



EFFECTS OF VITAMIN D PRETREATMENT ON THE NOX2, REDOX BALANCE AND NEURONAL MORPHOLOGY IN GERBILS EXPOSED TO TRANSIENT GLOBAL BRAIN ISCHEMIA

N. Petronijevic, M. Velimirovic, G. Jevtic Dozudic, T. Stojkovic, T. Nikolic, M. Zivkovic, N. Puskas, A. Mircic, I. Zaletel, V. Selakovic

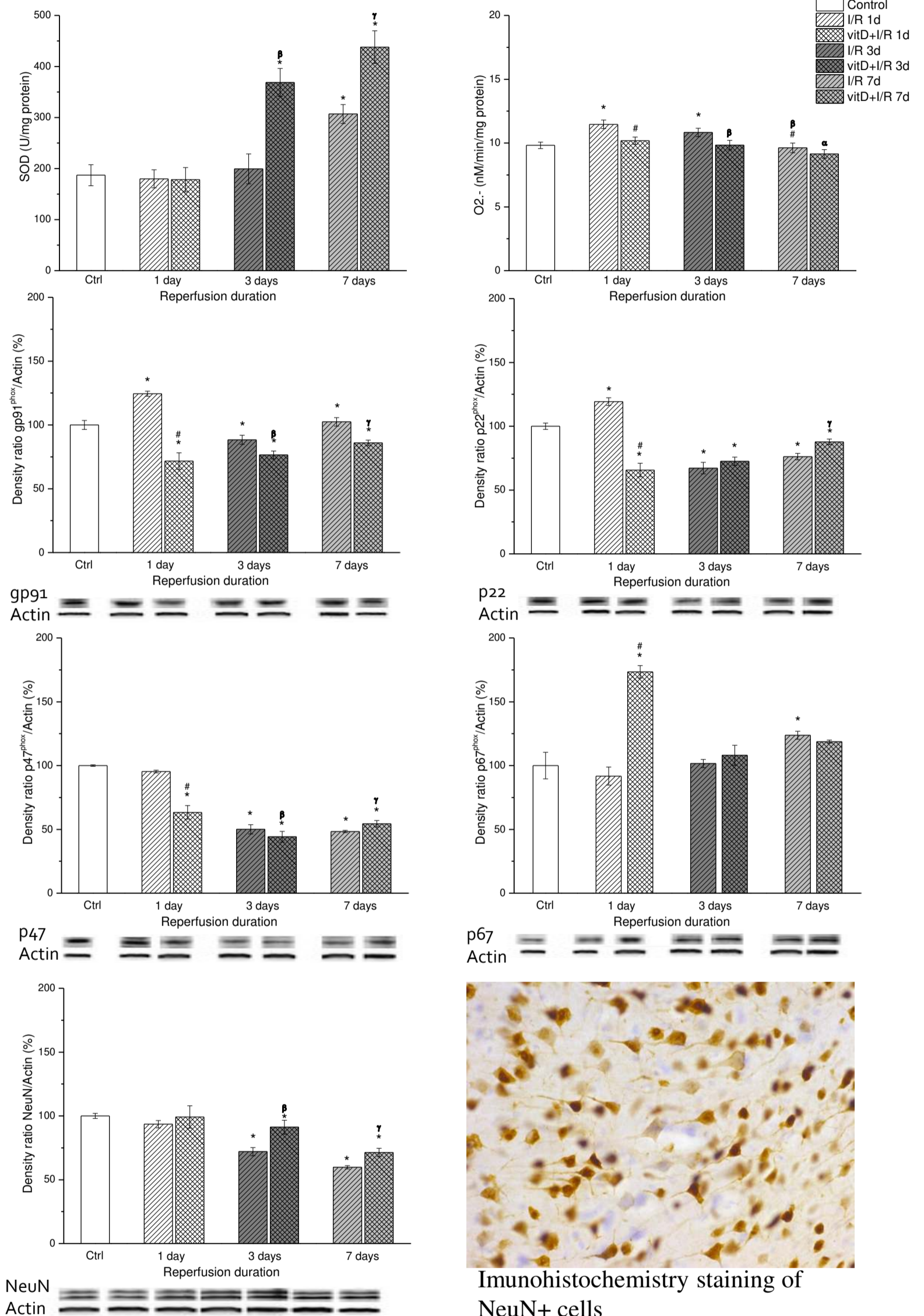
Introduction: Cerebral ischemia and reperfusion (I/R) are followed by the production of free radicals. The source could be NADPH oxidase isoform NOX2. Vitamin D is known to be neuroprotective.

The aim: Investigation of the effects of vitamin D pretreatment on the superoxide anion ($O_2^{\cdot-}$) production, superoxide dismutase (SOD) activity, expression of NOX2 and NeuN and morphology of neurons in the hippocampus of gerbils exposed to transient global brain ischemia.

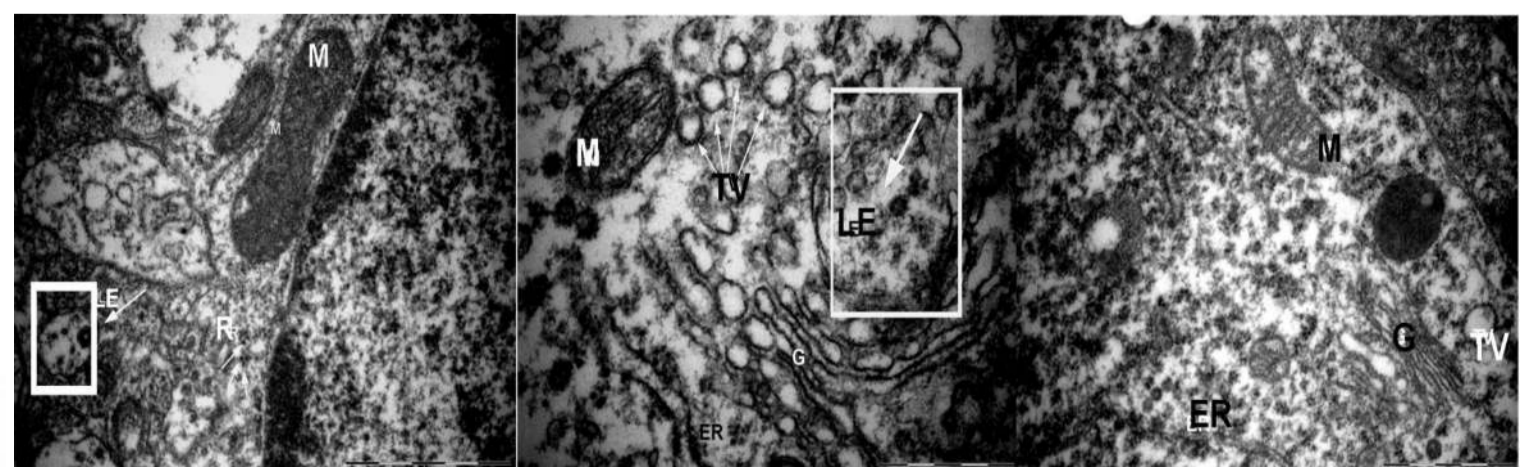
Material and methods: Gerbils were divided into seven groups: control, groups I/R1, I/R3 and I/R7 subjected to 10 minutes ischemia (ligation of both common carotid arteries) and reperfusion for 1, 3 or 7 days respectively, and groups vitD+I/R1, vitD+I/R3 and vitD+I/R7 that were pretreated with vitamin D seven days prior to corresponding I/R. Content of $O_2^{\cdot-}$ and SOD activity were determined by spectrophotometry. NOX2 and NeuN expression were determined by Western blot and immunohistochemistry and neuron morphology by transmission electron microscopy (TEM).

Conclusion: Vitamin D regulates production of $O_2^{\cdot-}$ and SOD activity after I/R and ameliorates effects on the expression of NOX2. Vitamin D protects cell structure in early phases of I/R injury.

Results



* p<0,05 compare to control group # p<0,05 compare to I/R 1d group
β p<0,05 compare to I/R 3d group γ p<0,05 compare to I/R 7d group



The TEM analysis of neurons. (A) Neuronal cell from control group shows late endosomes (LE) with intact membrane, ribosomes (R) and mitochondria (M) (bar = 1μm). (B) A Neuron after I/R showing late endosome with membrane rupture, accumulation of transport vesicles (TV) and more ribosomes (Bar = 0.5μm). (C) Neuron cell after vitD+I/R procedure showing intact endosome, some vesicles, Goldgi apparatus (G) and rough endoplasmic reticulum (ER) (Bar = 1μm).