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EFFECTS OF ACUTE HYPOXIA AND HYPER-OXIA ON ANTIOXIDANT DEFENSE SYSTEM IN GIILS OF CARPS (Cyprinus carpio L.)

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In our experiments common carps (Cyprinus carpio L.) were acclimated to an oxygen concentration of 7.31mg/L water (100%) - control (C) group. Two experimental groups (H1, H2) were exposed to acute hypoxia. One of them was exposed to a progressive reduction of O₂ to 4.10mg/L (55%) for 1¼ hours and another 1.95mg/L (25%) for 2 hours, respectively. The fourth experimental group (HP) was exposed to a progressive increase of oxygen concentration O₂ to 10mg/L (150%) for 15 minutes, respectively. In our experiment was determinated antioxidant defense system enzymes activity (superoxide-dismutase (SOD), catalase (CAT) glutathione-peroxidase (GSH-Px), and glutathione-reductase (GR) and glutathione-S-transferase (GST) activity (Misra, H.P. and Fridovich, I. (1972), Beutler (1982), Tamure et al. (1982), Habig et al. (1974), Glatzle et al. (1974).

Our results show that stress caused by acute hypoxia and hyperoxia significantly changes the activities of antioxidant defense system enzymes, superoxide-dismutase (SOD), glutathione-peroxidase (GSH-Px), and glutathione-reductase (GR) in carps gills. Acute prolonged hypoxia (H2) caused a cetrain degree of adaptation in respect to catalase (CAT) and glutathione-Stransferase (GST) activity.

A35.

FUNCTIONAL EXAMINATION OF M1 MUS-CARINIC RECEPTORS IN ISOLATED RAT UTERUS

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Muscarinic receptors subtypes of uterine smooth muscle was examined in many previously researches. In some references we found that there is M2 muscarinic receptors(Pennefather,1994), in some other M4, or simultaneously both M2 and M3 receptors(Eglen, 1989;1991), other authors believe that there are new population of muscarinic receptors undefinitely till now(Boxall at all.1998).

Aim: According to presence of significally functional differences in reacting of myometrium in various species, our investigation in this text will be refer to research to M1 subtype of muscarinic receptors in rat myometrium.

Materials and methods: Experiment was accomplished on isolated rat uterus. Experimental investigation is comprised 18 female rats Wistar race ,average weight of 150-180 gramme divided in 3 groups. Female rats were pretreated with 1mg/kg oestradiol intramusculary,24 hours before experiment. In an organ bath with De Jalon's solution for isolated organ we added:1.in control experimental series (6rats) acetylcholine(acetylcholine hydrochloride) in increasing log concentrations 1.3x10 ⁹M - 4x10⁻⁶M; 2.in second experimental series (6rats) 5 min. before acetylcholine we added pirenzepine, an M1muscarinic recept antagonist(pirenzepine dihydrochloride mononitrate) in concentration of 9x10⁻⁸M;3.in third experimental series (6rats) 5 min. before acetylcholine we added pirenzepine in concentration of $2.7 \times 10^{-7} \text{M};$

Results: In control experimental series, acetylcholine in increasing log concentrations provokes concentration dependent uterus contraction (r= 0.87; P<0.05).ED50 value for acetylcholine is 4.9x10⁻⁸M. In second experimental series acetylcholine in presence of pirenzepine in concentration of 9x10⁻⁸M provokes concentration depending uterus contraction (r=0.86; P, 0.05). ED50 value in this case is 2.3 x10⁻⁸M. In third experimental series acetylcholine in presence of pirenzepine in concentration of 2.7x10⁻⁷M is stronger and more over concentration depending provokes uterus contraction(r=0.91; P<0.01). ED50 value in this case is 9.3x10⁻⁹M.

Conclusion: Our results of experiment showed that the effect of acetylcholine in presence of pirenzepine was potented. It means that the effect of acetylcholine in presence of pirenzepine in concentration of 9x10⁻⁸M is twice, and in presence of 2,7x10⁻⁷M pirenzepine concentration is quintuple strongly than this in control series. The data suggest that population of M1 subtypes of muscarinic receptors exist in the rat uterus, and they are involved in regulation of miometrial contractility.

A36.

IMMUNE BLOOD KILLER CELLS. THEIR ORI-GIN AND SIGNIFICANCE

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Killer cells (K cells) play a very important role in immune defense, especially in mechanisms of none-specific antitumor immunity, which emphasize their therapeutic implications. They represent probably the most enigmatic feature of the non-specific antitumor immunity. The exact origin of this cell population is still under extensive investigation.