Physical rehabilitation therapy and research of lipid spectrum in students with vertebro-basilar disease within

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Manual and underwater massage of the cervicobrachial girdle were performed every other day in the amount of twenty sessions. Manual therapy was used in the form of the postisometric relaxation of all muscle groups of the cervicobrachial girdle with subsequent mobilization of those parts of the spine, where changes in the form of functional blocks in the intervertebral segments were detected by the manual and instrumental examination once a week for 2.5 months. Tractions of the cervical spine were performed if medically required under the subsequent X-ray control using a Gleason loop (15 sessions). Physical therapeutic exercises were assigned to all patients with the measurement of individual functional physical abilities using step-ergometric tests of functional classes 1 and 2. The training was carried out within the framework of stage 1 for 2–3 months. The criterion for completing was the reduction of HR (heart rate) at the usual loads from peak to plateau formation per 20–35 cardiac contractions.

The tissues lipid spectrum study was carried out according to the method modified by Prof. P.V. Stapay (Institute of Animal Biology, NAAS) [9]. The following procedure is based on the classic Folch method and permits to obtain the largest amount of lipids from the tissues. For this purpose, the hair lock on the hind head was cut off in such a way that when re-cut, after 3 months, the newly grown hair would be obtained. The selected material was washed with shampoo in warm water, then in cold physiological saline and dried using filter paper.

For the extraction of lipids, a mixture of chloroform and methyl alcohol solvents was used in a volume ratio of 1:2. The material was first thoroughly cut very small with scissors, and then mulled in liquid nitrogen in a metallic mortar to a powdered state. The amount of the crushed hair weighing 0.5 g was filled with an extraction mixture and homogenized with a teflon piston. The homogenate was transferred into the test tube and left for extraction for 12 hours at the room temperature. Later, the homogenate was filtered through a fat-free paper filter. The precipitate was washed three times with small portions of the extraction mixture. Extracts were combined. To remove water-soluble, i.e. non-lipid admixture, 0.74M KCl solution was added to the extract in the amount of 1/4 of the lipid extract total volume. The content was shaken up and left for sedimentation until the complete stratification. The upper water-methanol layer, lipids-free, is sucked off with a water-jet pump.

For the study, a lower chloroform layer containing lipids was used. It was filtered again through a fat-free paper filter into measuring tubes and brought the volume of 10 ml with the extraction mixture. Usually the weight method is used to determine the total amount of lipids in biological objects. However, it is suitable mainly for objects with high lipids content, which is not always possible. Therefore, we suggest the self-developed simple and easily reproducible method stipulating the use of concentrated sulfuric acid.

The amount of 0.5 ml lipid extract is introduced into pure and dry test tubes with pipettes. After preliminary evaporation of the extraction mixture under the exhaust hood, 5 ml of concentrated sulfuric acid is added to the test tubes and boiled for 15 minutes in a water bath. The content of the tubes is thoroughly mixed with a glass rod, cooled and measured using a blue light filter, using a 10 mm thick cuvet. Simultaneously, a blank sample with 5 ml of sulfuric acid is studied. The total amount of lipids is calculated by means of the calibration curve.

Lipid fractions were determined by the thin-layer chromatography method. Glass plates sizing 20x6 cm, thoroughly washed in soda solution and chrome mix, were dried at room temperature. As an adsorbent, silica gel with particles of 120x140 mesh (10–40 μm) was used. Silica gel was rubbed in a mortar with gypsum, which content was no more than 5%. The suspension was immediately poured onto a dry glass plate and distributed on its surface with the layer thickness of 0.3 mm. The dried plates were placed into the thermostat for 45 minutes at 110°C for activation.

The tested sample was applied to the starting line in the concentration of 2 mg per 0.05 ml of the solvent. The test plate was placed in a chromatographic chamber with a solvent (the mobile phase, a mixture of petroleum and diethyl ether in the ratio of 4:1). To saturate the chamber with the vapor of the solvent, its side walls and the bottom are covered with filter paper. Upon reaching the finish mark, the chromatograms are transferred to the exhaust hood at the temperature of 110°C.

Persistent color spots appear on silica gel. The chromatograms are placed in a glass jar, densely closed with bits of crystalline iodine. The developing process time is 10–20 minutes. After evaporation of iodine, the extract is harvested into pre-weighted bottles, and after complete removal of the solvent (with vacuum), the weighing bottles are weighed. The calibration curve is built for each individual class and their number is determined in the tested sample.

Results and discussion

The value of different lipids fractions, as can be seen from Table, in patients with VBI did not change unidirectionally. The amount of polar lipids before treatment was 0.17±0.004 mmol/l, and after the treatment it increased to 0.19 mmol/l. The Student’s coefficient was 2.9 at the variation-statistical data processing. In the correlation data analysis before and after the treatment there was a high degree of unidirectional changes, which is confirmed by the respective correlation coefficient – 0.7. Changes in the amount of cholesterol esters in the hair of patients compared to the polar lipids were less statistically reliably pronounced. Free fatty acids have undergone very slight changes, which are not statistically reliable.

Triacylglycerides have undergone already more significant changes. Before treatment, their amount in tissues was 0.39±0.07 mmol/l. After the treatment, their content in these tissues was 0.32±0.01 mmol/l with Student’s coefficient of 0.89. Such a sufficiently significant growth is statistically reliable (p<0.05). In the correlation analysis, mostly unidirectional changes were detected in over a third of the studied cases.
### Table. Content of lipid fractions in the patients’ hair

<table>
<thead>
<tr>
<th>Lipid fractions before and after treatment</th>
<th>Amount</th>
<th>p</th>
<th>t</th>
<th>r</th>
</tr>
</thead>
<tbody>
<tr>
<td>Polar lipids</td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Before treatment</td>
<td>0.17±0.004</td>
<td>&lt;0.05</td>
<td>2.9</td>
<td>0.7</td>
</tr>
<tr>
<td>After treatment</td>
<td>0.19±0.005</td>
<td></td>
<td></td>
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<tr>
<td>Cholesterol esters</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Before treatment</td>
<td>0.42±0.007</td>
<td>≥0.05</td>
<td>1.9</td>
<td>0.87</td>
</tr>
<tr>
<td>After treatment</td>
<td>0.44±0.008</td>
<td></td>
<td></td>
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<tr>
<td>Free fatty acids</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before treatment</td>
<td>0.35±0.03</td>
<td>≥0.05</td>
<td>1.9</td>
<td>0.4</td>
</tr>
<tr>
<td>After treatment</td>
<td>0.36±0.04</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Triacyglyceride</td>
<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Before treatment</td>
<td>0.39±0.07</td>
<td>≤0.05</td>
<td>0.89</td>
<td>0.3</td>
</tr>
<tr>
<td>After treatment</td>
<td>0.32±0.01</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Free cholesterol</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Before treatment</td>
<td>0.42±0.009</td>
<td>&lt;0.05</td>
<td>2.4</td>
<td>0.89</td>
</tr>
<tr>
<td>After treatment</td>
<td>0.39±0.008</td>
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</table>

**Fig.** Lipids content changes before and after the treatment performed.

Free cholesterol before treatment was 0.42±0.009 mmol/l. After treatment, its value dropped to 0.39±0.008 mmol/l, which is statistically reliable. Free fatty acids at cerebral circulation disorder therapy (CCDT) have undergone very small, statistically unreliable changes. The control group served as a group of healthy adolescents numbering 20 persons (12 girls and 8 boys aged 15–18 years) in which the content of lipid fractions in hair was distributed as follows: polar lipids – 0.24 mmol/l, cholesterol esters – 0.45 mmol/l, free fatty acids – 0.37 mmol/l, triacylglycerides – 0.31 mmol/l, free cholesterol – 0.375 mmol/l.

When applying physical therapy, we see some changes in most of the lipid fractions in the hair. As it can be seen from figure 1, there was a statistically reliable increase in the polar lipids amount after treatment, and a statistically reliable decrease in free cholesterol and triacylglycerides. At the same time, the value of free cholesterol and triacylglycerides reached the normal level in the control group of the studied patients.

Such a result has its scientific explanation and practical significance. Considering that the polar lipids fraction is nonatherogenic, whereas free cholesterol and triacylglycerides contribute to atherogenesis [8], the increase in the polar lipids amount and the decrease in the amount of triacylglycerides and free cholesterol after treatment can be interpreted as a positive effect of the performed physical therapy on this vascular pathology regression.

In the objective and instrumental examination of adolescents after the performed treatment, a clear positive dynamics of pathological symptoms regression was observed. Namely: in most of the examined patients, periodic dizziness, headache, visual impairment, vertebral listhesis and smoothed lordosis disappeared, in some cases the symptoms became very slight. On the ultrasound scans of extracranial vessels after treatment, the signs of extravasal compression of the vertebral arteries were not observed in more than half of the patients, while in the rest they became significantly less pronounced.

**Prospects for further researches**

Conduct research into the effectiveness of physical therapy using theoretical and methodological and pedagogical analysis in a comprehensive study of aspects of correction of vertebral-basilar syndrome in students.
Conclusions

1. The obtained data indicate a certain value of the tissue lipids content disorder in the pathogenesis of the vertebralbasilar insufficiency syndrome.

2. Correction of the cervical spine vertebral disorders by means of physical therapy leads to the elimination of extravascular compression signs and the restoration of lipid metabolism in tissues, which is accompanied by improvement of the clinical condition in patients and by the neurological deficiency regression.

Literature


References


The purpose of the work was to study the effect of physical therapy on the course of vertebrobasilar syndrome in adolescents.

Materials and methods. In the complex of physical therapy, methods of therapeutic exercises, treatment, massage, traction-manual therapy, and electrophysiotherapeutic procedures were used. The tissues lipid spectrum study was carried out according to the modified Folch method and permits to obtain the largest amount of lipids from the tissues.

Results. The therapeutic effect of physical therapy was manifested not only in the normalization of lipid metabolism, but in reducing the severity of the main clinical disease manifestations, which makes expedient the use of this therapy.

Conclusions. Correction of the cervical spine vertebral disorders by means of physical therapy leads to the elimination of extravascular compression signs and the restoration of lipid metabolism.

Key words: physical rehabilitation therapy, vertebrobasilar insufficiency syndrome, tissue lipid balance.