INFLUENCE OF CHRONIC EXPOSURE TO WEAK VARIABLE MAGNETIC FIELD ON ANTIOXIDANT ACTIVITY IN RATS WITH EXPERIMENTAL INFLAMMATION. G. Cieslar¹, J. Mrowiec¹, J. Zalejska-Fiolka², E. Birkner², S. Kasperczyk², A. Sieron¹. ¹Chair and Clinic of Internal Diseases, Angiology and Physical Medicine, Silesian Medical Univ, PL-41902 Bytom, Poland, ²Chair and Dept of Biochemistry, Silesian Medical Univ, PL-41808 Zabrze, Poland.

Objectives The aim of the study was to estimate the influence of chronic exposure to weak variable magnetic field used in magneto-stimulation on activity of some antioxidant enzymes in rats with experimental inflammation. Methods: Experimental material consisted of 128 male Wistar rats weighting 180-200 g. Weak variable magnetic field of saw-like shape of impulse, at a frequency of basic impulse 180-195 Hz and induction of 60 µT generated by device for magneto-stimulation Viofor JPS (Poland) basing on ion cyclotron resonance phenomenon was used. All animals were randomly divided into 4 groups (32 animals each). In first group whole body exposure to magnetic field lasting 36 minutes daily for 14 consecutive days was made. In second - control group sham-exposure without generating magnetic field inside of applicator lasting 36 minutes daily for 14 consecutive days was made. Rats in third group were injected with 50 ul of 5% solution of formaldehyde in region of right hip and after 24 hours were subjected to the same exposure cycle as in first group. The animals in fourth - control group were also injected with 50 µl of 5% solution of formaldehyde in region of right hip and then after 24 hours were subjected to the same sham-exposure cycle as in second group. In all groups at 7th and 14th day of repeated exposures or sham-exposures and at 7th and 14th day after the end of a cycle of exposures every time a part of animals (8 rats from each group) was exsanguinated in ether narcosis. In obtained blood and hemolysates of erythrocytes samples contents of some antioxidant activity indicators: activity of catalase (CAT), glutathione peroxidase (GPX) and superoxide dysmutase (SOD) in erythrocytes, activity of isoenzymes of superoxide dysmutase (Mn-SOD and ZnCu-SOD)) in serum as well as serum concentration of malondialdehyde (MDA) was determined by means of spectrophotometric and kinetic methods. In the statistical evaluation ANOVA analysis with subsequent post-hoc Mann-Whitney's U test were used. Summary of results: The activities of antioxidant enzymes as well as serum malondialdehyde concentration in particular groups of rats are presented in table 1. In magnetic field-exposed group a significant decrease in activity of most of analyzed antioxidant enzymes both in erythrocytes and serum during exposure cycle was observed as compared to a group of rats with experimental inflammation, in which these activities were significantly increased comparing to control group. Besides in both magnetic field-exposed groups a significant decrease in malondialdehyde serum concentration during exposure cycle was obtained. Conclusion: Chronic exposure to weak variable magnetic field used in magnetostimulation basing on magnetic resonance phenomenon causes a beneficial effect antioxidant reactions in course of experimental inflammation in living organisms.

| | ivity of some antioxidate in all groups of rats /td> | | | | |
|-----------------------------|--|----------------------------------|----------|-----|--|
| Parameter | Group | Day of exposure or sham-exposure | | | |
| | | 7 day of exposure cycle | avnocura | 1 - | he 14 day after the of end of exposure cycle |
| Activity of CA erythrocytes | T in | • | | · | • |

| [IU/mgHb] | Control | 197,3 | 168,9 | 152,0 | 158,4 |
|--|-------------------------------|---------|---------|---------|---------|
| | Inflammation | 142,4** | 117,8** | 211,1 | 102,9** |
| | Magnetic field | 178,8 | 112,3** | 105,5* | 88,0** |
| | Magnetic field + inflammation | 177,9 | 118,5** | 115,3** | 97,6** |
| Activity of GPX in erythrocytes [IU/gHb] | Control | 139,7 | 79,6 | 118,2 | 60,2 |
| | Inflammation | 271,2** | 164,4** | 138,3 | 73,2 |
| | Magnetic field | 141,3 | 107,0* | 103,1 | 118,7** |
| | Magnetic field + inflammation | 102,6 | 117,6** | 118,2 | 138,2** |
| Activity of SOD in erythrocytes | | | | | |
| [NU/gHB] | Control | 135,7 | 132,5 | 122,0 | 285,6 |
| | Inflammation | 117,3 | 99,6 | 185,9 | 141,4** |
| | Magnetic field | 97,1 | 169,9 | 220,7* | 172,5** |
| | Magnetic field + inflammation | 55,9 | 168,6 | 220,9* | 152,1** |
| Activity of Mn-SOD in serum [NU/ml] | Control | 8,2 | 12,0 | 13,2 | 7,7 |
| | Inflammation | 5,8* | 9,3 | 3,2** | 4,4 |
| | Magnetic field | 5,4* | 8,6 | 3,6** | 4,9 |
| | Magnetic field + inflammation | 3,4* | 5,7** | 3,0** | 4,4 |
| Activity of ZnCu- SOD in serum [NU/ml] | Control | 19,2 | 15,4 | 18,4 | 21,7 |
| | Inflammation | 24,3* | 20,4 | 24,7* | 22,5 |
| | Magnetic field | 21,9 | 21,9 | 26,1* | 26,1* |
| | Magnetic field + inflammation | 20,0 | 22,0 | 27,5* | 28,6* |
| Concentration of MDA [fÝmol/l] | Control | 6,3 | 6,2 | 4,1 | 5,0 |
| | Inflammation | 4,9 | 4,3 | 3,7 | 3,4* |
| | Magnetic field | 3,0** | 2,9** | 3,3 | 4,6 |
| | Magnetic field + inflammation | 3,9** | 2,9** | 3,5 | 4,6 |