Title: Intracerebroventricular injection of oxidant and antioxidant molecules affects long-term potentiation in urethane anaesthetized rats.

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Short title: ICV injection of oxidant and antioxidant molecules: effects on LTP
Production of superoxide anions in the incubation medium of hippocampal slices can induce long-term potentiation (LTP). Other reactive oxygen species (ROS) such as hydrogen peroxide are able to modulate LTP and are likely to be involved in aging mechanisms.

The present study explored whether intracerebroventricular injection of oxidant or antioxidant molecules could affect LTP in vivo. To this end, field excitatory post-synaptic potentials (fEPSPs) elicited by stimulation of the perforant pathway were recorded in the dentate gyrus of the hippocampal formation in urethane anaesthetized rats. N-Acetyl-L-cysteine, hydrogen peroxide (H$_2$O$_2$) or hypoxanthine/xanthine-oxidase solution (a superoxide producing system) were administrated by intracerebroventricular (ICV) injection. The control was represented by a group injected with saline ICV. Ten min after the injection, LTP was induced in the granule cells of the dentate gyrus (DG) by high frequency stimulation of the perforant pathway.

Neither the H$_2$O$_2$ injection or the N-Acetyl-L-cysteine injection caused any variation in the fEPSP at the 10-minute post-injection time point whereas the superoxide generating system caused a significant increase in the fEPSP. Moreover, at 60 min from tetanic stimulation, all treatments attenuated LTP compared with the control group.

These results show that ICV administration of oxidant or antioxidant molecules can modulate in vivo the LTP in the DG. Particularly, a superoxide producing system can induce potentiation of synaptic response. Interestingly, oxidants or antioxidants ICV injection prevented full expression of LTP compared to saline injection.
Main Body of Text

Reactive oxygen species (ROS) play a normal role as signalling molecules (Suzuki et al. 1997) and growth or pruning of the synaptic spine could be controlled in part by the balance in the synapse between neurodestructive pro-oxidants and neuroprotective antioxidants (Smythies 1999). In hippocampal slices, the induction of long-term potentiation (LTP) in CA1 region (Kamsler and Segal 2003a, Knapp and Klann 2002b) can be modulated by hydrogen peroxide and the increase or decrease of the potentiation is linked to the hydrogen peroxide (H$_2$O$_2$) concentration in the perfusion medium (Auerbach and Segal 1997, Kamsler and Segal 2003a, Katsuki et al. 1997, Pellmar et al. 1991). Moreover, peroxide scavengers (Knapp and Klann 2002a) and superoxide scavengers (Knapp et al. 1998, Klann 1998) can prevent full expression of LTP possibly by sequestration of superoxide anions. Notably, superoxide generation by xanthine oxidase activity can induce LTP in CA1 region in hippocampal slices (Knapp and Klann 2002a).

In vivo evidence shows that dietary manipulation with antioxidants and stress condition can also influence LTP (Alfarez et al. 2003, McGahon et al. 1999a, McGahon et al. 1999b, Shakesby et al. 2002, Vereker et al. 2001). Furthermore, reactive oxygen species (ROS) are possibly related to LTP impairment with aging and this process may be reversed by antioxidant-enriched diet (McGahon et al. 1999a, McGahon et al. 1999b, Murray and Lynch 1998, O'Donnell et al. 2000, Viggiano et al. 2006).

To advance in the understanding of the molecular mechanisms underlying LTP, it would be of great interest to support the results obtained from slices with an in vivo whole brain-model. The aim of the present study was to test the effects of ICV
injections of oxidant or antioxidant molecules on LTP induced in dentate gyrus (DG) in anaesthetized rats. We used male Sprague–Dawley rats weighing 250–300 g and housed at a controlled temperature of 22 ± 1 °C and humidity of 70% with a 12/12 h light–dark cycle from 07:00 to 19:00 hours. Laboratory standard food (Mil, Morini, Italy) and water were available at all times. All other parameters fulfilled the requirements of the ‘Guide for the Care and Use of Laboratory Animals’ by the National Research Council, implemented by EU and local rules.

Rats were anesthetized with 1g/kg urethane and placed in a stereotaxic apparatus with lambda 1 mm below bregma. The body temperature was monitored by a rectal thermometer and maintained at 37.0 ± 0.2 °C by an electrically shielded heating pad. In order to record evoked potentials in the hilus of DG, two small holes were drilled into the bone to reach the perforant pathway (pp) (7.5 mm posterior to the bregma, 4.2 mm lateral to the midline) with a unipolar stimulating electrode and the DG (3.5 mm posterior to the bregma, 2.5 mm lateral to the midline) with the recording electrode (Paxinos and Watson 1997). Two screws in the occipital bone were used as reference and ground. A third hole was made to insert a cannula into the lateral ventricle (0.4 mm posterior to the bregma, 1.7 mm lateral to the midline, 3.3 mm from the cranial theca). Recording and stimulating electrodes as cannula were always on the same side of the brain.

Stimulation and recording was achieved with a modular analog instrumentation (Neurolog, Digitimer, England) connected to a PC with a A/D-D/A converter (AT-MIO-XE50, National Instruments, Tx USA) to automatically follow all the stimulating/recording protocol with a custom software written under LabView.
environment (National Instruments).

The depth of recording and stimulating electrodes was optimized to maximize the amplitude of fEPSP evoked by a perforant path test shock.

For eliciting fEPSP, a series of eight monopolar pulses were generated with a frequency of 0.1 Hz. The pulse width was 100 µs and the stimulus intensity (200-400 µA) was adjusted to the value that evoked 50% of the maximum fEPSP amplitude. Amplitude (measured at 3.0 ms from the stimulus artefact; maximal amplitude never occurred before 3.8 ms) and slope (on the first millisecond of the rising phase) of the fEPSP were used to quantify the evoked potentials.

LTP was induced by stimulating the pp with 20 trains of 15 impulses with the same weight and amplitude as the test pulse. The frequency within the train was 200 Hz, and the distance between the trains was 5 s (Krug et al. 2001). Examples of fEPSP before and after LTP with measured parameters are shown in fig. 1.

Four groups of five animals each were subjected to the following procedure: 30 min recovery, 8-fEPSP (0.1 Hz) sampling, 5 µl intracerebroventricular (ICV) injection of one of the solutions described below, 10 min elapsed time, 8-fEPSP (0.1 Hz) sampling, LTP induction, 8-fEPSP (0.1 Hz) sampling after 60-min from the high frequency stimulation.

The solutions were: saline (group 1), N-Acetyl-L-cysteine (NAC) (100mM) (group 2), hydrogen peroxide (8.8 mM) (group 3), and hypoxanthine (1 mM)/xanthine-oxidase (0.82 U/ml) solution (a superoxide anions generating system) (group 4). The solutions were always prepared immediately before the injection.

At the end of the experimental session, animals were overdosed with urethane and perfused through the heart with 0.9% saline followed by 10% formalin in PBS. All
electrode sites were histologically verified in 50 µm frozen sections stained with cresyl violet.

One-way analysis of variance (ANOVA) was used to test significant differences between the experimental conditions. Paired t test was used to compare pre-treatment state with post-treatment state. Rejection level was fixed at p=0.05.

Fig. 2 shows percent of variation in fEPSP after 10 min from injection and after 60 min after high frequency stimulation displays the summary of the fEPSP data. The values are means ± SEM of five observations in each case.

The injection of hypoxanthine/xanthine-oxidase caused a significant increase of the fEPSP amplitude (112.8 ± 2.8) and slope (115.4 ± 3.8) [P < 0.01; paired t test]. On the other hand, ICV injection of saline, hydrogen peroxide (8.8 mM) or NAC did not affect the fEPSP amplitude and slope.

60 min after high frequency stimulation, significant differences appeared in fEPSP amplitude [F(3, 16) = 14.69; P < 0.01 one way ANOVA] and in fEPSP slope [F(3, 16) = 7.87; P < 0.01; one way ANOVA]. Unplanned pair-wise comparisons with Tukey’s method revealed that the potentiation in all the experimental groups were significantly lower than in the control group [saline group (amplitude = 207.0 ± 13.8; slope = 195.1 ± 11.4)] but there was no significant difference among the three experimental conditions [H₂O₂ group (amplitude = 131.0 ± 5.9; slope = 146.7 ± 7.6); the superoxide group (133.9 ± 11.0; 148.3 ± 7.0); NAC group (125.8 ± 7.7; 147.7 ± 7.1)].

These results show that ICV injection of oxidant or antioxidants molecule can modulate the synaptic transmission in DG in vivo. Hypoxanthine/xanthine-oxidase system (a system able to produce superoxide anions (Frederiks and Bosch 1997)) was able to
induce potentiation in DG while hydrogen peroxide and the antioxidant NAC attenuated LTP. Similar results were obtained in vivo by Vereker et al. (O’Donnell et al. 2000, Vereker et al. 2001) who showed that hydrogen peroxide is able to inhibit LTP induction (Vereker et al. 2001). These authors also suggested protective effects of antioxidants on aging and stress impairment in LTP, because the addition of antioxidants to the standard laboratory diet reverses the age-related deficit in LTP (Martin et al. 2002, McGahon et al. 1999a, McGahon et al. 1999b, Vereker et al. 2001). Our results show that ICV injection of an antioxidant species (NAC) has a deleterious effect on LTP. This result agrees with the evidence that antioxidants have inhibitory effects on LTP in vitro (Klann et al. 1998, Klann 1998, Thiels et al. 2000) and that the superoxide production is needed for LTP in CA1 in hippocampal slices (Knapp and Klann 2002a). Moreover, our results show that even if hypoxanthine/xanthine-oxidase system was able to induce potentiation, its effect on LTP induced by tetanic stimulation was deleterious because the overall potentiation after 60 min from tetanus was significantly smaller than in the control. It seems reasonable that superoxide produced by exogenous xanthine-oxidase activity leads to production of hydrogen peroxide with deleterious effects for LTP (Knapp and Klann 2002b), but this hypothesis needs further studies. The inhibitory effect of hydrogen peroxide on LTP is well known and Kamsler et al. proposed a dual role for H₂O₂ with high concentrations suppressing LTP and low concentration enhancing it (Kamsler and Segal 2003a, Kamsler and Segal 2003b). In our model, the H₂O₂ injection did not affect fEPSP slope and amplitude as in the in vitro slices preparation but had a considerable negative effect on the LTP induction by tetanic stimulation. These differences between in vivo model and in vitro slices preparation are not explainable with present data and also need further evaluations, nonetheless they
stress the importance of validating and corroborating *in vitro* data with *in vivo* models. Some observations should be done with regards to the ICV injection and anaesthesia. It was not possible to determine the distribution of the injected substances through the brain parenchyma therefore it was not possible to evaluate the effective concentration of the oxidants or antioxidants at the level of DG in the present model. It is also known that, under urethane anaesthesia, induction of LTP requires stronger tetanization protocol than in freely moving rats (Riedel *et al.* 1994). Therefore it could be interesting evaluating the effects of oxidants and antioxidants in freely moving rats.

Two limits of the present study should be pointed out. The first one is the absence of a dose-response curve that will be obtained by future experiments. In fact, it seems important to investigate about the effect of different ICV oxidants and antioxidants doses on LTP considering that H$_2$O$_2$ modulation of LTP depends on concentration (Kamsler and Segal 2003a, Kamsler and Segal 2003b). The second limit is related to the lack of a complete time course for the potentiation. This protocol was followed by the authors to simplify the design of the experimental session. Again, further studies are planned to obtain a complete time course for the effects of ROS ICV injection on LTP.

The principal finding of this study is that the ICV injection of oxidant and antioxidant molecules is able to modulate LTP in DG during acute recording *in vivo*. With the discussed limits, this finding supports and extends the validity of previous data obtained from *in vitro* slices preparation.
References


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Figure legend

Figure 1

Example of fEPSP before and after LTP induction. Arrow marks the stimulus artifact, grey square marks the portion of fEPSP used in evaluating slope (1 ms). At fixed latency (3.0 ms) from the start of stimulus artifact, amplitude of fEPSP was calculated. Calibration bars: 2 mV; 4 ms.
Figure 2

Percent of variation in fEPSP after 10 min from ICV injection of 5 µl of saline, hydrogen peroxide (8.8 mM), hypoxanthine (1mM)/xanthine oxidase (0.82 U/ml) (superoxide producing system) or N-Acetyl-L-cysteine (100 mM) and after 60 min after high frequency stimulation in the same groups. The values are means ± SEM of five observations in each case. Asterics mark significant differences with respect to saline group. See text for details.