The potential role of benfotiamine in the treatment of diabetic cardiac autonomic neuropathy

Abstract. Background. Cardiac autonomic neuropathy is a serious complication of diabetes mellitus that is strongly associated with approximately five-fold increased risk of cardiovascular mortality. Cardiac autonomic neuropathy manifests itself in a spectrum of things, ranging from resting tachycardia and fixed heart rate to the development of silent myocardial infarction. The significance of diabetic cardiac autonomic neuropathy has not been fully appreciated and there is no unified treatment algorithm. The purpose was to investigate the effects of benfotiamine on the heart rate variability, the corrected QT interval, QT dispersion and spatial QRS-T angle in patients with type 2 diabetes mellitus and cardiac autonomic neuropathy. Materials and methods. Thirty-two patients with type 2 diabetes mellitus and definite stage of cardiac autonomic neuropathy were allocated into two treatment groups: control (n = 15) received traditional antihyperglycaemic therapy; group 2 (n = 17) — benfotiamine 300 mg/day for three months in addition to standard treatment. Results. It was found that benfotiamine contributed to an increase in the sum of the squares of differences between adjacent normal-to-normal intervals, pNN50 (Δ% = +45.90 ± 7.91 %, p < 0.05), high-frequency component of heart rate variability during the active (Δ% = +25.80 ± 5.58 %, p < 0.05) and passive periods of the day (Δ% = +21.10 ± 4.17 %, p < 0.05), led to a decrease in the corrected QT interval (Δ% = −7.30 ± 1.36 %, p < 0.01), QT dispersion (Δ% = −27.7 ± 9.0 %, p < 0.01) and spatial QRS-T angle (Δ% = −24.4 ± 10.2 %, p < 0.01). Conclusions. The positive influence of benfotiamine suggests the feasibility of its administration to patients with type 2 diabetes mellitus and definite stage of cardiac autonomic neuropathy. Keywords: type 2 diabetes mellitus; cardiac autonomic neuropathy; benfotiamine; heart rate variability; corrected QT interval; spatial QRS-T angle
Hyperinsulinaemia can induce reversible prolongation of QTc (corrected QT for HR) in healthy subjects, hyperglycaemia and acute hypoglycaemia can induce the prolongation of QTc in both healthy and diabetic patients [11]. In patients with DM, prolongation of QTc was found during nocturnal hypoglycaemia that supports an arrhythmic basis for the dead in bed syndrome [12].

The spatial QRS-T angle, defined as the angle between the mean QRS and T vectors, indicates the main orientation of electrical heart activity during ventricular depolarization and repolarization, and has recently become an area of research interest. A wider QRS-T angle reflects an abnormal arrangement of ventricular repolarization and has been considered as a strong and independent risk factor for cardiac morbidity and mortality compared to other traditional cardiovascular risk factors and electrocardiographic (ECG) risk indicators such as the length of the QT interval [13–15]. The spatial QRS-T angle has recently been shown to be a strong and independent predictor of cardiac mortality in various groups of patients such as those with coronary artery disease (CAD), heart failure, T2DM, and elderly subjects [9, 15–17].

Therefore, the problem of effective treatment of CAN is particularly relevant. Pathogenetic treatment of CAN includes: balanced diet and physical activity; reducing insulin resistance; optimization of glycaemic control; treatment of dyslipoproteinaemia; correction of metabolic abnormalities in myocardium; prevention and treatment of thrombosis; use of aldose reductase inhibitors, gamma-linolenic acid, acetyl-L-carnitine, antioxidants, omega-3 polyunsaturated fatty acids, vasodilators, fat-soluble vitamin B1 (benfotiamine (BFT)), aminoguanidine; symptomatic treatment of concomitant diseases and syndromes (hypertension, coronary heart disease, heart failure, arrhythmias) and others [3, 18–21].

The purpose was to evaluate the effects of benfotiamine, a lipid-soluble thiamine derivative with higher bioavailability than that of thiamine, on the heart rate variability and the corrected QT interval, QT dispersion and spatial QRS-T angle in patients with type 2 diabetes mellitus and definite stage of cardiac autonomic neuropathy.

Materials and methods
To explore the effectiveness of some above-mentioned compounds, we examined 32 patients with T2DM and definite stage of CAN aged 50–59 years with disease duration of 1–6 years and median glycated haemoglobin A1c (HbA1c) 7.10 ± 0.40 %. Clinical characteristics of patients with T2DM and definite stage of CAN are given in Table 1.

CAN was diagnosed according to previously proposed criteria [1, 3].

The work was done according to the principles of the Declaration of Helsinki II and was approved by the medical ethics committee of Danylo Halytsky Lviv National Medical University (Minutes No. 1 dated August 29, 2016). All participants signed an informed consent prior to their inclusion in the study.

Patients with T2DM and definite stage of CAN were allocated into two treatment groups: group 1 (n = 15, controls) received traditional antihyperglycaemic therapy; group 2 (n = 17) — BFT 300 mg/day in addition to standard treatment. The duration of the therapy was 3 months.

The concentration of glucose in the blood was determined by the glucose oxidase method while HbA1c level was assessed using a highly sensitive method of ion exchange liquid chromatography with D-10 analyzer and Bio-Rad reagents (United States).

Resting 12-lead surface ECG with a paper speed of 25 mm/s and a signal size of 10 mm/mV was recorded in the morning. We performed resting ECG analysis, which included measurement of the following parameters: heart rhythm, HR, conduction intervals, and Holter-ECG (EC-3H recorder, Labtech, Hungary) analysis included measurement of 24-h ECG, circadian indexes and following

| Parameter                           | Patients with T2DM and definite stage of CAN (n = 32) |
|-------------------------------------|------------------------------------------------------|
|                                     | Controls (n = 15) | Benfotiamine (n = 17)            |
| Age, years                          | 55.33 ± 0.95     | 54.12 ± 0.65                     |
| Gender, n/%                         | Male             | 8/53.3                            | 10/58.8                           |
|                                     | Female           | 7/46.7                            | 7/41.2                            |
| Diabetes duration, years            | 3.60 ± 0.42      | 4.06 ± 0.36                      |
| BMI, kg/m²                          | 28.89 ± 0.16     | 26.66 ± 0.32                     |
| Medications, n/%                    | ACE inhibitors   | 12/80                             | 14/82.4                           |
|                                     | β-blockers       | 3/20                              | 4/23.5                            |
|                                     | Metformin        | 11/73.3                           | 11/64.7                           |
|                                     | Sulfonylurea     | 1/6.7                             | 1/5.9                             |
|                                     | Combined hypoglycaemic therapy | 3/20                           | 5/29.4                            |
|                                     | Hypertension, n/%| 12/80                             | 16/94.12                          |

Notes: BMI — body mass index; ACE — angiotensin-converting enzyme.
HRV parameters [22, 23]: standard deviation of all normal-to-normal (NN) intervals (SDNN), standard deviation of the means of all NN intervals for all 5-min segments of the entire recording (SDANNi), the square root of the means of all NN intervals for all 5-min segments of the entire recording (RMSSD), percentage of adjacent NN intervals that differ from each other by more than 50 ms (pNN50, %), the high-frequency component of HRV (HF), the low-frequency component of HRV (LF), the very low-frequency component of HRV (VLF), ratio of low to high frequency power components (sympathetic/parasympathetic ratio (LF/HF)). QTc was calculated by dividing the QT interval by the square root of the preceding NN interval time series (Bazett’s formula: QTc = QT/√NN) [24]. QT dispersion (QTd) was calculated as the difference between the maximum and minimum QTc. QRS-T angle is an ECG-derived measure of the difference in mean vectors of depolarization and repolarization. The absolute difference between the frontal QRS wave axis and T-wave axis was defined as frontal planar QRS-T angle. If such a difference exceeded 180°, the difference was calculated by subtracting from 180° [25].

Statistical analysis was based on the variation method using statistical parametric t-test, nonparametric Wilcoxon test, and Fisher’s, Pearson’s correlation coefficients. Data are presented as mean ± standard error of the mean (SEM). All tests were performed using the ANOVA (MicroCal Origin v. 8.0) software. Statistical significance was set at $p < 0.05$.

**Results**

We found that in patients with T2DM and definite stage of CAN, treatment hadn’t statistically significant influence on HbA1c level ($p > 0.05$).

The features of the time-domain HRV parameters in patients with T2DM and definite stage of CAN after BFT administration are given in Table 2.

Obtained results could prove that treatment with BFT in patients with T2DM and definite stage of CAN led to a significant decrease in the pNN50 and did not affect the SDNN, SDANNi and RMSSD (Table 2). Changes in the spectral heart rate variability parameters during the active period in patients with T2DM and CAN after 3 months of BFT therapy are given in Table 3.

Obtained results of this study could prove that treatment with BFT is accompanied by a significant increase in HF parameters during the active period compared to patients in control group (Table 3).

Changes in the spectral HRV parameters during the passive period in patients with T2DM and definite stage of CAN after 3 months of BFT therapy are given in Table 4.

### Table 2. Changes in the time-domain heart rate variability parameters in patients with T2DM and CAN after 3 months of benfotiamine therapy

| Parameter | Groups | Baseline | After treatment | Percentage change from baseline |
|-----------|--------|----------|----------------|---------------------------------|
| SDNN, ms  | Control group (n = 15) | 94.70 ± 4.84 | 91.30 ± 3.33 | -0.82 ± 2.73 |
|           | Benfotiamine (n = 17)  | 98.80 ± 2.73 | 103.8 ± 2.7 | +5.30 ± 1.64 |
| SDANNi, ms| Control group (n = 15) | 72.30 ± 4.08 | 74.70 ± 3.33 | +4.72 ± 2.67 |
|           | Benfotiamine (n = 17)  | 81.80 ± 3.18 | 87.40 ± 2.78 | +7.70 ± 2.33 |
| RMSSD, ms | Control group (n = 15) | 19.10 ± 1.36 | 18.70 ± 0.77 | +0.96 ± 3.67 |
|           | Benfotiamine (n = 17)  | 21.30 ± 1.42 | 23.10 ± 1.35 | +9.48 ± 2.37 |
| pNN50, %  | Control group (n = 15) | 3.90 ± 0.42 | 3.70 ± 0.23 | +7.10 ± 9.59 |
|           | Benfotiamine (n = 17)  | 4.94 ± 0.55 | 7.41 ± 1.06* | +45.90 ± 7.91 |

Notes: here and in Tables 3–5: the results are presented as absolute values and as percentage change from baseline ($\Delta\%$, mean ± SEM); * — $p < 0.05$ compared to baseline.

### Table 3. Changes in the spectral heart rate variability parameters during the active period in patients with T2DM and CAN after 3 months of benfotiamine therapy

| Parameter | Groups | Baseline | After treatment | Percentage change from baseline |
|-----------|--------|----------|----------------|---------------------------------|
| VLF, ms²  | Control group (n = 15) | 1,113.6 ± 98.1 | 1,103.9 ± 72.8 | +1.81 ± 2.63 |
|           | Benfotiamine (n = 17)  | 1,128.40 ± 80.34 | 1,241.2 ± 75.1 | +11.50 ± 1.99 |
| LF, ms²   | Control group (n = 15) | 390.80 ± 21.98 | 388.50 ± 14.74 | +1.60 ± 3.72 |
|           | Benfotiamine (n = 17)  | 414.8 ± 21.0 | 474.9 ± 21.9 | +16.80 ± 4.98 |
| HF, ms²   | Control group (n = 15) | 242.1 ± 18.5 | 244.5 ± 13.6 | +4.10 ± 4.15 |
|           | Benfotiamine (n = 17)  | 267.70 ± 17.61 | 326.40 ± 16.96* | +25.80 ± 5.58 |
| LF/HF     | Control group (n = 15) | 1.66 ± 0.06 | 1.63 ± 0.06 | -1.60 ± 3.07 |
|           | Benfotiamine (n = 17)  | 1.58 ± 0.04 | 1.47 ± 0.05 | -6.3 ± 2.9 |

Notes: here and in Table 4: total spectral power of all NN intervals between: VLF — 0.003 and 0.04 Hz; LF — 0.04 and 0.15 Hz; HF — 0.15 and 0.4 Hz.
It was found out that treatment with BFT is accompanied by a significant increase in HF parameters during the passive period compared to patients in control group (Table 4).

The features of the QTc, QTd and spatial QRS-T angle in patients with T2DM and advanced stage of CAN after treatment with BFT are given in Table 5.

Obtained results of this study could prove that prescription of BFT is accompanied by a more significant decrease in QTc, QTd and QRS-T angle compared to patients in control group (Table 5).

As a result of our studies, it was found that treatment with BFT contributed to a decrease in resting tachycardia (110 to 96 beats/min; p < 0.05), improvement of subjective feeling and an increase in tolerance to exercise loading. In addition, the majority of the patients with diabetic polyneuropathy (DPN) had a decrease and/or disappearance of pain, paresthesia, reduced frequency of muscle cramps, improvement and/or restoration of tactile, vibration and temperature sensitivity.

**Discussion**

Dysautonomia is a broad term that describes any disease or malfunction of the ANS. Evidence has been produced to indicate that various forms of mild to moderate vitamin deficiencies result in functional changes in the ANS. It is hypothesized that the predictable loss of efficiency in oxidative metabolism is the key to understanding the association of dysautonomia with many different diseases. Mild to moderate hypoxia and/or thiamine deficiency (TD) both give rise to exaggeration of centrally controlled mechanisms involved in all survival reflexes, mediated normally through a balanced reaction of the ANS and endocrine system. Together with dietary excesses, sugar consumption, particularly, appears to be responsible for triggering long-term disease related to the synthesis and use of cellular energy. Failure of the ANS cholinergic neurotransmission might follow from TD and/or other cofactors involved in glucose metabolism, exposing the organism to adrenal medullary release of epinephrine. Chronic exposure to moderate and severe hypoxia increases the activity of the ANS and adrenal medulla and TD induces an early functionally significant central muscarinic cholinergic lesion in rat studies [26].

Diabetes mellitus might be considered a thiamine-deficient state, if not in absolute terms then at least relative to the increased requirements deriving from accelerated and amplified glucose metabolism in non-insulin dependent tissues that, like the vessel wall, are prone to complications [27, 28]. The conventional indicator of thiamine sufficiency, erythrocyte transketolase (TKT) activity, is masked in clinical diabetes by increased levels of thiamine transporter-1 and thiamine transporter-2 proteins. The deficiency of thiamine in clinical DM may increase vascular fragility to the adverse effects of hyperglycaemia and, thereby, the risk of developing microvascular complications. Inhibition of TKT activity, and subsequent downregulation of the hexose monophosphate shunt, resulting in the accumulation of glyceraldehyde 3-phosphate, fructose 6-phosphate, and dihydroxyacetone phosphate, may be at least one mechanism in

| Parameter | Groups | Baseline | After treatment | Percentage change from baseline |
|-----------|--------|----------|----------------|--------------------------------|
| VLF, ms²  | Control group (n = 15) | 1,441.1 ± 106.7 | 1,439.3 ± 84.7 | +1.83 ± 2.47 |
|           | Benfotiamine (n = 17)  | 1,440.7 ± 90.9 | 1,572.4 ± 85.1 | +10.30 ± 1.98 |
| LF, ms²   | Control group (n = 15) | 509.10 ± 17.82 | 502.1 ± 13.0 | –0.60 ± 2.55 |
|           | Benfotiamine (n = 17)  | 524.1 ± 21.4 | 588.5 ± 25.8 | +12.70 ± 3.31 |
| HF, ms²   | Control group (n = 15) | 340.5 ± 25.3 | 326.70 ± 19.01 | –1.90 ± 3.38 |
|           | Benfotiamine (n = 17)  | 332.50 ± 18.39 | 396.6 ± 19.0 | +21.10 ± 4.17 |
| LF/HF     | Control group (n = 15) | 1.57 ± 0.08 | 1.59 ± 0.07 | +1.60 ± 1.92 |
|           | Benfotiamine (n = 17)  | 1.61 ± 0.05 | 1.50 ± 0.04 | –6.10 ± 2.75 |

| Parameter | Groups | Baseline | After treatment | Percentage change from baseline |
|-----------|--------|----------|----------------|--------------------------------|
| QTc, ms   | Control group (n = 15) | 433.40 ± 6.45 | 427.80 ± 4.72 | –1.10 ± 1.44 |
|           | Benfotiamine (n = 17)  | 423.10 ± 5.76 | 392.40 ± 7.74* | –7.30 ± 1.36 |
| QTd, ms   | Control group (n = 15) | 50.30 ± 4.53 | 46.00 ± 4.98 | –5.60 ± 6.97 |
|           | Benfotiamine (n = 17)  | 58.10 ± 3.94 | 39.40 ± 4.42* | –27.7 ± 9.0 |
| QRS-T angle, ° | Control group (n = 15) | 78.00 ± 6.44 | 69.70 ± 4.27 | –6.10 ± 5.52 |
|           | Benfotiamine (n = 17)  | 88.6 ± 6.4 | 59.30 ± 5.15* | –24.4 ± 10.2 |

Note: * — p < 0.01 compared to baseline.
the development of diabetes-induced vascular damage and other comorbidities [29–31].

Thiamine and its derivatives have been demonstrated to prevent the activation of the biochemical pathways (increased flux through the polyol pathway, formation of advanced glycation end products (AGEs) pathways, activation of protein kinase C (PKC), and increased flux through the hexosamine biosynthesis pathway induced by hyperglycaemia in DM). Thiamine definitively plays an important role in the diabetic endothelial vascular diseases (micro- and macroangiopathy), lipid disorders, retinopathy, nephropathy, cardiomyopathy, and neuropathy [32, 33]. Thiamine acts as a coenzyme for TKT and for the pyruvate dehydrogenase and alpha-ketoglutarate dehydrogenase complexes, enzymes which play a fundamental role in intracellular glucose metabolism. TKT and glucose-6-phosphate dehydrogenase, the rate-limiting enzymes of the pentose phosphate pathway, are inhibited in the diabetic heart under basal conditions [34].

Experiences from cardiology indicate that long-term increases in HRV and reduction in sudden cardiac death have only been shown with lipophilic agents that readily penetrate the blood nerve/blood brain barrier. In accordance with these observations, experimental data indicate a preventive effect of BFT on the development of CAN [35, 36]. High-dose therapy of thiamine and BFT inhibited AGES accumulation in the peripheral and reversed diabetic neuropathy potentially by reducing the levels of triose phosphates via activation of TKT [28, 37].

The evidence about positive thiamine effects was confined to hydrophobic thiamine metabolites that fulfill an important function under oxidative stress (OS) and nitr-o-syl stress. Thiamine protects nervous tissue probably by inhibiting nitric oxide (NO)-dependent tyrosine nitration and subsequent formation of dityrosine and interprotein tyrosine-tyrosine crosslink [31, 38]. Cardiac OS is involved in heart failure that is induced by thiamine deprivation in rats. These findings suggest that thiamine modules OS [33]. Nitric oxide synthase is an enzyme that is involved in the synthesis of NO, which regulates a variety of important physiological responses, including cell migration, the immune response, and apoptosis. Endothelial nitric oxide synthase and NO may play an important role in attenuating cardiac remodeling and apoptosis. Benfotiamine reduces OS and activates endothelial nitric oxide synthase to enhance the generation and bioavailability of NO, and it subsequently improves the integrity of vascular endothelium to prevent sodium arsenite-induced experimental vascular endothelial dysfunction [33, 38].

Thiamine supplementation can prevent hyperglycaemia-induced reduction in cell replication and proliferation as well as decreasing AGES formation. BFT has been shown to prevent an increase in markers of hexosamine biosynthesis pathway activity, intracellular AGES formation, intracellular PKC activity and the nuclear factor kappa B activation seen with in vitro hyperglycaemic damage [36]. Oral BFT in combination with the antioxidant alpha-lipoic acid treatment normalizes production of angiopoietin-2, a marker of increased intracellular methylglyoxal in endothelial cells, which contribute to AGES formation, and N-acetylglyceraldehyde-modified protein, a marker of hexosamine biosynthesis pathway activity [36]. Treatment with BFT has been shown to reduce activation of the polyol pathway of glucose metabolism and to increase TKT expression in the presence of hyperglycaemia [36]. Activation of AGE receptors in DM, found on cardiomyocytes, pericytes, and podocytes, stimulates post-receptor signaling, intracellular reactive oxygen species formation, and altered gene expression, leading to vascular damage [39].

Benfotiamine supplementation (100 mg/kg/day) for 14 weeks in streptozotocin-induced diabetic mice completely corrected hyperglycaemia-induced disruptions in Ca2+-homeostasis and mechanical functioning of cardiomyocytes [39].

In our previous investigations we have found that in patients with T2DM and definite stage of CAN, QRS-T (78.30 ± 1.95; p < 0.001), QTc (431.40 ± 2.94 ms; p < 0.001) and QTd (53.70 ± 1.49 ms; p < 0.01) were prolonged compared to patients without CAN [9]. An association between CAN and QT interval prolongation was demonstrated in many studies and it may predispose to sudden death in DM. Increased QTd was also suggested as a marker of diabetic autonomic neuropathy. Most of the data regarding QT interval and diabetic CAN are obtained in type 1 DM, with only few studies in T2DM [40].

The pathogenesis of QTc prolongation is multifactorial and includes imbalance in cardiac sympathetic innervation, intrinsic metabolic and electrolytic myocardial changes, left ventricular hypertrophy, CAD, and genetic factors could lead to QTc prolongation [41]. The day-night modulation of the QT/relative risk relation — on 24-h ECG recordings — was altered in CAN patients free of CAD, left ventricular dysfunction, or hypertrophy, with a reversed day-night pattern and an increased nocturnal QT rate dependence [34]. Reversible QTc prolongation may be induced by hyperinsulinaemia in healthy subjects, by hyperglycaemia and acute hypoglycaemia in both healthy and diabetic subjects [11, 41, 42]. In type 1 DM patients, prolonged QTc was shown to occur frequently during nocturnal hypoglycaemia and was associated with cardiac rate/rhythm disturbances. These findings support an arrhythmic basis for the dead in bed syndrome and possibly a provocative role of hypoglycaemia-induced sympathetic activation in cardiovascular events [1, 4, 12].

P. Valensi et al. demonstrated that changes in QTc can be considered as markers of cardiovascular autonomic dysfunction and as an important component in the potential prognostic value of the risk of arrhythmias [42]. Preserving the function of the parasympathetic nervous system in T2DM patients with CAN has a protective function, and the predominance of the sympathetic nervous system or the imbalance of LF/HF is harmful to the electrophysiological activity of the myocardium and may lead to changes in QRS-T [13, 14].

The spatial QRS-T angle was independently associated with glycaemic control, dyslipidaemia, and left ventricular myocardial performance in the diabetic subjects [13]. In the Rotterdam study, spatial QRS-T angle values ≥ 105° found in 20 % of patients with T2DM were associated with increased risk of cardiovascular mortality and sudden cardiac death. A spatial QRS-T angle < 75° was also significantly associated with increased risk for all clinical outcomes [43].
One recent study demonstrated that the spatial QRS-T angle is significantly wider in subjects with T2DM and CAN [14]. Moreover, presence and severity of CAN were the strongest predictors of the spatial QRS-T angle values. HRV parameters were significantly and independently associated with the spatial QRS-T angle, and explained almost 50% of its variability, suggesting the presence of a common pathophysiological ground linking the structural, functional and electrical myocardial disturbances in DM.

The results of our study showed that the prescription of BFT in the treatment of patients with T2DM and definite stage of CAN for 3 months contributed to a decrease in the QTc, QTd and QRS-T angle. Therefore, BFT may have therapeutic potential for neurological diseases by inhibiting inflammatory mediators and enhancing anti-inflammatory factor production [21, 33, 44].

The identification of the association of polymorphisms related to the genes of thiamine and TKT with DPN might be a first step in determining a DPN genetic risk profile with potential therapeutic repercussions. There is a moderate evidence from preclinical experimental models that high-dose thiamine and BFT: 1) inhibit hexose monophosphate, AGEs formation, and diacylglycerol-PKC through the TKT activation; 2) target various surrogate markers of hyperglycaemia-induced pathological processes; 3) can delay the progression of microangiopathic complications [45, 46].

Conclusions

1. The administration of benfotiamine for 3 months promotes an increase in parasympathetic link activity, particularly in pNN50, HF parameters during the active and passive periods of the day, and a decrease in the QTc, QTd, QRS-T angle. Obtained results suggest the feasibility of its use in the comprehensive treatment of patients with T2DM and definite stage of CAN.

2. The mechanism of BFT influence on diabetic CAN pathogenesis is not well-known. Thus, further investigations aimed at understanding the mechanism of action and confirming the beneficial effect of BFT on biochemical parameters, dynamics of independent cardiovascular tests, daily monitoring of electrocardiography, arterial wall stiffness parameters among patients with T2DM and definite stage of CAN are needed to validate this clinical findings.

Conflict of interests. There are no ethical/legal conflicts involved in the article. There are no conflicts of interest, including specific financial interests and relationships and affiliations relevant to the subject of this manuscript.

Participation of authors: V.A. Serhiyenko — collection of material, statistical data analysis, writing the manuscript text; V.B. Segin — management of the research, development of the study design, editing of the manuscript text; L.M. Serhiyenko — management of the research, development of the study design; A.A. Serhiyenko — management of the research, editing of the manuscript text. All authors contributed equally to the review. All authors have read and approved the final version of the manuscript.

Funding. No grant or funds have been received for this study.

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Pотенційна роль бенфотіаміну в лікуванні діабетичної автономної нейропатії серця

Резюме. Актуальність. Діабетична автономна нейропатія серця — серйозне ускладнення цукрового діабету, пов’язане з приблизно п’ятикратним підвищенням ризику серцево-судинної смертності. Автономна нейропатія серця проявляється широким спектром, розпочинаючи від тахикардії спокого і фіксованого серцевиття до розвитку безсимптомного інфаркту міокарда. Значення діабетичної автономної нейропатії серця до кінця не з’ясоване, також не існує єдиної алгоритму лікування.

Мета: дослідити вплив бенфотіаміну на стан варіабельності ритму серця, коригований інтервал QT, просторовий кут QRS-T у хворих на цукровий діабет 2-го типу з автономною нейропатією.

Матеріали та методи. Тридцять два пацієнта з цукровим діабетом 2-го типу та функціональною стадією автономної нейропатії серця були розподілені у дві групи лікування: контрольну (n = 15), яка отримувала стандартну цукрознижуючу терапію, та групу 2 (n = 17) — бенфотіамін 300 мг/добу доцільною додатково до традиційної терапії протягом трьох місяців.

Результати. Позитивний вплив бенфотіаміну свідчить про доцільність його призначення пацієнтом із цукровим діабетом 2-го типу та функціональною стадією автономної нейропатії серця.

Ключові слова: цукровий діабет 2-го типу; автономна нейропатія серця; бенфотіамін; варіабельність ритму серця; коригований інтервал QT; просторовий кут QRS-T

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Потенційна роль бенфотіаміну в лікуванні діабетичної автономної нейропатії серця

Резюме. Актуальність. Діабетична нейропатія серця — серйозне ускладнення цукрового діабету, зрослого з приблизно п’ятикратним підвищенням ризику серцево-судинної смертності. Автономна нейропатія серця проявляється широким спектром, розпочинаючи від тахикардії спокого і фіксованого серцевиття до розвитку безсимптомного інфаркту міокарда. Встановлено, що призначення бенфотіаміну викликало збільшення відсотка послідовних інтервалів NN, різниця між якими перевищує 50 мс, — pNN50 (Δ% = +45,90 ± 7,91 %, p < 0,05), високочастотної компоненти варіабельності ритму серця під час активного (Δ% = +25,80 ± 5,58 %, p < 0,05) та пасивного періодів доби (Δ% = +21,10 ± 4,17 %, p < 0,05), сприяло зменшенню коригованого інтервалу QT (Δ% = −7,30 ± 3,66 %, p < 0,01), дисперсії QT (Δ% = −27,7 ± 9,0 %, p < 0,01) та просторового кута QRS-T (Δ% = −24,4 ± 10,2 %, p < 0,01).

Висновки. Позитивний вплив бенфотіаміну свідчить про доцільність його призначення пацієнтам із цукровим діабетом 2-го типу та функціональною стадією автономної нейропатії серця.

Ключові слова: цукровий діабет 2-го типу; автономна нейропатія серця; бенфотіамін; варіабельність ритму серця; коригований інтервал QT; просторовий кут QRS-T

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