## 8TH INTERNATIONAL SYMPOSIUM ON ALS/MND

ABSTRACT TITLE:

## CONTENT OF NO REACTIVE -SH AND SUPEROXIDE DISMUTASE ACTIVITY IN CSF OF ALS PATIENTS

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The discovery of mutations in the CuZn superoxide dismutase (SOD) gene in familial ALS implicates oxidative stress in the pathogenesis of this disorder. Recent findings have shown that nitric oxide (NO) is not produced from nitric oxide synthase (NOS) without cooperative action of NOS and SOD, give an additional explanation for initiation of free radical chain reactions. It is possible that in case that not enough NO is produced in motor neurons other reactive species, nitrous oxide  $(N_2O)$  and hydoxylamine  $(NH_2OH)$ , and an excess of reactive oxygen species (ROS), such as hydrogen peroxide  $(H_2O_2)$ , can delay initiation of free radical chain reactions until capacity of antioxidative defense capacity is restrained due to inhibition of its components.

The aim of this study was to investigate the content of -SH groups and SOD activity in cerebrospinal fluid (CSF) both in ALS patients and control group, as well as to perform an *in vitro* test for CSF ·NO binding capacity, in same examines. Nitric oxide for *in vitro* testing was provided by adding 40% sodium nitrite solution in the solution of 10% ferrous sulfate in 50% sulfuric acid.

Our results showed significant increase -SH content in CSF of ALS patients in comparison with control group ( $114 \pm 29$  nmol -SH/mg proteins in CSF of ALS patients compared with  $37 \pm 12$  nmol -SH/mg proteins in CSF of controls)(p<0.05). After *in vitro* saturation of CSF with NO, we noted the significant decrease of -SH content which was not found in control group. We also found that SOD activity in CSF of ALS patients was higher than in control group ( $16.1 \pm 2.8$  U/ mg CSF proteins of ALS patients compared to  $10.2 \pm 2$  U/ mg CSF proteins of control group).

Submission for	Platform presentation (20 min /12 stokes)	
	Communication (10 min. platform presentation/6 slides)	
	Poster presentation	

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