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Research Article

COMPARISON OF BASELINE CHARACTERISTICS AMONG CASES AND CONTROLS TO ASSESS THE LEVELS OF VITAMIN D SERUM

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Abstract:

Background: According to an idea, a relationship between non-alcoholic greasy liver ailment (NAFLD) and serum nutrient D levels exist. Be that as it may, the unequivocal job and instruments are obscure.

Objective: To study serum vitamin D levels in patients with NAFLD was the aim of this study.

Place and Time of Study: Services hospital, Lahore from Feb 2018 to Jan 2019.

Methods: Analysis of NAFLD depends on ultrasound (U.S) discoveries after prohibition of other potential reasons for NAFLD and interminable liver infections. For this investigation, fifty patients with NAFLD and another 50 solid volunteers without NAFLD or any clinically clear liver ailments were enlisted. Serum 25(OH) nutrient D levels were estimated utilizing ELISA based test.

Results: Serum 25 (OH) nutrient D levels is conversely connected with weight list (BMI), all out cholesterol, triglycerides, low thickness lipoprotein (LDL) and age. There is measurably critical diminishing of serum 25(OH) nutrient D levels in patients with NAFLD than those without NAFLD.

Conclusion: It is concluded that Serum 25(OH) vitamin D level and NAFLD has inverse relation. **Keywords:** NAFLD, Fatty liver, Vitamin D.

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INTRODUCTION:

NAFLD has the probability of movement to nonalcoholic steatohepatitis (NASH), cirrhosis and hepatocellular carcinoma (HCC). The accurate instrument of NAFLD advancement is obscure. NAFLD is the most widely recognized liver infection in western countries [1]. It is normally connected with insulin obstruction and metabolic syndrome [2-3]. Nutrient D may have a job being developed of immune system ailments and provocative conditions through creation of incendiary cytokines. Nutrient D has numerous capacities in numerous frameworks of the human body, including muscles, bone, heart, gut, liver and safe framework [4-7]. An expanding proof is indicating a nearby connection between 25 (OH) nutrients D and NAFLD. A few examinations found a noteworthy relationship between nutrient D insufficiency and heftiness, metabolic disorder, type 2diabetes and insulin resistance [6-8]. Along these lines, to explore the conceivable relationship among NAFLD and serum 25(OH) nutrient D levels was the point of this examination.

METHODS:

We selected 50 patients with NAFLD (cases), and 50 sound volunteers without NAFLD or any ceaseless liver infection as controls, cross coordinated with patients in age and sex. This investigation is a cross sectional examination. It was carried out in order to examine the relationship among NAFLD and serum 25(OH) nutrient D.

Inclusion criteria:

All the patients selected were 18 years and above. All participants were the patients of NAFLD.

Detailed history, clinical examination and BMI calculation were done for every patient. All subjects (cases and controls) are admitted to Services hospital, Lahore.

Exclusion criteria:

with Wilson's disease The Patients or hemochromatosis were not selected for the study. Patients with renal failure and clinical, radiological or laboratory evidence of chronic liver disease were excluded. Patients who are taking drugs contain or affecting vitamin D level including: pure vitamin D drugs or multivitamins containing vitamin D formulas were expelled. People with hepatitis B or C infection and Alcohol users: daily consumption of more than 20 grams for males and 10 grams for females were not selected. This study does not include children.

Laboratory evaluation:

After a mid-night fasting, laboratory tests such as fasting blood sugar, lipid profile, complete blood count, liver functions (AST, ALT, serum bilirubin, INR, total protein and albumin) and renal function were carried out for all the patients.

Assessment of NAFLD:

Ultrasound examination was finished utilizing a Bmode arched test from 2.5-5 MHz on a Mindray DP-2200 ultrasound machine (Shenzhen, China). Finding of NAFLD was finished relying upon ultrasound imaging of the liver [9]. NAFLD was reviewed on a scale from 0-3 as indicated by Saverymuttu et al [10], where 0 is alludes to nonappearance of NAFLD, 1 mellow, 2 moderate and 3 serious NAFLD. Ultrasound examination was finished by a similar administrator who was visually impaired for the consequences of research center information and serum 25 (OH) nutrient D levels-for each subject around the same time of clinical examination and lab appraisal.

Vitamin D level:

Serum is isolated. It is put away in - 25° C for couple of days. It is an ELISA based test. By utilizing The Algeria 25-OH Vitamin D3/D2 Test Strip (ORGENTEC Diagnostika GmbH Carl-Zeiss-Straße 49-51 Mainz – Germany), 25(OH) nutrient D levels were estimated.

Ethical consideration:

Verbal assent was taken from people who took an interest in the examination. Servises hospital-Faculty of drug moral board of trustees endorsed the investigation.

Statistical analysis:

Multivariable direct relapse examination was performed to distinguish free hazard elements of nutrient D. Gauge statistic, clinical, and research center qualities were recorded as numbers and rate for clear cut information and means and standard deviation for constant information. Understudy's Ttest was utilized to think about aftereffects of constant factors among gatherings and chi square test for straight out factors. P-esteem thought about noteworthy if < 0.05. Measurable investigation was finished utilizing SPSS form 22 (IBM SPSS Inc., Chicago, US) for windows 10.

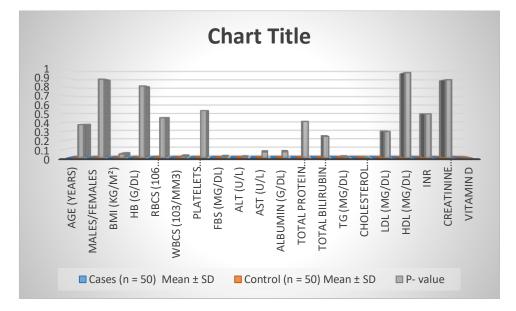
RESULTS:

Grade 1 NAFLD was available in 13 patients (26%), grade 2 out of 24 patients (48%), and grade 3 out of 13 patients (26%). Pattern attributes of patients and control were surrendered (Table 1). No noteworthy contrasts among guys and females in serum 25(OH)

nutrient D levels (Table 2). Patients with NA	\FLD
demonstrated higher white platelet tally, BMI, fa	asting

glucose (FBS), ALT, triglycerides and absolute cholesterol levels than controls.

Table 1: Baseline characteristics of cases and controls.				
	Cases $(n = 50)$ Control $(n = 50)$			
	Mean ± SD	Mean ± SD	<i>P</i> -value	
Age (years)	49.88 ± 9.126	47 ± 8.769	0.374	
Males/Females	27/23	26/24	0.886	
BMI (Kg/m ²)	26.18 ± 4.03	23.41 ± 2.64	0.045	
HB (g/dl)	12.11 ± 1.44	11.99 ± 1.033	0.809	
RBCs (106 cells/mm3)	4.64 ± 0.55	4.48 ± 0.64	0.453	
WBCs (103/mm3)	7.69 ± 1.92	6.13 ± 1.28	0.019	
Platelets (103/mm3)	266.08 ± 74.35	282 ± 59.48	0.533	
FBS (mg/dl)	135.28 ± 93.51	93.50 ± 19.89	0.013	
ALT (U/L)	54.42 ± 36.79	24.46 ± 10.43	0.008	
AST (U/L)	43.97 ± 27.53	27.30 ± 9.32	0.067	
Albumin (g/dl)	4.23 ± 0.34	4.02 ± 0.21	0.07	
Total protein (g/dl)	6.99 ± 0.67	6.79 ± 0.69	0.409	
Total bilirubin (mg/dl)	0.61 ± 0.24	0.51 ± 0.23	0.242	
TG (mg/dl)	200.25 ± 140.13	74.70 ± 34.74	0.008	
Cholesterol (mg/dl)	215.38 ± 50.81	150.30 ± 29.37	0	
LDL (mg/dl)	131.94 ± 33.42	145.12 ± 42.37	0.296	
HDL (mg/dl)	54.39 ± 17.41	54.70 ± 15.05	0.959	
INR	1.03 ± 0.09	1.06 ± 0.15	0.496	
Creatinine (mg/dl)	0.79 ± 0.28	0.77 ± 0.25	0.879	
Vitamin D	18.76 ± 14.37	40.36 ± 22.24	0	



BMI: body mass index, HB: Hemoglobin, RBCs: red blood cells, WBCs: white blood cells, FBS: fasting blood sugar, ALT: alanine transferase, AST: Aspartate transferase, TG: Triglycerides, LDL: low density lipoprotein, HDL: high density lipoprotein, INR: international normalized ratio.

NAFLD and 25(OH) vitamin D:

Complete cholesterol levels were conversely connected with serum 25(OH) nutrient D levels and

LDL levels were emphatically connected with serum 25(OH) nutrient D levels, generally no different elements influencing serum nutrient D levels. Serum

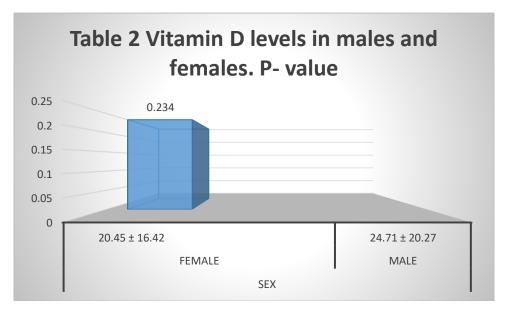
25 (OH) nutrients D levels were essentially decreased in patients with NAFLD than those without NAFLD (18.76 versus 40.36 p esteem 0.000) (Table 1). No noteworthy distinction in serum 25 (OH) nutrient D levels between patients with grade 1 NAFLD and those without NAFLD. Then again, there are noteworthy contrasts between patients with grade 2 and 3 on one hand and those with grade 1 and 0 then again. Serum 25(OH) nutrient D levels diminished with the expansion in the NAFLD grade (Table 3). Multivariable direct relapse investigation was utilized to evaluate indicators of serum 25 (OH) nutrients D (Table 4).

DISCUSSION:

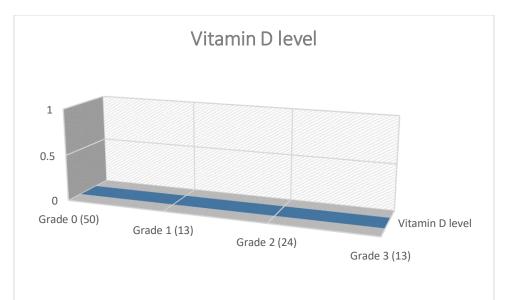
Patients in our investigation had typical engineered work, with no huge contrasts between NAFLD patients and control bunch in egg whites, INR or bilirubin levels. Along these lines, low dimensions of 25 (OH) nutrients D in NAFLD bunch in our examination isn't because of manufactured inadequacy of nutrient D by the liver.

Table 2: Vitamin D levels in males and females.

Vitamin	D (ng/dL)	Mean ± SD	P- value
	Female	20.45 ± 16.42	
Sex			0.234
	Male	24.71 ± 20.27	



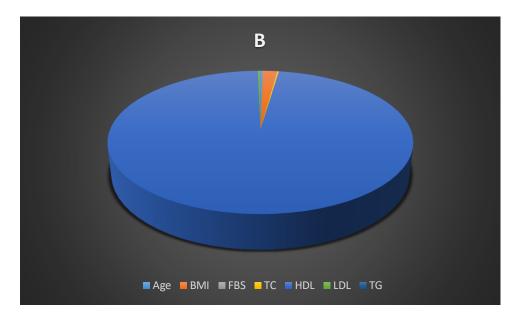
	Grade 0 (50)	Grade 1 (13)	Grade 2 (24)	Grade 3 (13)
Vitamin D level	40.36 ± 22.24 a	35.56 ± 20.83 a	13.39 ± 2.06 b	$12.74 \pm 5.56 \text{ b}$



a, b: there are significant differences between groups with different letters but not among groups with the same letter.

Determinant	B	S. E	Beta	<i>P</i> -value	
Age	-0.245	0.284	-0.121	0.395	
BMI	-1.274	0.697	-0.281	0.075	
FBS	-0.034	0.044	-0.16	0.447	
TC	-0.144	0.055	-0.435	0.013	
HDL	-71	0.156	-0.067	0.65	
LDL	0.203	0.07	0.401	0.006	
TG	0.021	0.03	0.157	0.493	

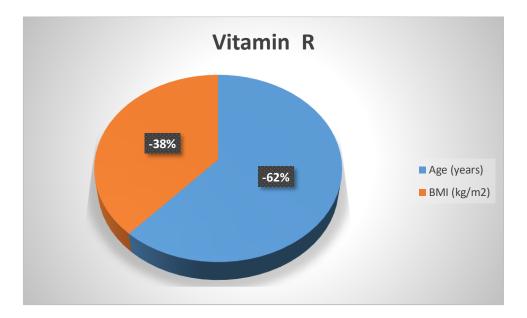
Table 4 Multivariable linear regression analysis of determinants of vitamin D level, Dependent variable: vitamin D.



BMI: body mass index, FBS: fasting blood sugar, TC: total cholesterol, HDL: high density lipoproteins, LDL: low density lipoprotein, TG: triglycerides

	Vitamin D (ng/dl)		
	R	P- value	
Age (years)	-0.3	0.034	
BMI (kg/m2)	-0.186	0.01	

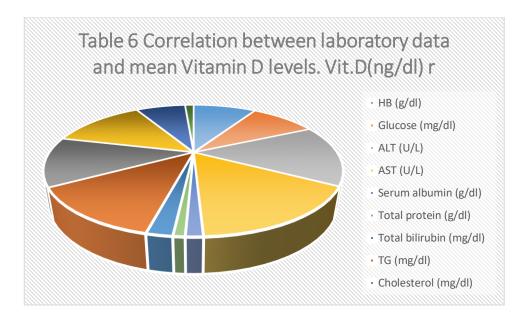
Table 5 correlation between Age & sex and mean vitamin D levels.



BMI: body mass index.

 Table 6 Correlation between laboratory data and mean Vitamin D levels.

	Vit.D(ng/dl)	
	r	P- value
HB (g/dl)	0.223	0.12
Glucose (mg/dl)	-0.232	0.109
ALT (U/L)	-0.383	0.006
AST (U/L)	-0.409	0.003
Serum albumin (g/dl)	-0.037	0.8
Total protein (g/dl)	-0.025	0.862
Total bilirubin (mg/dl)	0.056	0.7
TG (mg/dl)	-0.324	0.022
Cholesterol (mg/dl)	-0.309	0.029
LDL (mg/dl)	0.329	0.02
HDL (mg/dl)	0.176	0.22
Serum creatinine (g/dl)	-0.031	0.83



HB: hemoglobin, ALT: alanine transferase, AST: aspartate transferase, TG: triglycerides, LDL: low density lipoprotein, HDL: high density lipoprotein is expected to decrease due to synthetic dysfunction of the liver[11]. However, patients in our study had normal synthetic function, with no significant differences between NAFLD patients and control group in albumin, INR or bilirubin levels. So, low levels of 25 (OH) vitamin D in NAFLD group in our study is not due to synthetic deficiency of vitamin D by the liver. Also, vitamin D deficiency may induce NAFLD by impairing hepatic lipid metabolism[12]. Patients with vitamin D deficiency found to have high rates of insulin resistance, metabolic syndrome and inflammatory mediators including IL-4, IL-6 and TNF- α [13,14]. It was reported that vitamin D receptors widely exist in liver tissue with negative association between vitamin D receptors expression and necro-inflammatory grades of NASH[15]. Vitamin D may be sequestrated in the adipose tissue in obese patients[16]. Also, NAFLD patients may have a sedentary life with low sunlight exposure and nutritional imbalance.

Additionally, nutrient D lack may prompt NAFLD by disabling hepatic lipid metabolism [12]. The component by which 25 (OH) nutrients D may prompt NAFLD isn't clear. The liver proselytes' nutrient D to its dynamic structure, 25 (OH) nutrients D, so in liver maladies the 25 (OH) nutrient D levels is required to diminish because of manufactured brokenness of the liver [11]. In our investigation, patients with NAFLD demonstrated altogether diminished serum 25(OH) nutrient D levels than those without NAFLD. Serum 25(OH) nutrient D levels were observed to be logically

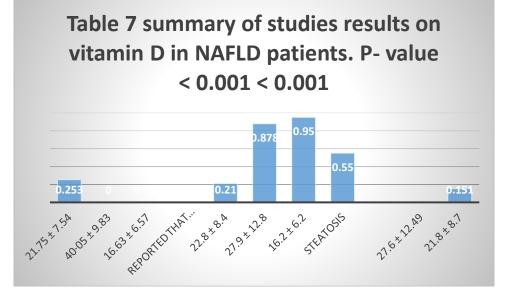
diminished with the expansion in the ultrasound grades of NAFLD. Nutrient D lack may prompt NAFLD by impeding hepatic lipid metabolism [12]. NAFLD patients may have a stationary existence with low daylight introduction and dietary lopsidedness. Patients with nutrient D lack found to have high rates of insulin opposition, metabolic disorder and fiery middle people including IL-4, IL-6 and TNF- α [13-14]. It was accounted for that nutrient D receptors broadly exist in liver tissue with negative relationship between nutrient D receptors articulation and necrofiery levels of NASH [15]. Nutrient D might be sequestrated in the fat tissue in stout patients [16]. Barchetta et al [18] 2011, found solid relationship between hypovitaminosis D and NAFLD which was autonomous on age, sex, BMI, lipid profile or glucose level. Chung et al [19] 2016, found that serum nutrient D levels were corresponded contrarily with NAFLD in a portion subordinate way autonomously on known NAFLD hazard factors. Our outcomes are in concurrence with Targher et al [17] 2007 who discovered huge reduction of nutrient D levels in patients with NAFLD than sound controls, which was related with NAFLD histo-pathological highlights. In our examination, nutrient insufficient patients (serum nutrient D under 20 mg/mL[20]) were fundamentally more established in age, more were females, with essentially higher aminotransferases levels than typical nutrient D subjects (serum nutrient D in excess of 30 mg/mL[20]). They additionally have essentially more elevated amounts of triglycerides and absolute cholesterol than subjects with typical dimensions of nutrient D (Table 8). Nelson et al [20] 2016, found that low dimensions of nutrient D are related with high danger of NASH in patients with NAFLD. Our

outcomes are likewise in concurrence with Zhai et al [21] 2016, Wang et al [22] 2016, Yang et al [23] 2017 and Eliades et al [8] meta-examination 2013 who revealed that nutrient D inadequate patients have 1.26-overlay expanded hazard for NAFLD than those with adequate nutrient D. In any case, our outcomes are in opposite with Li et al [24] 2013, who found no

noteworthy contrasts between patients with NAFLD and those without NAFLD in serum nutrient D levels. Additionally, Patel et al [25] 2016, Ha et al [26] 2017, De Paula et al [27] 2017 and Park et al [28] 2017 found no relationship between nutrient D levels and NAFLD (Table 7 abridges these investigations).

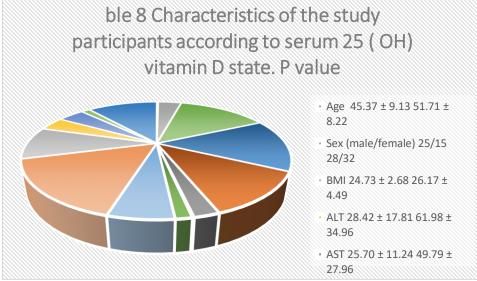
Study	NAFLD	Controls	P- value	
Targher et al 2007	51.0 ± 22	74.5 ± 15	< 0.001	
Barchetta et al 2011	14.8 ± 9.2	20.5 ± 9.7	< 0.001	
Chung et al 2016	21.75 ± 7.54	22.02 ± 8.43	0.253	
Zhai <i>et al</i> 2016	40.05 ± 9.83	40.98 ± 10.80	< 0.01	
Yang <i>et al</i> 2017	16.63 ± 6.57	18.44 ± 7.30	0.011	
Eliades <i>et al</i> 2013 meta-analysis	³ Reported that vitamin D deficient patients have 1.26-fold increased risk for NAFLD than vitamin D sufficient patients			
Li <i>et al</i> 2013	22.8 ± 8.4	22.1 ± 8.1	0.21	
Patel et al 2016	27.9 ± 12.8	27.7 ± 11.9	0.878	
Ha <i>et al</i> 2017	16.2 ± 6.2	16.2 ± 6.5	0.95	
De Paula <i>et al</i> 2017	Steatosis NASH 27.6 ± 26.5 ± 12.49 7.38 ± 10.00000000000000000000000000000000000	28.66 ± 8.40	0.55	
Park et al 2017	21.8 ± 8.7	21.5 ± 8.1	0.151	

Table 7 summary of studies results on vitamin D in NAFLD p	patients.
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	NVD	VDD	P value
Age	45.37 ± 9.13	51.71 ± 8.22	0.014
Sex (male/female)	25/15	28/32	0
BMI	24.73 ± 2.68	26.17 ± 4.49	0.213
ALT	28.42 ± 17.81	61.98 ± 34.96	0
AST	25.70 ± 11.24	49.79 ± 27.96	0.001
Bilirubin	0.58 ± 0.23	0.59 ± 0.24	0.813
Albumin	4.17 ± 0.26	4.19 ± 0.37	0.856
Total protein	6.90 ± 0.66	6.98 ± 0.69	0.687
FBS	103.00 ± 48.59	140.55 ± 98.92	0.139
TG	112.00 ± 82.00	213.84 ± 148.28	0.009
Total cholesterol	168.84 ± 35.30	222.90 ± 53.37	0
LDL	145.43 ± 32.29	127.92 ± 35.93	0.089
HDL	57.30 ± 16.36	52.71 ± 17.12	0.354
INR	01.04 ± 0.11	1.03 ± 0.10	0.946
Creatinine	0.82 ± 0.25	0.77 ± 0.28	0.53
HB	12.38 ± 1.21	11.90 ± 1.43	0.228
Platelets	284.74 ± 68.66	259.77 ± 72.41	0.234
WBCs	6.76 ± 1.33	7.75 ± 2.12	0.077
RBCs	4.56 ± 0.56	4.64 ± 0.58	0.64

Table 8 Characteristics of the study participants according to serum 25 (OH) vitamin D state.



NVD: normal vitamin D, VDD: vitamin D deficiency, BMI: body mass index, ALT: alanine transferase, AST: aspartate transferase, FBS: fasting blood sugar, TG: triglycerides, LDL: low density lipoprotein, HDL: high density lipoprotein, INR: international normalized ratio, HB: hemoglobin, WBCs: white blood cells, RBCs: red blood cells.

These conflicting outcomes among studies might be identified with contrasts in the examined populace, wholesome and ecological components.

REFRENCES:

- Patel YA, Henao R, Moylan CA. Guy CD, Piercy DL, Diehl AM, Abdelmalek MF. Vitamin D is Not Associated With Severity in NAFLD: Results of a Paired Clinical and Gene Expression Profile Analysis. *The American journal of gastroenterology* 2016; **111**: 1591-1598. [DOI: 10.1038/ajg.2016.406]
- Ha Y, Hwang SG, Rim KS. The Association between Vitamin D Insufficiency and Nonalcoholic Fatty Liver Disease: A Population-Based Study. *Nutrients* 2017; 9: [DOI: 10.3390/nu 9080806]
- De Paula FVL, Ramalho LNZ, De Paula FJA, Martinelli A. Low vitamin D level is not associated with severity of nonalcoholic fatty liver disease in morbidly obese patients. *Journal* of Hepatology 2017; 66: S157-S157. accession number: WOS: 000401056600326
- Park D, Kwon H, Oh SW, Joh HK, Hwang SS, Park JH, Yun JM, Lee H, Chung GE, Ze S, Park JH, Bae Y, Lee A. Is Vitamin D an Independent Risk Factor of Nonalcoholic Fatty Liver Disease?: a Cross-Sectional Study of the Healthy Population, *Journal of Korean medical science* 2017; **32**: 95-101. [DOI: 10.1038/ ajg.2016.406]
- Browning JD, Szczepaniak LS, Dobbins R, Nuremberg P, Horton JD, Cohen JC, Grundy SM, Hobbs HH. Prevalence of hepatic steatosis in an urban population in the United States: impact of ethnicity. *Hepatology* 2004; **40**: 1387-1395. [DOI: 10.1002/ hep.20466]
- Musso G, Gambino R, Cassader M, Pagano G. A meta-analysis of randomized trials for the treatment of nonalcoholic fatty liver disease. *Hepatology* 2010; **52**: 79-104. [DOI: 10.1002/hep.23623
- Targher G, Day CP, Bonora E. Risk of cardiovascular disease in patients with nonalcoholic fatty liver disease. *N Engl J Med.* 2010; 363: 1341-1350. [DOI: 10.1056/NEJMra0912063]
- Wang Y, Zhu J, D. HF. Where is the vitamin D receptor? Archives of biochemistry and biophysics. 2012; 123-133. [DOI: 10.1016/ jabb.2012.04.001]
- Holick MF, Vitamin D deficiency, *N Engl J Med.* 2007; **357**: 266281. [DOI: 10.1056/NEJMra070553]
- Ding C, Gao D, Wilding J, Trayhurn P, Bing C. Vitamin D signalling in adipose tissue, *Br J Nutr* 2012; **108**: 1915-1923. [DOI: 10.1017/S0007114512003285]
- 11. Targher G, Scorletti E, Mantovani A, Byrne CD. Nonalcoholic fatty liver disease and reduced serum vitamin D(3) levels. *Metab Syndr Relat*

Disord 2013; **11**: 217-228. [DOI: 10.1089/ met.2013.0044]

- 12. Kwok RM, Torres DM, Harrison SA. Vitamin D and nonalcoholic fatty liver disease (NAFLD): is it more than just an association? *Hepatology* 2013; **58**: 1166-1174. [DOI: 10.1002/hep.26390]
- Xia MF, Yan HM, He WY, Li XM, Li CL, Yao XZ, Li RK, Zeng MS, Gao X. Standardized ultrasound hepatic/renal ratio and hepatic attenuation rate to quantify liver fat content: an improvement method. *Obesity (Silver Spring)* 2012; 20: 444-452. [DOI: 10.1038/oby.2011.302]
- Saverymuttu SH, Joseph AE, Maxwell JD. Ultrasound scanning in the detection of hepatic fibrosis and steatosis. *British Medical Journal* 1986; **292**: 13-15. [PMC: 1338970]
- Lips P. Vitamin D physiology. *Prog Biophys Mol Biol* 2006; **92**: 4-8. [DOI: 10.1016/j.pbiomolbio. 2006.02.016]
- Norman AW. From vitamin D to hormone D: fundamentals of the vitamin D endocrine system essential for good health. *The American Journal* of Clinical Nutrition 2008; 88: 491S-499S. [PMCID : 18689389]
- 17. Alvarez JA, Ashraf A. Role of vitamin d in insulin secretion and insulin sensitivity for glucose homeostasis. *International Journal of endocrinology* 2010; 2010: 351385. [DOI: 10.1155/2010/351385]
- Roth CL, Elfers CT, Figlewicz DP, Melhorn SJ, MortonGJ, Hoofnagle A, Yeh MM, Nelson JE, Kowdley KV. Vitamin D deficiency in obese rats exacerbates nonalcoholic fatty liver disease and increases hepatic resistin and Toll-like receptor activation. *Hepatology* 2012; **55**: 1103-1111. [DOI: 10.1002/ hep.24737]
- Barchetta I, Carotti S, Labbadia G, Gentilucci UV, Muda AO, Angelico F, Silecchia G, Leonetti F, Fraioli A, Picardi A, Morini S, Cavallo MG. Liver vitamin D receptor, CYP2R1, and CYP27A1 expression: relationship with liver histology and vitamin D3 levels in patients with nonalcoholic steatohepatitis or hepatitis C virus. *Hepatology* 2012; 56: 2180-2187. [DOI: 10.1002/hep. 25930]
- Earthman CP, Beckman LM, Masodkar K, Sibley SD. The link between obesity and low circulating 25-hydroxyvitamin D concentrations: considerations and implications. *Int J Obes (Lond)* 2012; **36**: 387-396. [DOI: 10.1038/ijo. 2011.119]
- Targher G, Bertolini L, Scala L, Cigolini M, Zenari L, Falezza G, Arcaro G. Associations between serum 25-hydroxyvitamin D3 concentrations and liver histology in patients with

non-alcoholic fatty liver disease. *Nutr Metab Cardiovasc Dis* 2007; **17**: 517-524. [DOI: 10.1016/j.numecd. 2006.04.002]

- Barchetta I, Angelico F, Del Ben M, Baroni MG, Pozzilli P, Morini S, Cavallo MG. Strong association between non alcoholic fatty liver disease (NAFLD) and low 25(OH) vitamin D levels in an adult population with normal serum liver enzymes. *BMC Med* 2011; **9**: 85. [DOI: 10.1186/1741-7015-9-85]
- Chung GE, Kim D, Kwak MS, Yang JI, Yim JY, Lim SH, Itani M. The serum vitamin D level is inversely correlated with nonalcoholic fatty liver disease. *Clin Mol Hepatol* 0216; 22: 146151. [DOI : 10.3350/cmh.2016.22.1.146]
- 24. Nelson JE, Roth CL, Wilson LA, Yates KP, Aouizerat B, MorganStevenson V, Whalen E, Hoofnagle A, Mason M, Gersuk V, Yeh MM, Kowdley KV. Vitamin D Deficiency Is Associated With Increased Risk of Non-alcoholic Steatohepatitis in Adults With Non-alcoholic Fatty Liver Disease: Possible Role for MAPK and NF-kappaB? *The American Journal of* gastroenterology 2016; **111**: 852-863. [DOI: 10.1038/ajg. 2016.51]
- 25. Zhai HL, Wang NJ, Han B, Li Q, Chen Y, Zhu CF, Chen YC, Xia FZ, Cang Z, Zhu CX, Lu M, Lu YL. Low vitamin D levels and non-alcoholic fatty liver disease, evidence for their independent association in men in East China: a cross-sectional study (Survey on Prevalence in East China for Metabolic Diseases and Risk Factors (SPECT-China). *Br J Nutr* 2016; **115**: 1352-1359. [DOI: 10.1017/S0007114516000386]
- Wang D, Lin H, Xia M, Aleteng Q, Li X, Ma H, Pan B, Gao J, Gao X. Vitamin D Levels Are Inversely Associated with Liver Fat Content and Risk of Non-Alcoholic Fatty Liver Disease in a Chinese Middle-Aged and Elderly Population: The Shanghai Changfeng Study. *PloS one* 2016; 11: e0157515. [DOI: 10.1371/ journal.pone.0157515]
- Yang BB, Chen YH, Zhang C, Shi CE, Hu KF, Zhou J, Xu DX, Chen X. Low vitamin D status is associated with advanced liver fibrosis in patients with nonalcoholic fatty liver disease. *Endocrine* 2017; **55**: 582-590. [DOI: 10.1007/s12020-016-1152-x]
- Li LH, Zhang L, Pan SY, Wu XH, Yin XY. No Significant Association Between Vitamin D and Nonalcoholic Fatty Liver Disease in a Chinese Population. *Digestive diseases and sciences*. 2013; **58**: 2376-2382. [DOI: 10.1007/s10620-013-2658-1]