

## **DYSLIPIDEMIA IMPACT IN PRESENTATION OF CARDIOVASCULAR DISEASES OF PATIENTS WITH DIABETES MELLITUS AND DRUG TREATMENT OF DYSLIPIDEMIA**

**1.Nexhibe Nuhii**<sup>1</sup>, Doc. Dr. Sci. Med. Lutfi Zylbeari<sup>1, 2</sup>, 3.Dr. Driton Selmani<sup>1</sup>, 4. Dr. Ardiana Murtezani<sup>1</sup>

1. State University of Tetovo, Faculty of Medical Sciences, Pharmacy, Tetovo, Macedonia
2. Special Private Hospital For Nefrology And hemodialysis ,, Vita Medical Group "- Tetovo, Macedonia

**Abstract:**One of the major risk factors of early atherosclerosis and the introduction of thrombolytic processes and cardiovascular disease in patients with diabetes mellitus in addition to known factors (arterial hypertension, MIA syndrome, smoking, sedenterity, oxidative stress, psycho-stress, cytokines, etc ...) in recent years are counted also lipid abnormalities such as dyslipidemia or rather, are diabetic dyslipidemia. In patients with Diabetes Mellitus (DM type 1 and DM type 2 ) is proven and documented that there is a high positive correlation between hyperglycemia, glycosylated hemoglobin (HbA1c) and high lipid concentration values (LDL-ch and TG) and decrease in HDL-ch concentrations micro and macrovascular consequences, cardiovascular disease (CVD), retinopathy and diabetic nephropath(1) There are verifiable evidence that patients with insulin-dependent DM or treated with oral therapy are candidates with potential risk of cardiovascular diseases, peripheral vascular diseases, stroke compared with the healthy population. In the plasma of patients with DM were detected besides high concentrations of: blood glucose, glycosylated hemoglobin (HbA1c) were also detected high concentrations of LDL-ch and triglycerides and low concentrations of HDL-ch which further help the occurrence of cardiovascular disease (CVD) and coronary atherosclerosis complications (2). Aim of the paper work was to verify and document, role and correlation of lipid disorders (dyslipidemia) and hyperglycemia in the pace of progress and the appearance of cardiovascular diseases in patients with Diabetes Mellitus type.1 and the type 2 compared with healthy control individuals . The paper also aimed to influence positive effects of statins family in the treatment of hypercholesterolemia in patients with diabetes mellitus type 1 and type 2. In our patients treated with statins at the dose of 40 mg per day with duration of 3 months and reached a target of reducing the LDL cholesterol by 30-38%. The research was prospective cohort (, cross-section ") Totaly are included  $N^0 = 240$  examiners of whom 120 were patients of diabetes mellitus (DM 75 with tip1 while 45 were with DM type 2) while 120 individuals were healthy you served as group controllers. For examination was used 5+ (5) ml of venous blood taken from the vein in the patient lying position in order to avoid possible variations and the influence of the position of patients on lipid fraction values (9- 12%) which occur if the blood of patients is taken from the horizontal position. Dyslipidemia in diabetic patients with diabetes is present at the initial stages of an outbreak of the disease so its drug treatment in the early stages should be the primary postulate of physicians with which obviously would help the prevention and reduction of presentation of CVD

**Index Term:** Diabetes Mellitus (DM), blood glucose (Gl), the glycosylated hemoglobin (HbA1c), Lipids profiles, statins.

## 1 INTRODUCTION

Diabetes is one of the most massive diseases in the modern world with a tendency to increase the size of large and mostly appears in the developed and developing world (3). Diabetes is counted as the fourth cause of mortality in developed countries. A large number of studies have verified that epidemiologic regulation and control of sugar concentrations significantly reduced the rate of incidence of cardiovascular diseases (CVD) cerebral-vascular insults therefore the American Association for Diabetes (AAD) always provides guidance and recommendations on control and regulation of high values of glycemia and examination of glycosylated hemoglobin(HgbA1c) in patients with DM with which measures also reduce the risk of CVD, myocardial infraction and mortality of this group of patients. The control of hyperglycemia and glycohemoglobine (HgbA1c-average value of glycemia within three months) represents one of the primary measures in pursuit of the pace of progress to diabetes, so regular controls tracking and balancing of diabetes with dyslipidemia in the early stages of the disease, obviously would influence the prevention of the appearance of early atherosclerotic processes in coronary, cerebral and peripheral arteries .We always control glycemia and HgbA1c in patients with diabetes mellitus respecting the recommendations of AAD.

Nexhibe Nuhiu- State University of Tetovo, Faculty of Medical Sciences, Pharmacy, Tetovo, Macedonia  
Doc. Dr. Sci. Med. Lutfi Zylbeari -State University of Tetovo, Faculty of Medical Sciences, Pharmacy, Tetovo, Macedonia, Special Private Hospital For Nefrology And hemodialysis ,, Vita Medical Group "- Tetovo, Macedonia

Dr. Driton Selmani-State University of Tetovo, Faculty of Medical Sciences, Dentistry, Tetovo, Macedonia

Dr. Ardiana Murtezani-State university of Tetovo, Faculty of Medical Sciences, Medical Faculty, Tetovo, Macedonia

Recent years the incidence of unregulated diabetes and diabetic nephropathy and not only in the US and Europe but also the Balkans has an increase of 38% -42% which is due to: unregulated treatment of diabetes, psychostress, adiposity unrespected hygiene and dietary measures, excess consumption of fatty foods and disregard of ordinated therapy, smoking, physical inactivity, oxidative stress etc. Therefore, in recent years doctors always suggests that measurement and monitoring of blood glucose and lipid control to be one of the goals and measures mandatory for doctors of primary and secondary practice to what will be considerably decreased the incidence of SKV. So in the initial stages of presentation of Diabetes (DM) have dyslipidaemia and dyslipoproteinemia disorders with increased concentrations of LDL-ch, TG and HDL-ch reduction compared with patients with other diseases, so early examination of these disorders can significantly affect the prevention of the appearance of cardiovascular diseases (CVD (6,7). There are documented facts that the disorders of blood glucose and HgbA1c everytime in patients with DM are also associated with disturbance of lipid and therefore we decide to make our paper examinations lipid profile (total Cholesterol (CHT), Triglycerides(TG) Total lipid (TL), HDL and LDL-ch)], glycemia (GI) and glycosylated hemoglobin (HbA1c) in patients with diabetes-insulin users and patients treated with oral therapy. Patients with Diabetes Mellitus (DM) are at higher risk for early atherosclerosis and its consequences to the cerebrovascular system, cardiovascular and peripheral artery atherosclerosis compared with healthy population (4,5). Besides lipid abnormalities patients with DM have the disturbance of apolipoproteins. Apolipoproteines are integral protein of lipoproteinemic macromolecule specific to each class of them (8). Are related to lipid molecule using hydrophobic properties of fatty acids from phospholipids and polar part of the polypeptide chain (the process of inter-ionic reaction between phospholipids couples and opposite-charged amino acid alpha-helix electric to apoproteine. As factors underlying the appearance of cardiovascular diseases, cerebrovascular and early atherosclerosis in patients with DM apolipoproteines have an important role in metabolic disorders (9-11). Genetic factors of cardiovascular diseases, cerebrovascular and sclerotic processes are counted: the disruption of reverse transport of HDL-ch, cumbersome expression of B-receptors compared with E-receptors, reducing the conversion of VLDL to IDL

and LDL ch (12). The function of apolipoproteins is that they allow plasma lipid hydrosolubility in water (C<sub>h</sub>, TG, FL) of macromolecular complex-forming hydrosoluble lipoprotein (apolipoproteins) that are transported by the blood. The exact pathogenesis of diabetic dyslipidemia is not yet known; however, a large number of evidence suggest that insulin resistance has a central role in the development of this pathological phenomenon. The main cause of diabetic dyslipidemia is the release of fatty acids by increasing insulin-resistant fat and increased flux of free fatty acids in the liver in the presence of adequate stores of glycogen, which is still draining triglycerides encourages production, which in turn stimulates its secretion apolipoproteins-B (apo-B), Lp (a). and VLDL cholesterol. Diabetes mellitus – type 1 and generally well controlled rarely is associated with hyperlipidemia except diabetic ketoacidosis often associated with hypertriglyceridemia due to the increased release of tissue fatty acids (13-17). Pathological consequences of hypertriglyceridemia mostly appear to lipoprotein metabolism and early atherosclerotic manifestation. Anytime Diabetes is associated with high risk of cardiovascular disease (CVD). Management of diabetic dyslipidemia is a key element in a multi-factorial approach to prevent the occurrence of CVD in patients with diabetes. Patients with diabetes have a higher absolute risk of coronary disease presenting as patients without diabetes equally but with coronary disease, acute myocardial infarction and congestive heart failure, high prevalence of mortality(18,19) Lipid disorders ie diabetic dyslipidemia (atherogenic dyslipidemia) are always manifested by increased levels of triglycerides and LDL cholesterol and reduced level of cholesterol proatherogen-HDL-ch. Diabetic dyslipidemia is often helped by insulinemic resistance and is

present even before the diabetes. Small dense particles of LDL are more atherogenic due to their high sensitivity by increasing oxidative modification and the growth of taking the fat from the arterial wall. Overall, 30-40% of patients with diabetes suffer from diabetich dyslipidemia. All current national guidelines (NCEP- National Cholesterol Education Program) on the treatment of diabetich dyslipidemia as main target values have reduced the TG and LDL-ch and they suggest for LDL-c values from 100 to 70 mg / dl (20,21,22) as the optimal value for preserving the risk of coronary disease. NCEP recommendations association 2005 for the start of treatment of diabetes dyslipidemia of hypercholesterolemia namely with statin should be started when the values of LDL-ch are > 100 mg / dl to gain target effects of treatment with decreases in LDL-ch of 30-40 %, no pre-Liner LDL cholesterol levels, thus the lower the degree of risk of CVD. Results of many studies on the treatment of diabetic dyslipidemia and verified results have proven very successful during treatment with statine. In the case treatment with statin did not give proper effect to then preferably combined therapy, statin and niacin or statin with holestipol or holestiramin or fibrates with but any means combination niacin and fibrates between statins family due to the harmful effects of myositis or rhabdomyolysis consequences (23-27). Improvement and regulation of blood glucose values regardless of the type of dyslipidemia treatment has shown positive effects in improving lipid values. Beneficial effects in improving lipid abrevations in tip2 diabetic patients with oral therapy have shown metformin and rapaglinid treatments. There is documented evidence of these drug's influence on the improvement of diabetes and lipid disorders is closely linked with reduced levels of triglycerides and increased HDL-ch values (28,29,30)

## 2 Material and Methods Used

The research was prospective cohort („ cross-section ") Totally are included N<sup>0</sup> = 240 examiners of whom 120 were patients of diabetes mellitus (DM 75 with tip1 while 45 were with DM type 2) while 120 individuals were healthy you served as group controllers. For examination was used 5+ (5) ml of venous blood taken from the vein in the patient lying position in order to avoid possible variations and the influence of the position of patients on lipid fraction values (9- 12%) which occur if the blood of patients is taken from the horizontal position. Blood taken for examination inserted into the vial with a few drops heparin (5ccm serum) were sent for analysis in the

laboratory of Clinical Hospital of Tetovo and parallelly from a vial from the same patient was sent to the Institute of Clinical Laboratory in Skopje, in order to be verified and calibrated results obtained. Of the patients with DM (120) -54 (45%) of them were girls with an average age: 56.40 □12.80 but- 66 (55%) were male, with an average age: 59.50 □14:50 years. Group controller sound examination (voluntary blood donors) also were 54 (45%) women and 66 (55%) men with an average age identical: 15:00 □58.60 years. Of the total number of patients = N<sup>0</sup> = 120 with Type-1 diabetes mellitus (DM Tip1 th insulin dependent) were 75

while 45 were patients with Type-II diabetes mellitus (DM type 2 th treated with oral hypoglycemic), table number 1 .. Patients who were insulin dependent are counted as Type-1 while patients independent of insulin but with oral therapy, count as type-2 DM. So together with examination of concentrations of lipid profile, glycemia and the glicosylated hemoglobin (HbA1c) we made the determination of BMIx (Body Mass Index-table . no. 4). In all patients and the control group were analyzed lipid values of blood glucose and hemoglobin that is glycosylated within 3

months. The methods of determining the concentrations of lipid profile, blood glucose (GI) and HbA1c are identified in the table of number 2. As a reference value for GI and HbA1c values were taken according to criteria proposed by the World Health Organization (WHO) - {(GI = 3.5-6.5 mmol / l, (HbA1c% = 4.4% -6.6% T All analyzes are provided according to the study protocol and detected in the laboratory of the Institute of Clinical Laboratory of the University Clinical Center of the Medical Faculty in Skopje.

Table number 1: Reference Values and methods by authors whose blood glucose concentrations are determined, HbA1c and Lidids profiles are Presented in table 1.

Parameters Examined	Reference Values	Authors
LT	4-10g / l	ZOLLNER & Kirsch <sup>(74)</sup>
TG	0.68-l, 70 mmol / l	Buccola G. & H. David <sup>(75)</sup>
TCH	3, l, 5.2 mmol / l	CC. Allain et al <sup>(76)</sup>
LDL-ch	<3,4mmol / l, danger of adults:> 4.1 mmol / 1	Friedewalde & Fredricks on <sup>(77)</sup>
HDL-ch	> 1,6mmol / 1, danger of adults: <0.9 mmol / 1	WARNICKE G. et al <sup>(78)</sup>
Glicemia (GI)	3.5-6.5 mmol / L	Turbidimetric, camera-Cobas Integra 400
HbA1 c%	4.4-6.6%	Turbidimetric, camera-Cobas Integra 400

Table number 2: Presentation of diabetes patients under therapy

Tot. patients- N° = 120	DM type 1 (insulin-dependent)	DM type 2 (oral hypoglycemic)
	75	45

Table number 3: Distribution of patients by sex and age average

Gender	Number	The average age
Men	66 (55%)	58.60 □ 15:00
Women	54 (45%)	58.60 □ 15:00

Table number 4: Distribution of the control group average by gender and age

Gender	Number	The average age
Men	66 (55%)	56.40 □ 12.80
Women	54 (45%)	59.50 □ 14:50

The average age of patients was male gender =  $56.40 \pm 12.80$ , while female sex was =  $59.50 \pm 14.50$ , the average age difference between male and female according to statistics is nonsignificant  $p = 0.0005$ , which indicates a homogeneous groups (tab. 2)

Table. 5: Distribution of patients according BMIx: male = 75 and female = 45

BMIx	Male	Female
Poor Feeding	18	10
Normal feed	28	15
More feed	24	12
Obesity instance II-a	5	8

According table number 4. Differences between patients according to statistics is *nonsignificated*  $p < 0.0005$  and shows that working for homogeneous groups of patients.

### 3 Statistical processing of material examined

Values obtained of the blood glucose, HbA1c% and lipids (Kol.Total, TG, HDL-ch, LDL-ch) and control group are presented with mean values and standard deviation  $X \pm SD$ . In the results were also calculated correlation coefficient "r" statistical value of  $p \leq 1\%$  ( $p < 0.0001$ ). Statistics comparative lipid parameters between the two groups were analyzed to test the so-called

Studentov „t" while for examples dependent or independent and non-parametric tests were used tests: Mann-Whitney-U. significant statistics differences between the group of patients and control group obtained values of the parameters of lipids, glycemia and HbA1c% were analyzed to test the so-called „Anonova Two-Factor" statistical Worth „p" lesser of 5 %, namely  $p < 0.0005$ .

#### Results obtained:

The results obtained from the examination of blood glucose, HbA1c, lipid, (Kol.Total, TG, HDL-ch, LDL-ch) and the results obtained from the control group are presented in tables.2 and 3. Tables itself noted that the two groups of patients (DM Type-1 and DM-type 2) are verified high concentrations of lipids and HbA1% with significant statistical difference for  $p < 0.0001$ , compared with control group. Between values obtained of patients (with DM Type-1 and Type-2 DM) did not notice any significant difference facts that are consistent with many other studies (31,32). Lipid parameters presented a significant increase of the concentrations of: LDL-ch and TG and low concentrations of HDL-ch of the two groups of

patients with DM compared with the results from acquired by the group controller.

Table number 6: Presentation of the average Values of the Parameters analyzed to Examine patients with type 1 DM - the insulin-dependent  $N^0 = 75$  before treatment with hypopolypemic therapy.

Parameters	Number of patients	Average	Minimum	Maximum	$\pm$ SD
HbA1c%	76	10.80	6.80	13.80	5.80

Glycemia	75	9.60	7.6	14:00	3.20
TL (Total Lipids)	75	7.80	3.90	9.60	2.15
TG	75	3.70	1.26	4:50	0.80
TCH (Tot.Cholseterol)	75	5.60	2:50	7.90	2.14
HDL-ch	75	1:00	0.70	2.10	0.86
LDL-ch	75	4.60	4.20	5.80	0.94

Table number 7: Presentation of the average Values of the Parameters analyzed to Examine patients with type 2 DM dependent N<sup>o</sup> = 45 (oral hypoglocemic) -Before hippolypemic THERAPY TREATMENT.

Parameters	Number of patients	Average	Minimum	Maximum	± SD
HbA1c%	45	8.10	6.90	8.90	1.28
Glycemia	45	7.80	7:00	9:00	0.90
TL (Total Lipids)	45	7:50	5:40	12:50	3:40
TG	45	3:50	2:50	4.60	0.85
TCH (Tot.Cholseterol)	45	5.70	5.10	7.90	2.60
HDL-ch	45	1.12	0.80	2.30	0.60
LDL-ch	45	4.70	3.90	6.80	0.80

Table number 8: Presentation of the ***Mann-Whitney U-test*** for the Difference of the Values of the Parameters analyzed patients with DM type 1 and type 2 DM

Parameters	U	Z	p-level
Glycemia	6780.000	0.46895	0.860246
HbA1c%	8265.000	0.48280	0.006842
LT	1131.000	-0.13778	0.890417
<b>TG</b>	<b>655500</b>	<b>-3.25744</b>	<b>0.001124</b>
Cholesterol	1091.500	0.39693	0.691421
<b>HDL-ch</b>	<b>687800</b>	<b>-3.42614</b>	<b>0.001240</b>
<b>LDL-ch</b>	<b>8156.000</b>	<b>-3.456800</b>	<b>0.001460</b>

Was Recorded qual Difference Between the average seething of patients with DM type 1 and type 2 DM is josinjifikant for  $p < 0.005$ , Significant Difference Was Recorded only at: **TG** ( $p = 0.0011$ ), **HDL-ch** ( $p = 0.001124$ ) and **LDL -CH** ( $p = 0.00146$ )

Table number 9: Presentation of the average Values of the Parameters Examined in patients with DM Type 1, Type 2 DM and control group

P atients with - DM Type 1 and Type 2 DM						Controls Group		
Saw ERS	Number	Average	Minimum	Ma ximum	± SD	Average	± SD	p
<b>T L</b>	1 20	7. 80	2. 40	12.60	2.8 0	6. 40	0.60	<b>0 .0001</b>
<b>TG</b>	<b>1 20</b>	<b>3.85</b>	<b>2. 50</b>	<b>4. 80</b>	<b>0. 80</b>	<b>1. 28</b>	<b>0.63</b>	<b>0.0001</b>
<b>T Ch</b>	1 20	5. 80	4. 60	7. 40	0.9 2	4.9 0	1.2 4	<b>00:02 50</b>
<b>HDL-ch</b>	<b>1 20</b>	<b>1:03</b>	<b>0. 50</b>	<b>1. 15</b>	<b>0. 82</b>	<b>1.60</b>	<b>0. 60</b>	<b>0.0001</b>
<b>LDL-ch</b>	<b>1 20</b>	<b>4.20</b>	<b>3:40</b>	<b>5.4 0</b>	<b>0.9 5</b>	<b>3. 50</b>	<b>1.0 2</b>	<b>0.0001</b>
<b>Glycemia</b>	1 20	8. 60	4.90	9.80	4.6 5	5. 60	2. 10	<b>0.0001</b>
<b>HbA1c%</b>	1 20	8. 60	5. 80	12. 40	3. 90	7.20	3. 80	<b>0.0001</b>

Table 9: shows significant differences-p between the parameters examined between the patients with Diabetes mellitus (type 1 and type 2) and the control group. The difference which appears between the average values of the examined parameters of the two groups is significant statistic except total cholesterol values differ with  $p > 0.0005$ ). The values of the parameters examined LT, TG and LDL-ch, are higher of patients with DM-1 and DM-Tip Tip 2 with  $p < 0.0001$ , compared with control group. Lower values of patients with DM type 1 and type 2 DM compared with the control group were recorded only in HDL-ch for  $P < 0.0001$ .

Table number 10: Indicates significant differences between the examined parameters of patients with diabetes mellitus (type 1 and type 2) and the control group after 3 months after treatment with statins.

Parameters	Number of patients	Average	Minimum	Maximum	± SD	Controls group.Average	± SD
Glycemia	120	8.10	6.80	8.90	1.25	6:40	0.60
HbA1c%	120	7.60	7.100	8.70	0.80	<b>1.28</b>	<b>0.63</b>
<b>LT</b>	<b>120</b>	<b>7.80</b>	<b>5:40</b>	<b>7.10</b>	<b>1:40</b>	<b>4.90</b>	<b>1.24</b>
<b>TG</b>	<b>120</b>	<b>2.80</b>	<b>2.20</b>	<b>2.90</b>	<b>0.80</b>	<b>1.60</b>	<b>0.60</b>
<b>Cholesterol</b>	<b>120</b>	<b>5.70</b>	<b>4:50</b>	<b>5.60</b>	<b>1.20</b>	<b>3:50</b>	<b>1:02</b>
<b>HDL-ch</b>	120	1.18	0.80	2.70	12:50	5.60	2.10
<b>LDL-ch</b>	120	4:00	3.90	4.20.	0.60	7.20	3.80

From the table itself noted that the total lipid, triglycerides, total cholesterol and LDL-ch after treatment with statins doses of 12 weeks 1 tablet of 40 mg in the evening have significant reduction of their concentrations with  $p = 0.0001$  while the HDL fraction ch noticed a remodeling to increase its concentration, which testifies to the positive effects of statin for a double effect and the regulation of LDL hypercholesterolemia but also in increasing proatherogen HDL-ch concentration

#### 4 DISCUSSION:

Treatment of diabetic dyslipidemia recent years often by the concentrations of LDL-ch with decreased cholesterol values American Diabetes Association (ADA American Diabetes Association) of proatherogen (*HDL-ch*). In particular, patients with diabetes tend to have a significant increase of oxidized cholesterol (LDLox) and a higher percentage of particles ,, dyslipidemia in patients with diabetes. There are documented foam cells "which are highly susceptible to oxidation at high facts that the patients with diabetes from lipid fractions most risk consequences of submitting the Cardiovascular diseases often manifest hypertriglyceridemia (concentration increase (CVD, acute myocardial infarction, angina pectoris stable and of triglycerides-TG) and hypercholesterolemia-increased unstable coronary insufficiency ...). A large number of cohort

studies suggest that dyslipidemia and concentrations of elevated TG, LDL-ch and reduced concentrations of HDL-ch are at high positive correlation and independent predictor of CVD risk (33). In recent study by group of patients 5963 from ages > 40 years with dyslipidemia and diabetes treated with statins its verified a reduction and a decrease in LDL-ch for 22% and significant reduction in symptoms of CVD appearances (34). Observational studies of ADA American Diabetes Association together with friends Medical Nutrition Therapy -MNT- have verified

that patients who have used more healthy diet and increased physical activity (normal body weight) had decreased the triglycerides and LDL-ch to increase levels of HDL cholesterol and have had less symptoms of CVD (35,36,37). A large number of clinical studies for effects of treatment of diabetic dyslipidemia targeting the scope of medication therapy (statins, fibrates, niacin holoestipol, holestiramin) as target values for effective treatment have been proposed: LDL-ch <2.60 mmol / l in HDL cholesterol are = 1.02 mmol / l), and triglycerides levels are = 1.7 mmol / l). The females HDL-ch levels may be higher due to estrogens. Recommendations for treatment of dyslipidemia are always followed on the basis of recommendations and consensus proposed by the ADA and NCEP-National Cholesterol Education Program (38).

Hypertriglyceridemia may be a risk factor for CVD in people with initial diabetes. Initial hypertriglyceridemia therapy is consistent with dietary preventive measures such as: changes of way of life, weight loss, increased physical activity, limited consumption of saturated fats, reducing carbohydrates consumption, and reducing alcohol consumption, balancing diabetes (oral therapy or insulinemic) and then if the aforementioned measures do not show proper effects to then start therapy with medication of fibrates (gemfibrozil, fenfibrat, Clofibrat etc.) or in the cases of high hypertriglyceridemia fibrates may be combined with Niacin (<2 g / day. Often the clinicians presented the question of when and in which value of Tg should start treating hypertriglyceridemia? Decision to initiate pharmacological therapy depends on the judgment of the clinician - it must be between triglyceride levels from 2:30 to 4:50 mmol / l). The therapeutic combination of statins family and fibrates is prohibited due to the extremely high side effects of myositis and rhabdomyolysis. In case of high dyslipidemia these combinations are preferred therapy, statins with nicotinic acid, the statins with holestiramin or holestipol, fibrates with nicotinic acid, fibrates with holestiramin and holestipol, nicotinic acid with holestiramine or Holetipol. Choosing statins family should

depend mainly on lowering LDL necessary to achieve the goal of LDL-ch value of <100 mg / dL [2.60 mmol / l]). The use of statin therapy with high dose (eg 80 mg) to treat dyslipidemia in patients with high levels of LDL- ch and TG also shall be limited to because of side effects (increased transaminases and pain muscle) and therefore to these patients therapy should be started with the dose of 40 mg once a day and be accessed and then normalized target values after dosage laboratory examination shall be reduced to 20 mg per day. Patients with type 1 diabetes who are in good controled glycemia tend to have

normal levels of lipoprotein, unless they are overweight. Contained lipoprotein may be abnormal, but the effects of these anomalies in relation to CVD are unknown. Aggressive treatment of diabetic dyslipidemia decreases significantly the risk of CVD in patients with diabetes. The main purpose of therapy is to reduce the concentrations of LDL-ch to ≤ 100 mg / dL [2.60 mmol / l]. Initial pharmacological therapy consists and should be with the use of statins family. In case of submission of an intolerance to statins family then preferably be combined therapy also with other hypolipemics (such as niacin, holestipol, holestiramin, etc). Treatment of high levels of triglycerids be treated with fibric acid derivatives (gemfibrozil or fenofibrates) or niacin.

the result of unregulated diabetes we have manifestations of the disturbances in micro and macrovascular levels (39). There are documented facts that a large number of patients with DM are potential candidates for more comorbid conditions ranging from cardiovascular disease (ischemic heart disease, acute stroke infarction, angina pectoris unstable, left ventricular hypertrophy, congestive heart weakening, stroke, peripheral vascular disease, vascular complexity diabetic, diabetic retinopathy, diabetic nephropathy, etc. All of the aforementioned diseases are the main cause of frequent and morbidity and mortality of patients with unregulated diabetes (40-45) therefore the American Association of Diabetes always suggests the maintenance and regulation of normal glycemia values. Irregular checks and not balancing the glycemia is counted as one of risk factors for cardiovascular diseases and rapid progression of chronic renal damage in patients with diabetes whether they are insulin users or have oral hypoglycemic therapy (46-50). Numerous epidemiological studies and the American Association for Diabetes (AAD) have verified and documented that the regulation and regular check of glycemia decrease the risk of cardiovascular disease and myocardial infarction and their complications with which is reduced the rate of mortality of diabetic patients. Concentration of the hemoglobin that is



glycolysilated HgbA1c (which represents the average value of endogenous triglycerides) and decrease of HDL (lipoproteins glycemia within three months) is calculated as above standard that removes cholesterol from the blood, also referred to as " risk assessment of CVD in patients with clearing factor ") Increased cholesterol, endogenous triglyceride DM (53,54,55). American Association for diabetes ( ADA LDL and VLDL and HDL reduction, separately or combined American Diabetes Association)always calls and suggests for between them form the phatobiochemic and pathophysiologic mandatory screening of hemoglobin glycolysilated values in basis of birth and acceleration of the atherosclerotic process order to appropriately make decisions for treatment of diabetes that damages mostly large caliber medium caliber arteries in in order to reduce further diabetic complications [56 57]. The clinical practice known as atherosclerosis (63). Treatment of results of the acquired from lipid profile showed a high disorder diabetic dyslipidemia recent years often by the American for both groups of patients examined (also those with Type 2 Diabetes Association (ADA American Diabetes Association) has also those with DM-Tip.2) that complies with all studies on been the topic of discussion by proposing dietary and disorders profiles of lipoproteins in patients with DM. In the therapeutic measures on the management of dyslipidemia in presentation of the CVD and mortality rates in diabetic patients patients with diabetes mellitus. Patients with type 2 diabetes are exopt increased sugar level also affect many other factors spotential candidates to four fold risk of cardiovascular disease as: metabolic imbalance lipoapoprotein Apo-B and Lp (a), presentation (CVD) compared with the population which suffers disordered metabolism of carbohydrates, disorder of coagulation other diseases. There are documented facts that the factors, arterial hypertension, smoking, secondary patients with diabetes from lipid fractions most often manifest hyperparathyroidism, sedenterity,, oxidative stress hypertriglyceridemia (concentration increase of triglycerides-TG) etc. (58). Chronic hyperglycemia, combined with dyslipidemia and hypercholesterolemia-increased concentrations of LDL-ch hyperapolipoproteinemia increase the risk of morbidity and with decreased cholesterol values of proatherogen (*HDL-ch*). In mortality from cardiovascular diseases in uremic patients with particular, patients with diabetes tend to have a significant diabetes treated with terminal chronic hemodialysis. Beside increase of oxidized cholesterol (LDLox) and a higher disorder of carbohydrate metabolism diabetes as a chronic percentage of particles ,, foam cells "which are highly metabolic disorder impairs and other substances .Thus during susceptible to oxidation at high risk consequences of submitting diabetes predominates unraveling protein metabolism that is the Cardiovascular diseases (CVD, acute myocardial infarction, expressed by decreases in total protein level in the blood, and angina pectoris stable and unstable coronary insufficiency ...). A its special ingredients, such as: Albumins and globulins and large number of cohort studies suggest that dyslipidemia and globulins ingredients such as: alpha globulins, especially gamma concentrations of elevated TG, LDL-ch and reduced beta globulins which are protective antibodies for the concentrations of HDL-ch are at high positive correlation and organism (59,60.61, 62).). Protein breakup is clinically independent predictor of CVD risk(64) In recent study by group manifested with curbing of body growing.In diabetics there are patients 5963 from ages> 40 years with dyslipidemia and sensitive turbulences of lipids values .As we know the main lipids diabetes treated with statins its verified a reduction and a are : cholesterol, triglycerides, phospholipids and free fatty acids increase in LDL-ch for 22% and significant reduction in . These lipids in blood are not free but circulate with other symptoms of CVD appearances (65.66) . Observational studies substances as lipoproteins . First disorder of fatty metabolism of ADA American Diabetes Association together with friends diabetes is the increasing of lipolysis process (melting of fat Medical Nutrition Therapy -MNT- have verified that patients that occurs during the gluconeogenesis. This causes the who have used more healthy diet and increased physical activity increase in blood of free fatty acids which serve as the starting (normal body weight) had decreased the triglycerides and LDL- point for excess production of some biochemical substancesch to increase levels of HDL cholesterol and have had less which are called ketone bodies and therefore for the emergence symptoms of CVD (67,68,69). A large number of clinical studies of diabetes ketoacidosis. During diabetes by activation of many effects of treatment of diabetic dyslipidemia targeting the metabolic pathways, emerges the increase of the cholesterol scope of medication therapy (statins, fibrates, niacin holoestipol, and hypercholesterolemia and hypertriglyceridemia . On the cholestiramin) as target values for effective treatment have been other side for genetic reasons yet not finally clarified blood level proposed: LDL-ch are <2.60 mmol / l in HDL cholesterol are = rise occurs for some lipoproteins such as increased LDL 1.02 mmol / l), and triglycerides levels are = 1.7 mmol / l) . The (which carries blood cholesterol) of VLDL (which carries in blood HDL-ch levels may be higher due to

estrogens. Recommendations for treatment of dyslipidemia with nicotinic acid, fibrates with holestiramin and holestipol, always followed on the basis of recommendations and consensus proposed by the ADA and NCEP-National Cholesterol Education Program (70.71) . Initial hypertriglyceridemia therapy is consisted of dietary preventive measures such as: changes of way of life, weight loss, increased physical activity, limited consumption of saturated fats, reducing carbohydrates consumption , and reducing alcohol consumption, balancing diabetes (oral therapy or insulin therapy) and then if the aforementioned measures do not show proper effects to then start therapy with medication Group of fibrates (gemfibrozil, fenfibrat, Clofibrat etc.) or in the cases of hypertriglyceridemia fibrates may be combined with Niacin (1-2 g / day. Often the clinicians presented the question of when and which value of Tg should start treating hypertriglyceridemia Decision to initiate pharmacological therapy depends on the judgment of the clinician - it must begin between levels from 2:30 to 4:50 mmol / l). The therapeutic combination of statins family and fibrates is prohibited due to the extremely side effects of myositis and rhabdomyolysis. In case of high aggressive SKV. Aggressive treatment of diabetic dyslipidemia dyslipidemia these combinations are preferred therapy, with nicotinic acid, statins with holestiramin or holestipol, fibrates

## 5. Conclusion:

In conclusion we can say that the knowledge of mechanisms, etiopathogenesis, function and abnormalities on polymorphism and the negative impact of lipids (hypertriglyceridemia and hypercholesterolemia) and unbalanced glycemia of patients with diabetes mellitus (regardless of the type of diabetes) are among risky factors and independent in presentation CVD and premature atherosclerosis. Treatment and normalization of their highest values at the initial stages of the disease is of paramount importance and can significantly affect the prevention and deterrence pace of progress to early atherosclerotic processes and cardiovascular disease in these patients. Patients with diabetes (regardless of their type- insulin dependent diabetes mellitus or treated with oral hypoglycemic) are at same and high risk from the early appearance of atherosclerosis and cardiovascular disease. Therefore, improvement, balancing and regular checkups of diabetes and lipids with medicament therapy (statins, fibrates, niacin,

Holestipol, Holestiramina are the first step (per primam) in prevention and pace of progress and incidence of CVD and atherosclerotic processes . In treatment of uremic dyslipidemia in recent years a large number of studies have verified extremely high positive effects during treatment with statins (the dose of 40 mg) with what it seems is also contained and reduced the incidence of CVD presentation of diabetic patients and was also verified in our paper where we noticed a decrease in concentration of LDL-ch for 37% and 28-30% TG for facts that are consistent with other studies. We propose, based on preferences and consensus proposed by the American Association for Diabetes on the control of blood glucose, glycosylated hemoglobin ( HgbA1c) that treatment of diabetic dyslipidemia should be started in the initial stages of diabetes, no matter what type of diabetes what will be prevented visible appearances atherosclerotic phenomena (early atherosclerosis) in cardiovascular system, brain and peripheral arteries.

IJSER

## Literature:

1. Tarkun I, Cetiarslan B, Canturk Z. Lipoprotein(a) concentration in patients with type 2 diabetes mellitus without cardiovascular disease: relationship to metabolic and diabetic complications. *Nutr Metab Cardiovasc Dis* 2002 Jun; 12 (3) 127-131.
2. Ogbera O A, Azenabour O A. Lipoprotein(a), C-reactive protein and some metabolic cardiovascular risk factors in type 2 DM. *Diabetologia & Metabolic Syndrome* 2010; 2:51.
3. M. Bogoev. Diabetes Mellitus, Etiopatogenza, Mikrovaskularni komplikacii, Terapija. *Skopje-2003*; f:11-30.
4. Barbara Kolleris, Martin Auinger, Veronika Resing et al. Lipoprotein(a) as a predictor of Cardiovascular Disease in a Prospectively Followed Cohort of patients With Type 1 Diabetes. *Diabetes Care* July 2006, Vol.29, no.7;1661-1663 .
5. Shai I, Schulze MB, Manson JE, Stampfer MJ, Rifai N, Hu FB: A prospective study of lipoprotein (a) and risk of coronary heart disease among women with type 2 diabetes. *Diabetologia* 48, 2005; 2691-2692.
6. Maurus Marques de Almeida Holanda, Rosalia Gouveia Filizola, Maria Jose de Carvalho Costa et al. PLASMA LIPOPROTEIN (A) LEVELS –A comparison between diabetic and non diabetic patients with acute ischemic stroke. *Arq Neuropsiquiatr* 2004;62(2-A): 233-236.
7. Nawawi HM, Muhajir M, Kian YC, et al. Type of diabetes and waist-hip ratio are important determinants of serum lipoprotein(a) levels in diabetic patients. *Diabetes Res Clin Pract* 2002; 56: 221-227.
8. Alaupovic P, Kostner G, Lee DM, Conathy WL, Magnani HH. Peptide composition of human plasma apolipoproteins A, B and C. *Expos Annu Bioch* .Ponticelli C. et al. Lipid abnormalities in maintenance dialysis patients and renal transplant recipients. *Kidney Int Suppl*. 1978; 8: S 72.
9. Haas LB, Wahl PW, Sherrard DJ. A longitudinal study of lipid abnormalities in renal failure. *Nephron* 1983; 33: 145.
10. Somer JB. Et al. B. Lipoprotein lipids in chronic renal failure and hemodialysis: the influence of etiology and implication for atherogenesis. *Atherosclerosis* 1979; 34:353. *Med*. 1972;31:145-60.
11. Agarwal SK. et al. Prevalence of Chronic Renal Failure in adults in Delhi, India. *Nephrol Dial Transplant* 2005; (20) :1638-42. 12. Miida T, et al. LCAT-dependent conversion of pre  $\beta$ 1-HDL into  $\alpha$ -migrating HDL is severely delayed in haemodialysis patients. *J Am Soc Nephrol*. 2003;14:732-8.
12. Assmann G, Funke H. Genetische Diagnostik von Störungen des Lipoproteinstoffwechsels. *Lab Med*. 1992;16:369-74.
13. Scanu AM, Fless GM. Lipoprotein(a). Heterogeneity and biological relevance. *J Clin Invest*. 1990; 85:1709-15.
14. Taskinen MR (2003) Diabetic dyslipidaemia: from basic research to clinical practice. *Diabetologia* 46: 733–749.
15. Del Pilar Solano M and Goldberg RB (2005) Management of diabetic dyslipidemia. *Endocrinol Metab Clin North Am* 34: 1–25.
16. Chahal TJ and Ginsberg HN (2006) Diabetic dyslipidemia. *Endocrinol Metab Clin North Am* 35: 491–510.
17. Adiels M et al. (2007) Acute suppression of VLDL1 secretion rate by insulin is associated with hepatic fat content and insulin resistance. *Diabetologia* 50: 2356–2365.
18. Haffner SM, Lehto S, Ronnema T, Pyorala L, Laakso M: Mortality from coronary heart disease in subjects with type 2 diabetes and in non-diabetic subjects with and without prior myocardial infarction. *N Engl J Med* 339 : 229-234, 1998.
19. Miettinen H, Lehto S, Salomaa V, Mahonen M, Niemela M, Haffner SM, Pyorala K, Tuomilehto J, for the FINMONICA Myocardial Infarction Register Study Group: Impact of diabetes on mortality after the first myocardial infarction. *Diabetes Care* 21: 69-75, 1998.
20. Demacker PN, Veerkamp MJ, Bredie SJ, Marcovina SM, de Graaf J, Stalenhoef AF: Comparison of the measurements of lipids and lipoproteins versus assay for apolipoprotein B for estimation of coronary heart disease risk: a study in familial combined hyperlipidemia. *Atherosclerosis* 153 : 483-490, 2000

21. Executive Summary of the Third Report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III). *JAMA* 285:2486 - 2589,2001

22. Grundy SM, Cleeman JI, Merz CN, Brewer HB Jr, Clark LT, Hunninghake DB, Pasternak RC, Smith SC Jr, Stone NJ; the Coordinating Committee of the National Cholesterol Education Program; National Heart, Lung, and Blood Institute; American College of Cardiology Foundation, and American Heart Association: Implications of recent clinical trials for the National Cholesterol Education Program Adult Treatment Panel III guidelines. *Circulation* 110 : 227-239,2004.

23 American Diabetes Association: Standards of medical care in diabetes (Position Statement). *Diabetes Care* 28 (Suppl. 1):S4 - S36, 2005

24.. Colhoun HM, Betteridge DJ, Durrington PN, Hitman GA, Neil HA, Livingstone SJ, Thomason MJ, Mackness MI, Charlton-Menys V, Fuller JH: Primary prevention of cardiovascular disease with atorvastatin in type 2 diabetes in the Collaborative Atorvastatin Diabetes Study (CARDS): multicentre randomized placebo-controlled trial. *Lancet* 364:685 -696, 2004

25. Sever PS, Dahlof B, Poulter Wedel H, Beevers G, Caulfield M, Collins R, Kjeldsen SE, Kristinsson A, Mclnnes GT, Mehlsen J, Nieminen M, O'Brien E, Ostergren J, for the ASCOT investigators: Reduction in cardiovascular events with atorvastatin in 2,532 patients with type 2 diabetes. *Diabetes Care* 28 : 1151-1157,2005

26. Brown BG, Zhao XQ, Chait A, Fisher LD, Cheung MC, Morse JS, Dowdy AA, Marino EK, Bolson EL, Alaupovic P, Frohlich J, Albers JJ: Simvastatin and niacin, antioxidant vitamins, or the combination for prevent of coronary disease. *N Engl J Med* 345:1583 -1592, 2001

27. American Diabetes Association: Evidence-based nutrition principles and recommendations for the treatment and prevention of diabetes and related complications (Position Statement). *Diabetes Care* 26 (Suppl. 1):S51 - S61, 2003.

28. Pfeffer MA, Keech A, Sacks FM Cobbe SM, Tonkin A, Byington RP, Davis BR, Friedman CP, Braunwald E: Safety and tolerability of pravastatin in long-term clinical trials: prospective Pravastatin Pooling Project (PPP). *Circulation* 105:2341 - 2346, 2002

29. Heart Protection Study Collaborative Group: MRC/BHF Heart Protection Study of cholesterol lowering with simvastatin in 20,536 high-risk individuals: a randomized placebo-controlled trial. *Lancet* 360 : 7-22,2002

30. Gagne C, Bays HE, Weiss SR, Mata P, Quinto K, Melino M, Cho M, Musliner TA, Gumbiner B: Efficacy and safety of ezetimibe added to ongoing statin therapy for treatment of patients with primary hypercholesterolemia. *Am J Cardio* 90:1084 - 1091, 2002.

31. Ribault A, Drou MR, Letteulier C. et al. Determination of lipoproteina (a) concentration and lipoprotein(a) molecular weights in diabetic patients. *Diabetes Metab* 2000; 26: 107-112.

32. Lundstam U, Herlitz J , et al. Serum lipids, lipoprotein(a) level ,and apolipoprotein(a) isoforms as prognostic markers in patients with coronary artery disease. *I intern Med.* 2002; 251: 111-118.

33..Haffner SM: Management of dyslipidemia in adults with diabetes (Technical Review). *Diabetes Care* 21:160-178, 1998

33. Turner RC, Millns H, Neil HA, Stratton IM, Manley SE, Matthews DR, Holman RR: Risk factors for coronary artery disease in non-insulin dependent diabetes mellitus (UKPDS 23). *BMJ* 316:823-828, 1998

34. Heart Protection Study Collaborative Group: MRC/BHF Heart Protection Study of cholesterol-lowering with simvastatin in 5963 people with diabetes: a randomised placebo-controlled trial. *Lancet* 361:2005-2016, 2003

35. American Diabetes Association: Nutrition principles and recommendations in diabetes (Position Statement). *Diabetes Care* 27 (Suppl. 1):S36-S46, 2004

36. American Diabetes Association: Physical activity/exercise and diabetes (Position Statement). *Diabetes Care* 27:S58-S62, 2004

37. Grundy SM, Balady GJ, Criqui MH, Fletcher G, Greenland P, Hiratzka LF, Houston-Miller N, Kris-Etherton P, Krumholz HM, LaRosa J, Ockene IS, Pearson TA, Reed J, Smith SC, Washington R: When to start cholesterol-lowering therapy in patients with coronary heart disease: a statement for healthcare professionals from the American Heart Association task force on risk reduction. *Circulation* 95:1683–1685, 1997
38. NCEP Expert Panel on Detection, Evaluation and Treatment of High Blood Cholesterol in Adults: Executive Summary of the Third Report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III). *JAMA* 285:2486–2497, 2001
39. Lundstam U, Herlitz J, et al. Serum lipids, lipoprotein(a) level, and apolipoprotein(a) isoforms as prognostic markers in patients with coronary artery disease. *Intern Med*. 2002; 251: 111-118.
40. Lutfi Z. Profil na Dislipidemija i Apoproteinskite Aberaci kaj Pacienti Lekuvani so Povtoruvani Hemodijalizi Univerzitet „Sv. Kiril i Metodij“ Skopje Juni – 2009; *Doktorska Disertacija*
41. Koro CE, Bowlin SJ, Bourgeois N et al. Glycemic control from 1988 to 2000 among US adults diagnosed with type 2 diabetes: a preliminary report. *Diabetes Care* 2004; 27: 17–20.
42. Liebl A, Mata M, Eschwege E. Evaluation of risk factors for development of complications in Type II diabetes in Europe. *Diabetologia* 2002; 45: S23–28.
43. Saydah SH, Fradkin J, Cowie CC. Poor control of risk factors for vascular disease among adults with previously diagnosed diabetes. *JAMA* 2004; 291: 335–342.
44. American Diabetes Association. Tests of glycemia in diabetes (position statement). *Diabetes Care* 2004; 27: S91–S93.
45. Engelgau MM, Geiss LS, Saaddine JB et al. The evolving diabetes burden in the United States. *Ann Intern Med* 2004; 140: 945–950.
46. Wei M, Gaskill SP, Haffner SM et al. Effects of diabetes and level of glycemia on all-cause and cardiovascular mortality. The San Antonio Heart Study. *Diabetes Care* 1998; 21: 1167–1172.
47. Hanefeld M, Fischer S, Julius U et al. Risk factors for myocardial infarction and death in newly detected NIDDM: the Diabetes Intervention Study, 11-year follow-up. *Diabetologia* 1996; 39: 1577–1583.
48. Kuusisto J, Mykkanen L, Pyorala K et al. NIDDM and its metabolic control predict coronary heart disease in elderly subjects. *Diabetes* 1994; 43: 960–967.
49. Andersson DK, Svardsudd K. Long-term glycemic control relates to mortality in type II diabetes. *Diabetes Care* 1995; 18: 1534–1543.
50. Selvin E, Marinopoulos S, Berkenblit G et al. Meta-analysis: glycosylated hemoglobin and cardiovascular disease in diabetes mellitus. *Ann Intern Med* 2004; 141: 421–431.
51. Menon V, Greene T, Pereira AA et al. Glycosylated hemoglobin and mortality in patients with nondiabetic chronic kidney disease. *J Am Soc Nephrol* 2005; 16: 3411–3417.
52. Selvin E, Coresh J, Golden SH et al. Glycemic control and coronary heart disease risk in persons with and without diabetes: the atherosclerosis risk in communities study. *Arch Intern Med* 2005; 165: 1910–1916.
53. Dunn PJ, Cole RA, Soeldner JS et al. Reproducibility of hemoglobin A1c and sensitivity to various degrees of glucose intolerance. *Ann Intern Med* 1979; 91: 390–396.
54. Meigs JB, Nathan DM, Cupples LA et al. Tracking of glycosylated hemoglobin in the original cohort of the Framingham Heart Study. *J Clin Epidemiol* 1996; 49: 411–417.
55. American Diabetes Association. Standards of medical care in diabetes (position statement). *Diabetes Care* 2005; 28: S4–S36.
56. American Diabetes Association. Clinical practice recommendations. *Diabetes Care* 2006; 29: S3.
57. Collins AJ, Li S, Gilbertson DT et al. Chronic kidney disease and cardiovascular disease in the Medicare population. *Kidney Int Suppl* 2003; S24–S31).
58. Khaw KT, Wareham N, Luben R et al. Glycated haemoglobin, diabetes, and mortality in

men in Norfolk cohort of European prospective investigation of cancer and nutrition (EPIC-Norfolk). *BMJ* 2001; 322: 15–18.

59. Stump G.S.et.al. Alternations in Protein Metabolism in Diabetes Mellitus. At book; Joslin's Diabetes Mellitus. 14th Edition. Edited by C. Ronald

60. Kahn et al. Lippincott Williams and Wilkins, 2005: 275-290.

61. Kahn S.E. The Pathophysiology of Type II (Non-insulin-Dependent) Diabetes Mellitus. At book: Ellenberg and Rifkins Diabetes Mellitus 5th Edition. Edited by Daniel Porte Jr, M.D. and al. Appelton and Lange, 1997; 487-512.

62. Schaefer E.J.et.al. The Diagnosis and Management of Lipoprotein Disorders. At book: Medical Management of Diabetes Mellitus. Edited by Jack L. Lealy et.al. Marcel Dekker, Inc. 2000; 499-526.

63. Howard B.W.et.al. The Pathophysiology and Treatment of Lipid Disorders in Diabetes Mellitus. At book; Joslin's Diabetes Mellitus. 13th Edition. Edited by C. Ronald Kahn et.al. Lea and Febiger. 1994; 372-396.

64. Haffner SM: Management of dyslipidemia in adults with diabetes (Technical Review). *Diabetes Care* 21:160–178, 1998.

65. Turner RC, Millns H, Neil HA, Stratton IM, Manley SE, Matthews DR, Holman RR: Risk factors for coronary artery disease in non-insulin dependent diabetes mellitus (UKPDS 23). *BMJ* 316:823–828, 1998

66. Heart Protection Study Collaborative Group: MRC/BHF Heart Protection Study of cholesterol-lowering with simvastatin in 5963 people with diabetes: a randomised placebo-controlled trial. *Lancet* 361:2005–2016, 2003

67. Rubins HB, Robins SJ, Collins D, Fye CL, Anderson JW, Elam MB, Faas FH, Linares E, Schaefer EJ, Schectman G, Wilt TJ, Wittes J: Gemfibrozil for the secondary prevention of coronary heart disease in men with low levels of high-density lipoprotein cholesterol: Veterans Affairs High-Density Lipoprotein Cholesterol Intervention Trial Study Group. *N Engl J Med* 341:410–418, 1999

68. American Diabetes Association: Nutrition principles and recommendations in diabetes (Position Statement). *Diabetes Care* 27 (Suppl. 1):S36–S46, 2004

69. American Diabetes Association: Physical activity/exercise and diabetes (Position Statement). *Diabetes Care* 27:S58–S62, 2004.

70. Grundy SM, Balady GJ, Criqui MH, Fletcher G, Greenland P, Hiratzka LF, Houston-Miller N, Kris-Etherton P, Krumholz HM, LaRosa J, Ockene IS, Pearson TA, Reed J, Smith SC, Washington R: When to start cholesterol-lowering therapy in patients with coronary heart disease: a statement for healthcare professionals from the American Heart Association task force on risk reduction. *Circulation* 95:1683–1685, 1997

71. NCEP Expert Panel on Detection, Evaluation and Treatment of High Blood Cholesterol in Adults: Executive Summary of the Third Report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III). *JAMA* 285:2486–2497, 2001

72. American Diabetes Association: Detection and management of lipid disorders in diabetes (Consensus Statement). *Diabetes Care* 16:828–834, 1993

73. Haffner SM, Lehto S, Rönnemaa T, Pyörälä K, Laakso M: Mortality from coronary heart disease in subjects with type 2 diabetes and in nondiabetic subjects with and without prior myocardial infarction. *N Engl J Med* 339:229–234, 1998.

74. Zölner N, Kirchs KZ. Fotometriska-oboena metoda. *Ges Exp Med.* 1962; 135: 545.

75. Bucola G, David H. Quantitative determination of serum triglycerides by use of enzymes. *Clin Chem.* 1973; 19:476-82.

76. Allain CC, Poon LS, Chan CS, Richmond W. Enzymatic determination of total serum cholesterol. *Clin Chem.* 1974; 20:470-5.

77. Friedewald WT, Levy RJ, Fredrickson DS. Estimation of concentration of low density lipoprotein cholesterol without the use of the preparative ultracentrifuge. *Clin Chem.* 1972; 18:499-502

**Adress of the autors**  
**M-r. NEXHIBE NUHIU**  
**E-mail: nexhibe.nuhii@unite.edu.mk**

IJSER