

**RESEARCH OF THE BEHAVIORAL REACTIONS OF RATS AFTER
AN INTRANASAL ADMINISTRATION OF MODIFIED FRAGMENT OF NEUROPEPTIDE Y**

Havrylov I.O. Kuchmuk D.A., Oderiy I.V.

Scientific supervisor: prof. Zagayko A. L.

National University of Pharmacy, Kharkiv, Ukraine

gavrilov.i.ok@gmail.com

Introduction. Neuropeptide Y (NPY) is a regulatory compound that performs many functions in the body, including the processes of food consumption, reaction to stress, learning, etc. A modified NPY fragment, which is an analogue of the functionally active NPY site, was studied in this work.

Aim. Study of the effect of a low molecular weight fragment of neuropeptide Y on the behavior of rats after intranasal administration.

Materials and methods. The experiment was performed on 30 rats, divided into 5 equal groups. The intact group received water intranasally, the 2nd, 3rd, 4th — a solution of the test compound in a dose of 0.1, 0.15 and 0.2 mg / kg. Comparison drug – Stresam "Bicodex" group 5, was administered per os (22 mg/kg). The following tests were performed: the light/dark test (LDT) and a forced swimming test (FST).

Results and discussion. In the light/dark test for group 2, the time spent in the light chamber increased by 30% compared to the control, but was less than the Stresam group (41%). The time in the light chamber for groups 3 and 4 is not significantly different. Latency for the rate to enter into the dark for groups of 0.1; 0.15 and 0.2 mg/kg was higher, respectively, in 2.4, 5.4 ($p < 0.05$) and 1.9 times, compared with control. The latent time of group 2 was relatively equal to group 5. The number of peeps out and transitions between groups did not differ significantly. In FST, the total time of immobilization was dose-dependently reduced compared with control (by 10% – 0.1 mg/kg, 27% – 0.15 mg/kg). The immobilization time of the 4 groups is less than the control by 49% ($p < 0.05$) and the Stresam group by 18%. Latent immobilization time increased for groups 2, 4 and 5 in 1.4, 2 ($p < 0.05$) and 1.8 times, but for 3 groups it did not change significantly. The number of immobilizations, in comparison with control, for the 4 and 5 groups does not differ, and for the second it increased by 43%, and 32% for the third.

Conclusions. The modified NPY fragment in the light/dark test showed anxiolytic property, less in strength than in Stresam. In a forced swimming test the studied peptide significant showed the ability to delay and reduce the duration of the onset of despair behavior, which may indicate antidepressant-like activity.

STUDY OF THE MODIFIED FRAGMENT OF NEUROPEPTIDE Y INTERACTION WITH COMPOUNDS AFFECTING CENTRAL NERVOUS SYSTEM ACTIVITY

Havrylov I.O., Lytkin D. V., Chumak E. I.

Scientific supervisor: prof. Zagayko A. L.

National University of Pharmacy, Kharkiv, Ukraine

gavrilov.i.ok@gmail.com

Introduction. Neuropeptide Y (NPY) is a biologically active peptide involved in the regulation of many processes in the human body. NPY receptors are widespread both in the brain and in peripheral organs. This determines neuropeptide's ability to affect on various processes such as food consumption, emotional state, vascular tone and so on. The studies were performed with a modified fragment of NPY, which is analogous of C-terminus – that is responsible for receptor binding.

Aim. Investigation of rats' behavioral responses under the conditions of intranasal administration of a modified NPY fragment and compounds that affect central nervous system (CNS) activity.

Materials and methods. The following drugs were used – caffeine-sodium benzoate («Caffeine-sodium benzoate», solution for injection 100 mg/ml, CJSC »Darnitsa Pharmaceutical Company», Ukraine) at a dose of 10 mg/kg, as a compound having psychostimulant and excitatory properties and gidazepam («Gidazepam IC®», tablets 0.05 g. «InterChem» SLC, Ukraine) at a dose of 15 mg/kg, as a compound having anxiolytic and sedative properties at this dose.

The studies were performed on 40 male rats weighing 150-180 g. Animals were divided into 5 equal groups. 1st — control animals that received a solvent (saline) intranasally (i.n.) for 40-45 minutes before the test, and after 20-25 minutes the solvent is intraperitoneal (i.p.). 2nd (caffeine) – animals that received i.n. solvent, and after 20-25 minutes solution of caffeine (10 mg/kg, i.p.). 3rd «caffeine+peptide» – peptide solution (0.1 mg/kg, i.n.) and after 20-25 minutes a solution of caffeine (10 mg/kg, i.p.). 4th (gidazepam) – received a solution of gidazepam 60 minutes before the experiment (15 mg/kg per os), and after 20 minutes the solvent was administrated i.n. 5th – «gidazepam+peptide received» gidazepam (15 mg/kg, per os.), and after 20 minutes peptide solution (0.1 mg/kg, i.n.).