образов, что приводит к существенному ухудшению качества жизни и результатов обучения.

Цель явилась разработка оптимального диагностического алгоритма для детей с ХГСО.

Материалы и методы исследования. Для достижения поставленной цели в отделение «врожденных и приобретенных заболеваний ЛОР органов» РСНПМЦПи в клинике ТашПМИ было обследовано 30 детей в возрасте от 3 до 17 лет, за период с 2017 по 2019. Мальчиков было 13, девочек – 17. Методика обследования включала: изучения анамнеза и жалоб больного, общий оториноларингологический осмотр без/с эндоскопией, отомикроскопия, слуховые тесты, изучения микрофлоры хронического воспалительного секрета, ОАЭ, тимпанометрия, тональная пороговая аудиометрия, МСКТ височной кости.

Результаты. Давность болезни составляла примерно 1-10лет. Наиболее частой жалобой больных и их родителей было гноетечения из уха у 23

детей (76,86%), запах из уха у 17(56,6%), снижение слуха у 20(66,7%) и головная боль у 14(46,7%). При аудиологическом исследовании характер аудиологической кривой практически у всех больных носил смешанный характер с различным костно-воздушным интервалом. Интраоперационно во всех случаях было выявлено, реккурантная холеастеатома у 25 (83%) больных, грануляции и полипы у 23 (77%). Катамнестические данные за оперированными ушами в течении 3 лет показало анатомические результаты у 93% больных, функционалные результаты у 73%.

Выводы. Больные ХГСО обращаются за медицинской помощью при наличии жалоб, воспалении или нагноении. Основным методом лечения больных ХГСО является хирургическое лечения, причем проведения одномоментное санирующего и реконструктивного операцию на ухе.

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THE FACTORS OF PATHOGENESIS OF NONALCOHOLIC STEATOHEPATITIS IN PATIENTS WITH CHRONIC KIDNEY DISEASE

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ФАКТОРЫ ПАТОГЕНЕЗА НЕАЛКОГОЛЬНОГО СТЕАТОГЕПАТИТА У БОЛЬНЫХ С ХРОНИЧЕСКОЙ БОЛЕЗНЬЮ ПОЧЕК

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ЧИННИКИ ПАТОГЕНЕЗУ НЕАЛКОГОЛЬНОГО СТЕАТОГЕПАТИТУ У ХВОРИХ НА ХРОНІЧНУ ХВОРОБУ НИРОК

Summary.

As a result of studies, it was found that patients with non-alcoholic steatohepatitis on the background of obesity are characterized by lipid distress syndrome with an increase in total cholesterol in blood, low density proatherogenic lipoprotein, deficiency of anti-atherogenic high-density lipoproteins. The role of nitrosative stress in the pathogenesis of nonalcoholic steatohepatitis, the confirmation of which is the increase in the concentration of nitrosothiols, peroxynitrite and other metabolites NO in the blood. Thus, with the comorbid flow of non-alcoholic steatohepatitis and chronic kidney disease of the I-II stage, on the background of obesity, a deeper lipid imbalance, hypercholesterolemia, including in the low density lipoprotein (1.8 times, p <0.05), a decrease in the content of high density lipoprotein (1.8 times, p <0.05), an increase in the atherogenicity index, and hyperproduction of the endothelium and lymphocytes from the progressive damage to the endothelium (an increase in the number of desquamated endothelial cells).

Резюме.

В статье приведены теоретическое обобщение результатов исследования липидного спектра крови и эндотелиальной дисфункции за коморбидности течения неалкогольного стеатогепатита с ожирением и хронической болезнью почек I-II стадии, сопровождается глубоким липидный дисбаланс, гиперхолестеролемия, в т.ч. в составе липопротеидов низкой плотности, снижение содержания липопротеидов высокой плотности, рост индекса атерогенности, а также гиперпродукция NO эндотелием и лимфоцитами с прогрессирующим повреждением эндотелия.

Резюме.

У статті наведено теоретичне узагальнення результатів дослідження ліпідного спектру крові та ендотеліальної дисфункції за коморбідного перебігу неалкогольного стеатогепатиту з ожирінням та

хронічною хворобою нирок I-II стадії, що супроводжується глибшим ліпідний дисбаланс, гіперхолестеролемія, у т.ч. у складі ліпопротеїдів низької густини, зниження вмісту ліпопротеїдів високої густини, зростання індексу атерогенності, а також гіперпродукція NO ендотелієм та лімфоцитами із прогресуючим ушкодженням ендотелію.

Introduction. The comorbidity of non-alcoholic steatohepatitis (NASH) and chronic kidney disease (CKD) on the background of obesity is often recently drawn to the attention of both practitioners and researchers [1, 2]. Schematically, the development of NASH can be presented in several stages: fatty infiltration of the liver, oxidative stress, mitochondrial dysfunction, TNF / endotoxin-mediated injury, aseptic inflammation, diffused liver fibrosis, development of liver-cellular insufficiency (LCI) [1,2,3]. The first place among the causes of the development of NASH is insulin resistance syndrome. NASH most often occurs in obesity (20-81%). The prevalence of NASH in the world is 10% (600 million people) [2,4,5]. In the last 5 years in Ukraine, the incidence of steatohepatitis has increased by 76.6%. In the 12-40% of patients with liver steatosis during 8-13 years, NASH develops with early liver fibrosis (LF).

The frequency of occurrence of NASH in patients with CKD is unknown. The mechanisms of their joint development are described in isolated works, which were conducted mainly in the experiments. Despite the fact that among various pathological processes in the internal organs that occur in the background of a metabolic syndrome - NASH is an extremely common disease, and quite often it occurs in patients with CKD, so far, this comorbidity remains a significant problem of the present and needs to be sufficiently studied.

The aim of the study – to find out the features of the lipid blood spectrum and the development of endothelial dysfunction in non-alcoholic steatohepatitis in patients with obesity of the 1st degree and chronic kidney disease of the 1st and 2nd stage.

Material and methods of research. 135 patients with non-alcoholic steatohepatitis (NASH) with comorbid obesity I degree and chronic kidney disease (CKD) of the 1st and 2nd stage, were examined. Patients were divided into 2 groups: of which 52 patients with non-alcoholic steatohepatitis with obesity I degree (group 1), 53 patients with non-alcoholic steatohepatitis with comorbid obesity of the 1st degree and chronic kidney disease of the I-II stage (chronic uncomplicated pyelonephritis with latent phase in subsiding exacerbation phase) (group 2). The control group consisted of 30 practically healthy persons of the corresponding age and sex. The average age of patients was (45.8 ± 3.81) years, men were 48, and women 57 persons.

The functional state of the endothelium was studied by the content of stable metabolites of nitrogen monoxide (NO) (nitrites, nitrates) In the blood by L.C.Green et al. The number of desquamated endothelial cells in the blood was determined by the method of J.Hladovec in the modification of N.N. Petrischev et al. The lipid blood spectrum was studied based on the content of common lipids (TL), total cholesterol, triacylglycerols (TG), low density lipoprotein (LDL) and high-density lipoprotein (HDL) (Danish Ltd, Lviv), and also calculated the index of atherogenicity (IA) by the formula: IA = total cholesterol / HDL. The degree of carbohydrate compensation was determined by the level of glycemia in the onset and 2 hours after glucose loading (glucose tolerance test) by the glucose oxidase method, the content of insulin in the blood (DRG System) - by the immunoassay (ELISA) method, the content of glycosylated hemoglobin (HbA1c) using standard sets of reagents (DanishLtd, Lviv) by the method of V.A. Koroleva. The degree of insulin resistance (IR) was determined by the size of the body mass index (BMI): body weight (kg) / height2 (m); The HOMA-IR index (D.R. Matthews et al.), which was calculated using the HOMA Calculator Version 2.2 Diabetes Trials Unit, University of Oxford (UK). The statistical analysis was performed using parametric and non-parametric criteria (Student, Pearson) on RS AMD Athlon 64 using Statistica 5.1 software (StatSoft, Inc., USA) and SPSS 10.0.5. Standard Version.

The diagnosis of NASH was established in accordance with the unified clinical protocol, approved by the order of the Ministry of Health of Ukraine No. 826 from 06.11.2014, in the presence of criteria for the exclusion of chronic diffuse liver disease of the viral, hereditary, autoimmune or medicinal genesis as causes of cholestatic or cytolytic syndromes, as well as the results of the USG survey. Diagnosis and treatment of CKD were performed according to the recommendations of the clinical guidelines of the State Institute "Institute of Nephrology, NAMS of Ukraine" (2012). The analysis of clinical manifestations of NASH and CKD of the 1st and 2nd stage, biochemical, laboratory parameters of the functional state of the liver, kidneys, endothelium, ultrasonographic data was studied in dynamics after 30, 90 days of treatment, as well as 3 months after treatment.

The statistical analysis of the results was carried out in accordance with the type of research carried out and the types of numerical data that were obtained. Distribution normality was verified using Liliefors, Shapiro-Uilka tests and the direct visual evaluation of eigenvalues distribution histograms. Quantitative indices having a normal distribution are represented as mean (M) ± standard deviation (S). Discrete values are

presented in the form of absolute and relative frequencies (percentage of observations to the total number of surveyed). For comparisons of data that had a normal distribution pattern, parametric tests were used to estimate the Student's t-criterion, Fisher's F-criterion. In the case of abnormal distribution, the median test, Mann-Whitney Rank U-Score, and Wilcox's T-criterion (in the case of dependent groups) were used for multiple comparison. Statistica for Windows version 8.0 (Stat Soft inc., USA), Microsoft Excel 2007 (Microsoft, USA) software packages were used for statistical and graphical analysis of the obtained results.

Results of the research and their discussion. Analysis of the lipid profile of the blood in patients with NASH and obesity showed a number of changes that differed depending on the presence of CKD (table 1). Indicators of concentration in blood of total lipids in patients of 1st and 2nd groups exceeded the norm by 26.4% and 34.2%, respectively, with a statistically significant difference between the groups ($p < 0.05$). The content of total cholesterol in blood indicated that it increased by 37.4 and 46.7 ($p < 0.05$) compared with PHPs in patients of 1st and 2nd groups. Changes in the concentration of TG in the form of a significant increase (respectively, 2.2 and 2.0 times ($p < 0.05$)) were recorded in the 1st and 2nd groups of patients.

Table 1

Indicators of lipid spectrum of blood, glycemia and endothelial dysfunction in patients with non-alcoholic stethohepatitis, obesity of the I-II degree and with comorbidity with chronic kidney disease of the I-II stage (M ± m)

Indicators, units measurement	Groups of patients surveyed		
	PHP	Group 1 NASH+Obesity	Group 2 NASH with CKD + Obesity
Total Cholesterol, mmol / 1	4,72±0,11	6,89±0,38*	6,93±0,39*/**
LDL, mmol / 1	2,54±0,02	4,05±0,022 *	4,58±0,04*/**
HDL, mmol / 1	1,28±0,05	0,72±0,02 *	0,76±0,04 */**
TG, mmol / 1	1,47±0,03	2,42±0,03*	3,19±0,07 */**
NO IN BLOOD, mmol / 1	15,32±1,225	30,49±1,318 *	40,51±1,173 */**
ET-1, pmol / 1	6,17±0,854	11,25±0,457 *	18,83±0,559 */**
DEC x104/L	3,03±0,204	3,87±0,123 *	5,80±0,127 */**
Notes: * - changes are probable in comparison with the index in PHP (P <0,05); ** - changes are probable when comparing the indices in patients with NASH (P <0.05);			

That is, the content in TG in the blood in the comorbid flow of NASH with CKD and obesity were significantly lower than in patients with NASH and obesity.

The study of blood concentrations of proatherogenic lipoprotein fractions indicated a number of changes: the concentration of LDL in the patients of the 1st group was 1.5 times higher than the control group ($p < 0.05$), and in patients of the 2nd group LDL increased in 1.7 times ($p < 0.05$). It is also necessary to point out that with the increase in the activity of cytolysis, the content of the cholesterol and LDL in the blood in NASH with comorbidity with CKD and obesity - increased, which may be an important prognostic factor in the progression of atherosclerosis in these patients. Concentration in blood of antiatherogenic lipoproteins - HDL in patients of both groups was significantly lower in comparison with control: in patients of the 1st group - in 1,5 times ($p < 0.05$), in 2nd group - 1,7 times ($p < 0.05$). As can be seen from the results of the study, the maximum suppression of HDL synthesis (Table 1) was observed in patients of the 2nd group, indicating a minimum level of protection of endothelial vessels from free radical aggression and atherogenic fractions of blood lipids. The result of these changes was a significant increase in the index of atherogenicity in patients of both groups of observation: the 1st group - 2.2 times, the 2nd group - 2.0 times with the maximum changes in the index in patients with NASH, CKD and obesity, which testifies on the one hand, the presence of significant risk factors for the progression of atherosclerosis in these patients on the background of obesity, and on the other - on the favorable pathogenetic situation with regard to the progress of NASH. Thus, the development of NASH in patients with CKD and obesity is accompanied by a significant disorder of dis-hyperlipidemia with the highest among groups comparing with the increase in the content of cholesterol and low-density proatherogenic lipoprotein, a possible decrease in high-density anti-atherogenic lipoprotein and an increase in the atherogenicity index.

In patients of 1st and 2nd groups, a slight increase in cardiac glycemia was observed at 9.3% and 14.8%, respectively ($p < 0.05$) compared with the control group. Analysis of indicators of postprandial glycemia, obtained during Gamma glutamyltransferase (GGT), in patients of 1st and 2nd groups also showed an increase in glucose content in 120 min after loading - respectively by 16.5% and 31.2% ($p < 0.05$) compared with indicators in the PHP group. Investigation of insulin content in blood on an empty stomach revealed hyperinsulinemia, which in patients of the 1st group exceeded the index in the PHP group by 1.9 times; in patients of the 2nd group insulin content exceeded the norm by 2.2 ($p < 0.05$) times .

The results of the study showed that in patients with NASH, a significant increase in the content of NO in the blood was detected in comparison with the index in PHP ($p < 0.05$) (Table 1) in group 1 - in 2,1 times, in the 2nd group - in 2,6 times ($p < 0.05$). The role of nitrosative stress in the pathogenesis of NASH was proved, the confirmation of which is the increase in the concentration of nitrosothiols, peroxynitrite and other

metabolites NO in the blood [2, 10]. Increased peroxynitrite formation due to the generation of NO by leukocytes is an important aspect of the damaging effect and inflammation process in NASH [3]. Pathological hyperproduction of NO by endothelial cells and leukocytes from inflammatory infiltrates in the liver contributes to the development of nitrosative stress in NASH. The established hypernitrate in blood may also be considered compensatory in response to hyperproduction of ET-1 in all observational groups. Thus, the content of ET-1 exceeded the index in PHP, respectively, in patients in the 1st group in 1.7 times, in the 2nd group - in 3.0 times (p1-2 <0.05). Confirmation of the presence of endothelial dysfunction (ED) in patients with NASH with CKD resulted in a probable growth of the number of desquamated endothelial cells (DEC) in the 2nd group of patients in 1.9 times (p2 <0.05). Generation by neutrophils during the exacerbation of NASH of a significant number of active forms of oxygen and nitrogen and hyperproduction of endothelial cells and endometrial lymphocytes with progressive damage to the endothelium (growth of DEC) leads to significant ED, accompanied by mosaic angiospasm of the arteries due to hyperproduction of ET-1 and parectic vasodilatation of the veins of the portal vein system because of the hyperproduction of NO.

Conclusions Thus, in patients with non-alcoholic steatohepatitis on the background of obesity lipid distress syndrome with an increase in total cholesterol in blood, low density proatherogenic lipoproteins, and a deficiency in anti-atherogenic high-density lipoproteins is characterized. In a comorbid flow of non-alcoholic steatohepatitis and chronic kidney disease stage I and II on the background of obesity a deeper lipid imbalance (hypertriacylglycerolemia (2.1 times, p <0.05), hypercholesterolemia (1.5 times, p <0.05), including in the low density lipoprotein (1.8 times, p <0.05), decrease in the content of high density lipoprotein (1.8 times, p <0.05), increase in the atherogenic index (at 2 , 7 times, p <0,05), as well as hyperproduction of the endothelium of the NO and lymphocytes with progressive damage to the endothelium (growth of DEC)) was confirmed.

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