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Dr. Liljana Ilievska, Skopje

Zagreb, Mart 19, 2006

Dear Dr Ilievska,

We thank you submitting your very interesting paper:

L Ilievska, E Ilievska (Skopje):

EVOKED POTENTIALS IN EVALUATION OF SOLCOSERYL (ACTIHAMYL) EFECTS IN THE TREATMENT OF CEREBRAL ATROPHIC PROCESSIES

for presentation in the Program of the 46th International Neuropsychiatric Pula Symposium.

I am glad to confirm, that the paper has been reviewed and accepted for publication in the Proceedings and for presentation in the PROGRAM of our Symposium. Your paper has to be presented at the Thursday, June 17, 2006, 15:00 - 16:00.

The relevant information on the Oral presentation you can find on our Web Site:

www.pula-symp.com

Kind regards, looking forward to see you in Pula!

Yours sincerely,

Boško Barac, Secretary General INPS, Coordinating Editor of the INPS "Proceedings"

EVOKED POTENTIALS IN EVALUATION OF SOLCOSERYL (ACTIHAMYL) EFECTS IN THE TREATMENT OF CEREBRAL ATROPHIC PROCESSES

L. Hievska, E. Hievska Clinic of Neurology, Clinical Centre, Skopje, R. Macedonia

Twenty-five patients with cerebral atrophic processes, either local or diffuse, of various etiology, proved by computerized tomography were investigated and treated. Eleven patiens, beside other symptoms, also had epiteptic seizures. The youngest patient was 43-year-old, the oldest 62, with mean age of 59 years.

All patients underwere psychologic testing prior to and after therapy with Actihamyl (Solcoseryl), and the ossults showed different degrees of mental function deficits. Also, EEG and evoked potential (visual and somatosensory) recordings were made in several instances, showing to be a good indicator in objaectivization of the Actihamyl action on cerebral metabolism.

Concerning the therapy, the patients administering this medication were divided into two groups:

- The first one (16 patients) adminsitered amp. Actihamy! within a period of six weeks:
- The socond one (9 patiens) who were examined by means of EVP after i.v. injection of Actihamil, when the acute effect was tested.

The contol group consisted of 16 patients with no neurologic disease, adminsitering placebo instead of a drag. There were no subjective or objective effects shown by means of EEGs and EVPs recording.