

**Publication****Acute stress induced modifications of calcium signaling in learned help-  
less rats****JournalArticle (Originalarbeit in einer wissenschaftlichen Zeitschrift)****ID** 107231**Author(s)** Velbinger, K; De Vry, J; Jentsch, K; Eckert, A; Henn, F; Müller, W E**Author(s) at UniBasel** [Eckert, Anne](#) ;**Year** 2000**Title** Acute stress induced modifications of calcium signaling in learned helpless rats**Journal** Pharmacopsychiatry**Volume** 33**Number** 4**Pages / Article-Number** 132-7

Previous reports have demonstrated reduced elevations of free intracellular calcium concentration in blood cells of depressed patients after various stimuli. Therefore, a disturbance of intracellular calcium (Ca<sup>2+</sup>) homeostasis has been postulated to be involved in the pathophysiology of mood disorders. It was the aim of the present study to investigate whether Ca<sup>2+</sup> signaling was affected in spleen T-lymphocytes of rats submitted to a learned helplessness paradigm, an animal model of depression with a high level of construct, face and predictive validity. In addition, we tested for effects of acute stress on the Ca<sup>2+</sup> signaling in helpless rats, as compared to non-stressed rats. It was found that mitogen-induced Ca<sup>2+</sup> signaling only tended to be reduced in helpless rats. However, when helpless rats were submitted to acute immobilization stress, Ca<sup>2+</sup> signaling appeared to be significantly blunted, whereas the same stressor did not affect Ca<sup>2+</sup> signaling in the non-helpless control rats. These acute stress-induced differences in Ca<sup>2+</sup> signaling were not paralleled by a differential increase in plasma corticosterone. It is hypothesized that blunted Ca<sup>2+</sup> signaling, as assessed in spleen T-lymphocytes of helpless rats, may be a correlate of the increased vulnerability of helpless rats to acute stressors.

**Publisher** Thieme**ISSN/ISBN** 0720-4280**edoc-URL** <http://edoc.unibas.ch/dok/A5253498>**Full Text on edoc** No;**Digital Object Identifier DOI** 10.1055/s-2000-11220**PubMed ID** <http://www.ncbi.nlm.nih.gov/pubmed/10958261>**ISI-Number** WOS:000088643700002**Document type (ISI)** Journal Article