8. pET System Manual 11th edition. – Merck; Germany, 2004.
9. Baneyx Fr. Recombinant protein expression in *Escherichia coli*. // Current Opinion in Biotechnology – 1999. – 10. – P.411-421.

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CHOLESTEROL-LOWERING ACTIVITY OF LACTIC ACID BACTERIA PROBIOTIC STRAINS *IN VIVO*

Cholesterol-lowering activity of probiotic strains of lactic acid bacteria genera Lactobacillus and Bifidobacterium in the in vivo experiments on the model of experimental hypercholesterolemia in mice was studied. It is established that the prophylactic scheme of introduction of probiotic cultures is more effective than therapeutic one for the manifestation of cholesterolase activity of probiotic cultures. The most effective were the cultures: L. acidophilus and B. bifidum, as well as the composition B. bifidum + B. longum. Cholesterol-lowering activity of the studied strains and their compositions in this experiment ranged between 40-78%. It is noted that cholesterolase activity of other studied strains was not lower, and in some cases, higher than that of most of the drugs currently used in cholesterinosis.

К е у в о р д с: probiotic lactic acid bacteria, cholesterol-lowering activity, cholesterolase activity, cholesterol-assimilating strains, cholesterinosis.

In recent years, the number of reports on the ability of lactic acid bacteria to lower serum cholesterol levels is increasing in the scientific literature. The ability of certain strains of normal microflora to assimilate and deconjugate precipitate bile acids as well as to destroy, bind and assimilate cholesterol is the basis of their cholesterol-lowering effect (the ability to reduce cholesterol levels) [5; 4; 7-9]. High cholesterol levels in the serum as a whole and in the low-density lipoprotein is a major risk factor for coronary heart disease and atherosclerosis, and cerebrovascular atherosclerosis, hypertension, cancer of different parts of the digestive tract and some other pathologic conditions [2]. Thus, the purpose of the study was to establish the cholesterol-lowering activity of the previously selected strains of probiotic *Lactobacillus* and *Bifidobacterium* genera in the *in vivo* experiments on the model of experimental hypercholesterolemia in mice.

Materials and Methods. Probiotic strains of genera *Lactobacillus* and *Bifidobacterium* isolated from associative culture in laboratory studies of fermented biological material: *Bifidobacterium bifidum* VK-1, *Bifidobacterium longum* VK-2, *Lactobacillus acidophilus* IMV B-7279, *Lactobacillus casei* IMV B-7280, *Lactobacillus bulgaricus* IMV B-7281 were used as subjects of the study.

The authors used in the experiments white mice weighing 16-18 and 18-20 g, male mice of the *Balb/c* aged 2.5 months and female mice *Balb/c* aged 3 months. Experimental hypercholesterolemia was simulated in mice by feeding the animals with high-calorie diet (Table 1) for a week. Crystalline cholesterol with chemical purity of > 99% (Sigma, USA) was added to the diet. This model allows raising the serum cholesterol levels in mice by $46.54 \pm 2.1\%$ at an average as compared with intact mice.

Two schemes of administration of the probiotic strains – the prophylactic and therapeutic ones

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were worked out in the study. According to the prophylactic scheme mice of the experimental group received per os 0.3 ml of freshly prepared suspensions of the freeze-dried probiotic cultures, their combinations in concentrations of 3×10^8 cells/ml, and mixed fodder during 4 days. On the fifth day the mice received high-calorie diet and continued to receive probiotic cultures every day until the end of the diet (seven days). On the first, third and seventh day since the beginning of high-calorie diet the level of total serum cholesterol in animals was determined [1]. Cholesterol-lowering activity (cholesteraze activity) calculated by a decrease of concentration of serum cholesterol in mice which received high-calorie diet and probiotic cultures or their combinations in comparison with the control group of mice, which received only high-calorie diet. Cholesterol-lowering activity was evaluated in per cents from the control group of mice.

Table 1

Composition of diet based on corn meal to feed mice

Components	Quantity, g
Corn meal	153.5
Butter	27.2
Wheat bran	72.6
Soybean meal	98.0
Kitchen salt	0.9
CaHPO ₄	5.4
CaCO ₃	3.3
Vitamins, macro- and microelements*	1.5

Comments: The diet consists of the following vitamins, macro- and microelements: riboflavin – 1.76 µg; pantothenic acid – 8.80 µg; niacin – 8.80 µg; vitamin B₁₂ – 8.80 µg; choline chloride – 176.00 µg; vitamin A – 1760 IU; vitamin D₃ – 176 IU; vitamin E – 4.4 IU; and also complex of macro- and microelements: selenium – 39.6 µg, iodine – 300 µg; iron – 19.8 µg; manganese – 11 µg, copper – 2.2 µg, zinc – 39.6 µg per 1 kg of feed.

The therapeutic scheme provided co-administration of high-calorie diet and probiotic cultures in the diet of mice in the same doses as in the prophylactic scheme. The blood samples were also taken from the animals to analyze the level of total cholesterol on the first, third and seventh day of the experiment, respectively.

Two control groups of mice were used: the first (control) group included the intact mice, which diet contained only the standard feed, the second one (control + diet) included mice which diet included only high-calorie products with no addition of probiotic cultures.

Results and Discussion. In the previous experiments it was shown that all the studied strains of lactic acid bacteria were probiotic with high resistance to aggressive conditions of the gastrointestinal tract [3].

The previous experiments have proved that the selected strains, as well as compositions based on them, have high cholesterol-lowering activity *in vitro* [11].

The cholesteraze activity of bacteria genera *Lactobacillus* and *Bifidobacterium* was determined in the study in the experimental model of hypercholesterolemia in mice. The results are shown in the figures (Fig. 1-6).

The data presented in Fig. 1. show that on the first day of using the therapeutic scheme introduction of probiotic cultures into mice weighing 18-20 g maximum of cholesteraze activity was observed for *L. acidophilus* – $31.15 \pm 1.4\%$, while the minimum for compositions of *L. acidophilus* + *L. casei* – $3.0 \pm 0.1\%$, and *L. casei* + *L. bulgaricus* – $4.32 \pm 0.2\%$. At the same time for other cultures cholesteraze activity ranged to 8.44-16.07%. On the 3rd day of observation maximum cholesteraze activity remained $37.11 \pm 1.6\%$ for *L. acidophilus*, the minimum value was $17.84 \pm 0.7\%$ for *L. casei*. The maximum value of cholesteraze activity on the 7th day of observation was shown by the culture *L. casei* as $62.28 \pm 2.5\%$, minimum – by the composition *L. acidophilus* + *L. casei* as $28.70 \pm 1.2\%$, respectively. For other cultures cholesteraze activity was almost the same and varied within $43.01 \pm 1.7\%$.

Fig. 2 shows cholesteraze activity for mice weighing 16-18 g under administration of probiotic bacteria according to the therapeutic scheme. The maximum values of cholesteraze activity were shown by *L. acidophilus* + *L. casei* as $43.38 \pm 1.5\%$, and by *B. bifidum* + *B. longum* as $64.78 \pm 2.7\%$

on the 1st, 3rd and 7th days, respectively. On the first day minimum values were $16.66 \pm 0.7\%$ for *L. casei* and $16.63 \pm 0.6\%$ for *L. casei* + *L. bulgaricus*, on the third and seventh day they were 35.19 ± 1.3 and $48.16 \pm 1.8\%$, respectively for *L. casei* + *L. bulgaricus*. On the seventh day of observation the average value of cholesteraze activity was $33.0 \pm 2.3\%$, which is by $13.32 \pm 0.5\%$ more than for mice weighing 18-20g. The obtained data suggest that it is easier to restore the organisms of young mice under the therapeutic administration of probiotic cultures, than the organisms of more mature mice.

Fig. 3 and 4 demonstrated the cholesteraze activity of probiotic cultures in male mice *Balb/c* aged 2.5 months in prophylactic and therapeutic schemes of probiotic cultures administration, respectively.

For prophylactic scheme of administration of probiotic cultures (Fig. 3) cholesteraze activity for all strains and their compositions were characterized by practically the same values and amounted to $33.63 \pm 1.4\%$ on the 1st day, $45.41 \pm 1.6\%$ on the 3rd day, and $65.29 \pm 2.6\%$ on the 7th day, respectively. The maximum value of cholesteraze activity on the 7th day of observation was characteristic of the culture *L. acidophilus*, it was $69.58 \pm 2.8\%$. *L. acidophilus* and culture composition of *B. bifidum* + *B. longum* were the most effective for administration by the prophylactic scheme. The same trend remained for the therapeutic scheme (Fig.4). On the first day of the experiment the average cholesteraze activity ranged from 8.68% for *L. casei*, to $19.87-20.98\%$ for *L. acidophilus*, *B. bifidum* + *B. longum*, *L. casei* + *L. bulgaricus*. The average value of cholesteraze activity on the third day of the therapeutic scheme was $35.11 \pm 1.4\%$. By the 7th day cholesteraze activity increased minimum to 56.5% for *L. casei* and *L. casei* + *L. bulgaricus*, maximum to $65.84 \pm 2.6\%$ for *B. bifidum* + *B. longum*.

Fig. 5 and 6 showed the data for cholesteraze activity of female mice *Balb/c* aged 3 months when using the prophylactic and therapeutic schemes of administering the probiotic bacteria, respectively. In this case, the prophylactic scheme administration also showed higher values of cholesteraze activity than the therapeutic one. For prophylactic scheme administration (Fig. 5) the cholesteraze activity amounted to $37.01 \pm 1.4\%$ on the 1st day, $57.11 \pm 2.3\%$ on the 3rd day, $68.37 \pm 3.0\%$ on the 7th day. The maximum of cholesteraze activity on the 7th day of observation was shown by the culture *L. acidophilus* as $78.04 \pm 3.0\%$ and by the composition *B. bifidum* + *B. longum* as $74.08 \pm 3.0\%$.

The therapeutic scheme (Fig. 6) was characterized by slightly lower average values of cholesteraze activity: $26.63 \pm 1.1\%$ for the 1st day, $38.57 \pm 1.5\%$ for the third day, and $58.82 \pm 2.4\%$ for the 7th day. The maximum value of activity was found for the culture *L. acidophilus* as $69.59 \pm 2.8\%$, for other cultures it varied between $51.75 - 60.03\%$ on the 7th day of observation.

Recent researches in this field conformed completely to these experimental data [10].

As can be seen from all figures, regardless of breed, age, sex, body weight of mice and administration scheme of probiotic cultures their cholesteraze activity increases to the seventh days of observation. It should also be noted that the prophylactic schemes of administration of probiotic cultures had higher values of cholesteraze of bacteria than therapeutic ones. This suggests that the prevention of disease is the best treatment.

L. acidophilus and *B. bifidum*, as well as composition *B. bifidum* + *B. longum* were the most effective cultures used for treatment of mice with hypercholesterolemia. At the same time, it should be noted that the cholesteraze activity of the other studied strains was not lower, and in some cases even higher than that of most of the drugs currently used in cholesterinosis, for example Lovastatin, Fluvastatin, Atorvastatin and others [6].

Cholesterol-lowering activity of the studied strains and their compositions in the experiment ranged between 40-78%. In the future it is planned to increase the percentage of cholesteraze activity by more detailed working out of administration schemes and doses of cultures, as well as the selection of combinations and ratios of strains in these combinations.

Thus, the selected cultures of lactic acid bacteria could potentially be used to create on their basis new probiotics to reduce serum cholesterol in humans. Probiotics that contain cholesterol-assimilating strains of lactic acid bacteria can efficiently complete the complex therapy of patients with cardiovascular, cancer and other diseases.

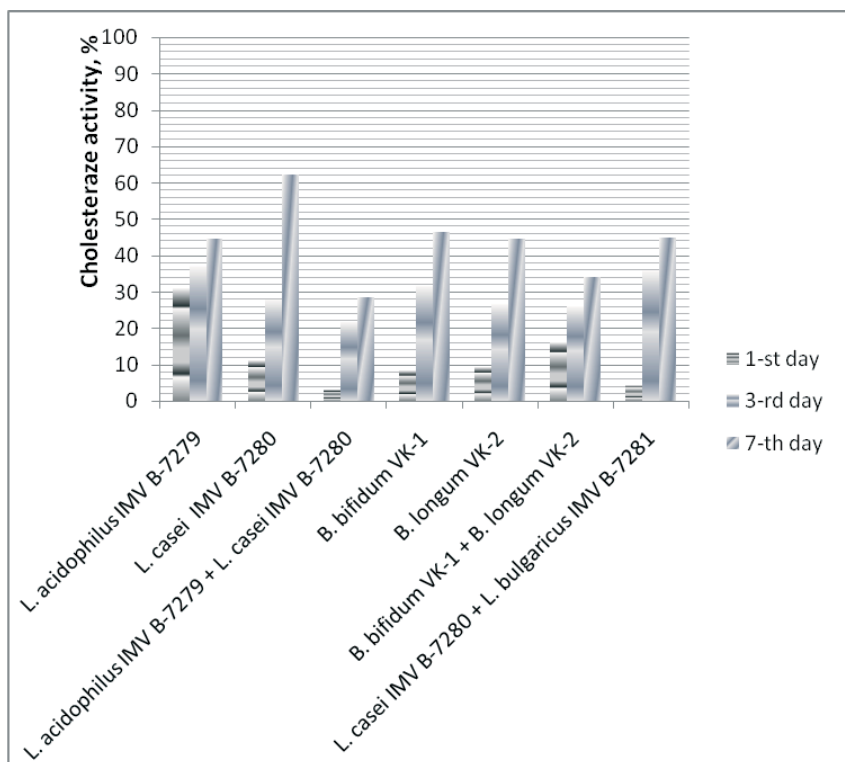


Fig.1. Dependence of cholesterolase activity of lactic acid bacteria and their compositions on observation time under therapeutic scheme administration of probiotic cultures ($P < 0.05$).

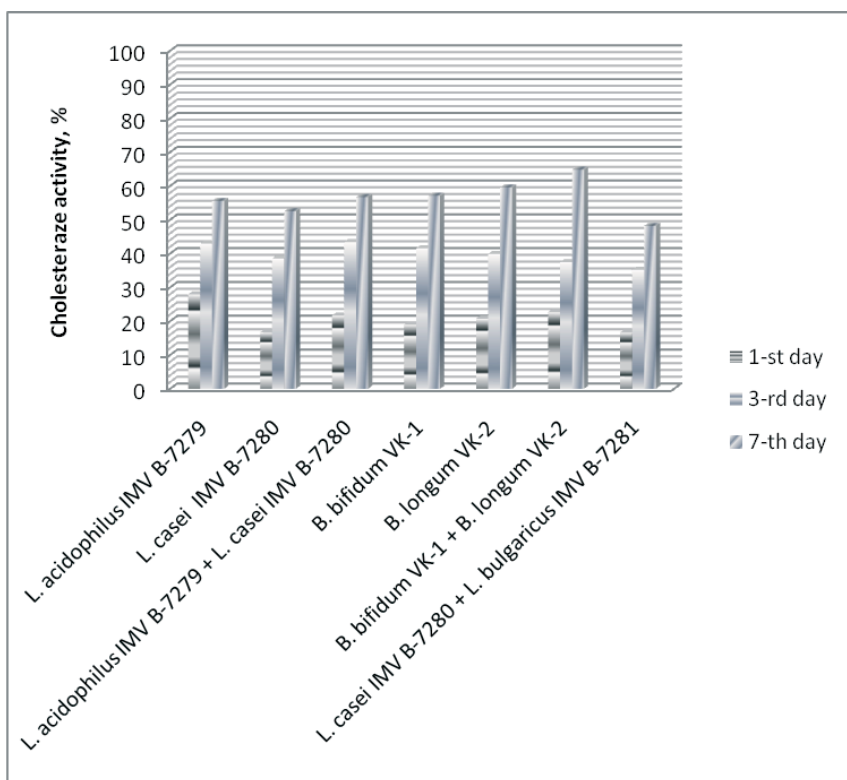


Fig.2. Dependence of cholesterolase activity of lactic acid bacteria and their compositions on observation time for mice 16-18 g under the therapeutic scheme administration of probiotic cultures ($P < 0.05$).

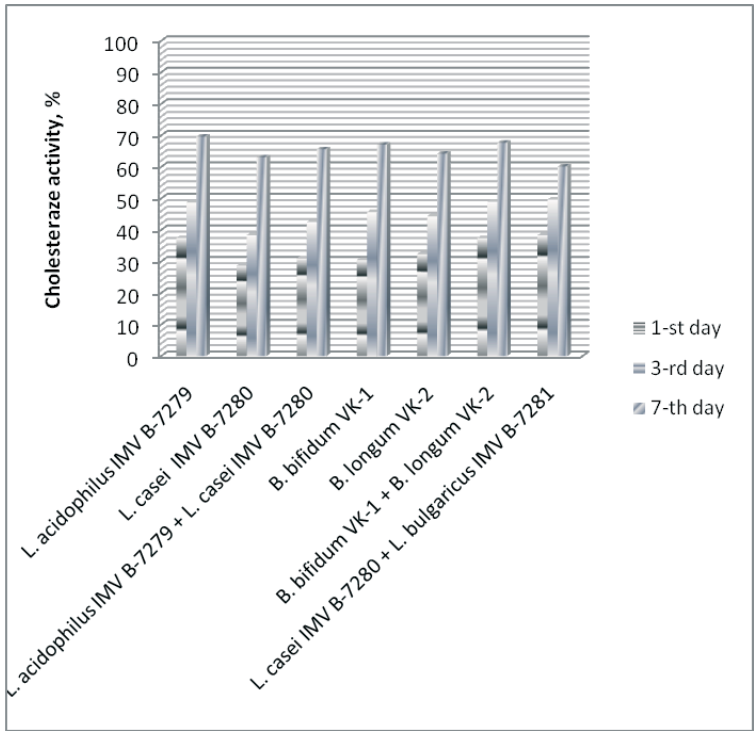


Fig.3. Dependence of cholesterolase activity of lactic acid bacteria and their compositions on the days of observation for male mice *Balb/c* aged 2.5 months when using the prophylactic scheme administration of probiotic cultures ($P < 0.05$).

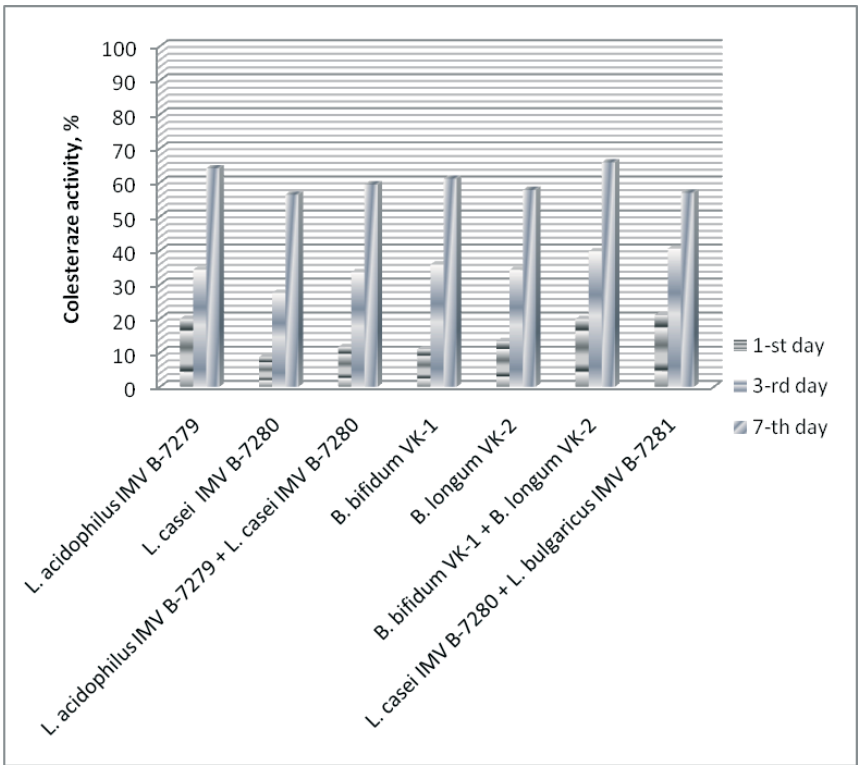


Fig.4. Dependence of cholesterolase activity of lactic acid bacteria and their compositions on the days of observation for male mice *Balb/c* aged 2.5 months when using the therapeutic scheme administration of probiotic cultures ($P < 0.05$).

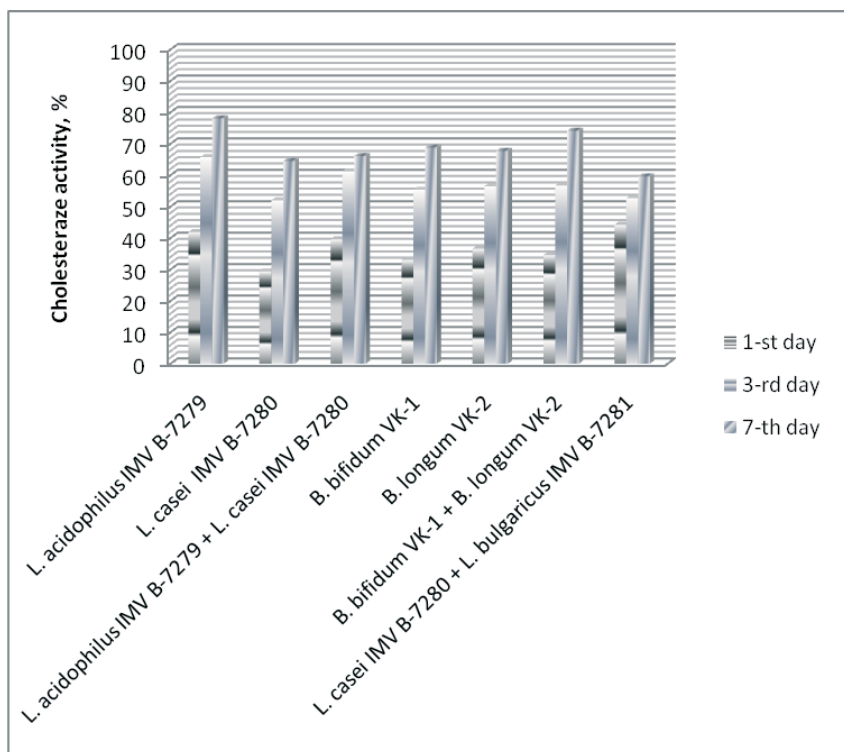


Fig.5. Dependence of cholesterolase activity of lactic acid bacteria and their compositions on the days of observation for female mice *Balb/c* aged 3 months when using the prophylactic scheme administration of probiotic cultures ($P < 0.05$).

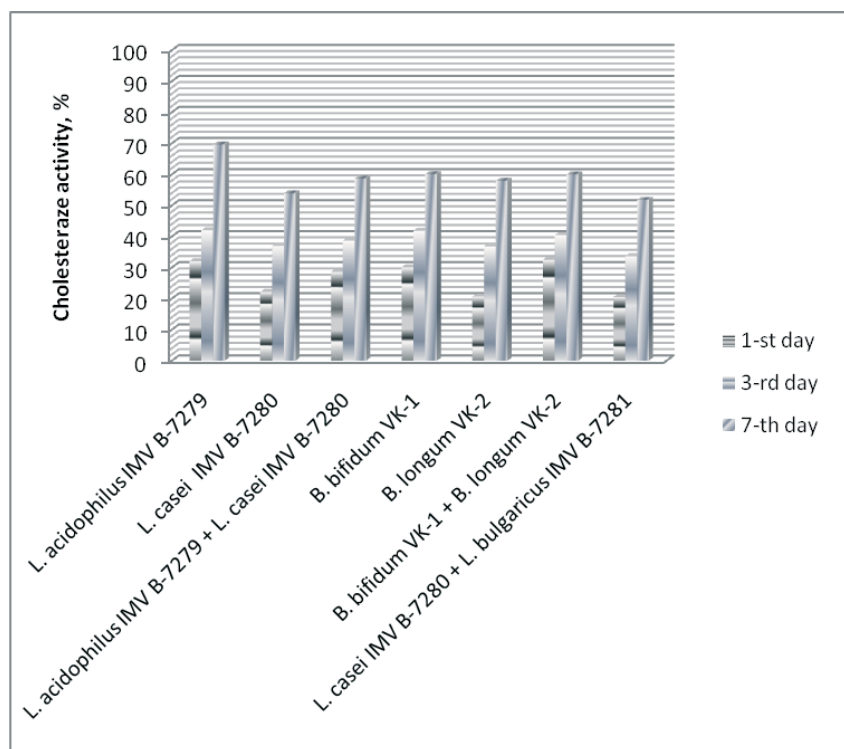


Fig.6. Dependence of cholesterolase activity of lactic acid bacteria and their compositions on the days of observation for female mice *Balb/c* aged 3 months when using the therapeutic scheme administration of probiotic cultures ($P < 0.05$).

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ГИПОХОЛЕСТЕРИНЕМИЧЕСКАЯ АКТИВНОСТЬ ПРОБИОТИЧЕСКИХ ШТАММОВ МОЛОЧНОКИСЛЫХ БАКТЕРИЙ *IN VIVO*

Резюме

Изучена гипохолестеринемическая активность пробиотических штаммов молочнокислых бактерий родов *Lactobacillus* и *Bifidobacterium* в опытах *in vivo* на модели экспериментальной гиперхолестеринемии у мышей. Установлено, что профилактическая схема введения пробиотических культур является более эффективной, чем лечебная, для проявления холестеразной активности культурами. Наиболее эффективными оказались культуры: *L. acidophilus* и *B. bifidum*, а также композиция *B. bifidum* + *B. longum*. Отмечено, что холестеразная активность остальных изученных штаммов была не ниже, а в некоторых случаях и выше, чем большинства лекарственных препаратов применяемых в настоящее время при холестеринозе.

Ключевые слова: пробиотик, молочнокислые бактерии, гипохолестеринемическая активность, холестерин-ассимилирующие штаммы, холестериноз.

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ГИПОХОЛЕСТЕРИНЕМИЧНА АКТИВНІСТЬ ПРОБІОТИЧНИХ ШТАМІВ МОЛОЧНОКИСЛИХ БАКТЕРІЙ *IN VIVO*

Резюме

Вивчена гіпохолестеринемічна активність пробіотичних штамів молочнокислих бактерій родів *Lactobacillus* та *Bifidobacterium* в досліді *in vivo* на моделі експериментальної гіперхолестеринемії у мишей. Встановлено, що профілактична схема введення пробіотичних культур є більш ефективною, ніж лікувальна, для прояву холестеразної активності культурами. Найбільш ефективними виявилися культури: *L. acidophilus* та *B. bifidum*, а також композиція *B. bifidum* + *B. longum*. Відмічено, що холестеразна активність інших вивчених штамів була не нижче, а в деяких випадках і вище, ніж у більшості лікарських препаратів, що застосовуються в теперішній час для лікування холестеринозу.

Ключові слова: пробіотик, молочнокислі бактерії, гіпохолестеринемічна активність, холестерин-асимілюючі штами, холестериноз.

1. Колб В.Г., Камышиников В.С. Клиническая биохимия (Пособие для врачей-лаборантов). – Минск: Беларусь, 1976. – 311 с.
2. Мосійчук С.М., Хоменко М.Б., Михайлова Т.С., Кігель Н.Ф., Карнов О.В. Пробиотики: можливість застосування при гіперхолестеринемії // Український медичний часопис. – 2006 – № 2 (52). – С. 10 – 23.
3. Старовойтова С.А., Лазаренко Л.Н., Авдеева Л.В., Полищук Е.И., Тимошок Н.А., Шинкаренко Л.Н., Бабенко Л.П., Мокрозуб В.В., Спивак Н.Я., Чейнеш А.В., Николайчук В.И. Поиск штаммов бактерий родов *Lactobacillus* и *Bifidobacterium* перспективных для создания пробиотиков // Науковий вісник Ужгородського університету. Серія Біологія. – 2009. – Вип. 26. – С. 216–219.
4. Brashears M.M., Gilliland S.E., Buck L.M. Bile salt deconjugation and cholesterol removal from media by *Lactobacillus casei* // J. Dairy Sci. – 1998. – **81**, №8. – P. 2103–2110.
5. Dambekodi P.C., Gilliland S.E. Incorporation of cholesterol into the cellular membrane of *Bifidobacterium longum* // J. Dairy Sci. – 1998. – **81**, N7. – P.1818–1824.
6. Erkelens D.W., Baggen M.G., Van Doormaal J.J. et al. Clinical experience with simvastatin compared with cholestyramine // Drugs. – 1988. – **39**, Suppl. 3. – P. 87–92.

7. Kimoto H., Ohmoto S., Okamoto T. Cholesterol removal from media by *Lactococci* //J. Dairy Sci. – 2002. – **85**, N 12. – P. 3182–3188.
8. Noh D.O., Kim S.H., Gilliland S.E. Incorporation of cholesterol into the cellular membrane of *Lactobacillus acidophilus* ATCC 43121 //J. Dairy Sci. – 1997. – **80**, N12. – P. 3107–3113.
9. Psomas E.I., Fletouris D.J., Litopoulou-Tzanetaki E., Tzanetakis N. Assimilation of cholesterol by yeast strains isolated from infant feces and Feta cheese //J. Dairy Sci. – 2003. – **86**, N 11. – P. 3416–3422.
10. Sirilun S., Chaiyasut C., Kantachote D., Luxanani P. Characterization of non human origin probiotic *Lactobacillus plantarum* with cholesterol-lowering property //African Journal of Microbiology Research. – 2010. – **4**, N 10. – P. 994–1000.
11. Starovoitova S., Kishko K., Lazarenko L., Shynkarenko L., Spivak M., Nikolaychuk M. Cholesteraze activity of new lacto- and bifidobacteria strains *in vitro* //Науковий вісник Ужгородського університету. Серія Біологія. – 2010. – Вип. 27. – С. 1–4.

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БИОЛОГИЧЕСКАЯ АКТИВНОСТЬ ЛИПОПОЛИСАХАРИДОВ *BUDVICIA AQUATICA*

*Впервые изучен жирнокислотный состав липидов А липополисахаридов (ЛПС) 6 штаммов *Budvicia aquatica* – представителей нового вида *Enterobacteriaceae*. Установлено присутствие жирных кислот с длиной углеродной цепи от C₁₂ до C₁₈. Все исследуемые штаммы *B. aquatica* содержали 3-гидрокситетрадекановую кислоту (23,1-43,8 %, в зависимости от штамма), которая была доминирующей и характерна для семейства *Enterobacteriaceae*. ЛПС исследуемых штаммов проявляли токсичность и пирогенность.*

Ключевые слова: *Budvicia aquatica*, липополисахарид, липид А, жирнокислотный состав, токсичность, пирогенность.

Важным фактором патогенности грамотрицательных бактерий является липополисахарид (ЛПС) – специфический гликополимер, который входит в состав наружной мембраны клеточной оболочки грамотрицательных бактерий и вместе с белками формирует ее внешний слой. Находясь в тесном контакте с мембранными белками, ЛПС обеспечивают целостность, стабильность и функциональное предназначение наружной мембраны микробной клетки. В силу своего поверхностного расположения ЛПС играет важную роль во взаимоотношениях бактериальной клетки с окружающей средой, а в случае патогенных микроорганизмов – с организмом-хозяином, в отношении к которому он проявляет себя как О-антиген и эндотоксин. Многие патофизиологические проявления грамотрицательных инфекций, в том числе эндотоксемия и бактериальный шок, ассоциированы с уникальными эндотоксическими свойствами ЛПС. Среди широкого спектра биологической активности ЛПС особое внимание исследователей привлекают их токсичность и способность активировать клетки иммунной системы. Результатом специфического взаимодействия ЛПС с клетками макроорганизма является биосинтез активных медиаторов-цитокинов, которые при низкой концентрации регулируют работу иммунной системы организма, а при высокой – вызывают развитие сложной сети токсических эффектов, таких как пирогенность, лейкопения, септический шок.

Поскольку липид А ЛПС погружен во внешнюю мембрану клеток бактерий, вероятно, он проявляет свои токсические эффекты, когда освобождается из размножающихся клеток в растворимой форме или когда в результате аутолизиса, действия комплемента, фагоцитоза или определенного типа антибиотиков происходит лизис клеток.

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