Sulfamethoxazole induces zinc changes at hippocampal mossy fiber synapses from pregnant rats

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Abstract. The accumulation of intracellular ionic zinc and pharmaceutical compounds, like the antibiotic sulfamethoxazole, may contribute to various neuropathologies. Sulfamethoxazole and the drug trimethoprim, are inhibitors of enzymes involved in the synthesis of tetrahydrofolate and also of carbonic anhydrases. The inhibition of the latter enzymes, which are localized both intra- and extracellularly and have a key role in pH regulation, causes alkalinization that is associated with higher spontaneous transmitter release. Intense synaptic stimulation causes the entry of released zinc into postsynaptic neurons, through glutamate receptor channels or voltage dependent calcium channels. The aim of this study was to evaluate the effect of sulfamethoxazole (180 µM) on basal postsynaptic zinc and to compare it with that caused by two depolarizing media, containing high potassium or tetraethylammonium, which may induce long term synaptic plasticity. The studies were performed in brain slices from gestating rats, at the mossy fiber synapses from hippocampal CA3 area, using the zinc indicator Newport Green. In the presence of KCl (20 mM) and sulfamethoxazole (180 µM) the zinc signals were enhanced, unlike in tetraethylammonium (25 mM). After sulfamethoxazole the tetraethylammonium evoked zinc signal had reduced amplitude. Thus, the data suggests that sulfamethoxazole enhances transmitter release affecting synaptic zinc physiology.

Keywords: Sulfamethoxazole — Synaptic zinc — KCl and TEA evoked depolarizations — Newport Green

Introduction

Antibiotics are frequently used on humans and animals in order to treat various bacterial infections. Disposal of unused pharmaceuticals and by-products is an important source of pharmaceutical pollution, which can affect groundwater and potentially drinking water (Kummerer 2003). The presence of antibiotics in the environment can modify the ecosystems and may favour the development of resistant bacteria (Boreen et al. 2004). The conventional wastewater treatments are not designed to remove pharmaceutical waste like antibiotics and therefore they are released in the environment, having been detected in surface waters at concentrations that reach values near 4 µM (Heberer 2002). Sulfamethoxazole (SMX) is a widely-used bacteriostatic antibiotic often prescribed along with the antibiotic trimethoprim that interferes with the production of dihydrofolate. SMX is used to treat various infections, such as venereal, gastrointestinal and respiratory infections (Kiellhofner 2005). It was observed that about 15% of the compound is excreted by the human body without being metabolically degraded (Hirsch et al. 1999; Schmitt et al. 2005). This antibiotic has been detected at concentrations of 70–150 ng/l (0.3–0.6 nM) and 200–2000 ng/l (0.8–8 nM)
in surface water and secondary wastewater effluents, respectively (Dodd and Huang 2004). It has been reported that the application of SMX can be associated, in very rare cases, with encephalopathy and psychosis, essentially in immune-compromised or elderly patients (Saidinejad et al. 2005). The neurotoxic effects result from the excellent penetration in the central nervous system, however, the exact mechanism of neurotoxicity is unknown (Grill and Maganti 2011).

SMX belongs to a generation of antibiotics, that includes the sulfa drugs, which have well known mechanisms of action and inhibit three types of enzymes. Two of them, dihydropteroate synthase and dihydrofolate reductase, are involved in the synthesis of tetrahydrofolate (Caspasso and Supuran 2014). The other type consists of the carbonic anhydrases (CA), found in all living organisms, which catalyze the reaction between carbon dioxide and water resulting in the formation of bicarbonate and protons (Supuran 2008). Thus, they are important intra- and extracellular components of the brain pH buffering system (Obara et al. 2008; Casey et al. 2010), their inhibitors being clinically used as anticonvulsant drugs (Thiry et al. 2008). It is normally considered that alkalization causes enhanced neuronal activity and synaptic transmission while acidification leads to lower neuronal response (Sinning and Hubner 2013). Synaptic vesicles become highly acidic during neurotransmitter loading, by the action of the vacuolar H⁺-ATPase, leading vesicle fusion to the insertion of these proton pumps in the presynaptic membrane (Jefferies et al. 2008). Thus, during synaptic activity, exocytosis involves the release of protons, accompanying transmitter release, causing a short but large acidification of the cleft medium, followed by a long and transient extrasynaptic alkalization (Sinning and Hubner 2013). In hippocampal synapses, this increase in extracellular pH, that can be up to 0.1–0.2 pH units, may occur in milliseconds. In the case of intense stimulation, and thus higher cellular metabolism, the cleft acidification may last for a longer period. At the Schaffer collaterals-CA1 pyramidal cell synapses, the specific CA inhibitor acetazolamide (10 mM), a permeant sulfonamide derivative, causes a large extracellular pH increase without changes in the corresponding electrophysiological responses, measured as field potentials (Chen and Chesler 1992). In the same work, using an inhibitor of extracellular CA, the membrane impermeant dextran bound sulfonamide (DBSA), similar pH enhancements were observed, having been proposed that they are due to postsynaptic H⁺ entry through cation channels (Chen and Chesler 1992), following the pH dependent activation of NMDA receptors (Sinning and Hubner 2013).

At glutamatergic synapses spontaneous transmitter release is diminished at lower intracellular pH, as found in experiments with the Na⁺-driven Cl⁻/HCO₃⁻ exchanger Slc4a8, an acid extruder mainly localized to nerve terminals, and is enhanced at higher internal pH (Sinning et al. 2011). On the postsynaptic site, acid extrusion is mediated by the electroneutral Na⁺/HCO₃⁻ co-transporter Slc4a7, whose expression increases during glutamate excitotoxicity (Sinning and Hubner 2013). Since SMX is cell-permeant and inhibits carbonic anhydrases that mediate rapid pH buffering, its application should cause an increase in intracellular pH and, consequently, higher glutamate and zinc release.

One of the most important features of the central nervous system is plasticity, which is mainly triggered by specific forms of synaptic activity. In particular, long-term potentiation (LTP), is considered to mediate learning and the formation of cellular memory (Malenka and Bear 2004; Nicoll and Schmitz 2005; Bliss and Collingridge 2013). An important form of LTP is expressed at the zinc enriched mossy fiber-CA3 pyramidal cell synapses, of hippocampal CA3 area, where zinc is co-released with glutamate during synaptic activity (Weiss et al. 2000; Li et al. 2001; Cho et al. 2003; Quinta-Ferreira et al. 2004; Frederickson et al. 2005; Ketterman and Li 2008; Paoletti et al. 2009; Khan et al. 2014). The hippocampal mossy fiber boutons include numerous vesicles that contain high concentrations of free or loosely-bound zinc (Wenzel et al. 1997; Frederickson et al. 2000; Hallermann et al. 2003; Rollenhagen and Lübbe 2010). It has been shown that intense zinc release may lead to neurotoxicity and neuronal disorders (Choi and Koh 1998; Frederickson et al. 2005; Sensi et al. 2011) the effect being evoked by zinc entry into neurons through voltage-dependent calcium channels (VDCCs) or via AMPA, kainate and NMDA receptor channels (Sensi et al. 1997; Marin et al. 2000). LTP can be evoked by intense electrical or chemical stimulation, in the latter case, through the application of extracellular potassium (Bernard et al. 1994; Roisin et al. 1997), tetrathyrammonium (TEA) or 4-aminoypyridine (Bancila et al. 2004; Suzuki and Okada 2009). Cell depolarization evoked by external potassium that can stimulate simultaneously all mossy fiber synapses (Zhao et al. 2012), and/or high external zinc, combined or not with potassium, evoked intracellular zinc enhancements in cultured cortical neurons and in hippocampal slices (Sensi et al. 1997; Marin et al. 2000; Ketterman and Li 2008). On the other hand, it has been shown that TEA evokes mossy fiber LTP causing also an inhibition of synaptic activity due to zinc binding to presynaptic K_ATP channels (Bancila et al. 2004; Suzuki and Okada 2009). Their activation by zinc may lead to cell hyperpolarization and thus to the subsequent reduction of neurotransmitter and zinc discharge, that should result in a decrease of postsynaptic zinc in agreement with our previous observations (Bastos et al. 2017a). With respect to the synaptic action of SMX, there is still no available data. The aim of this work was to investigate the effect of SMX on basal mossy fiber synaptic zinc signals, comparing it with that induced by KCl, and
also by TEA. For this purpose, we have recorded fluorescence signals from hippocampal slices, obtained from the brain of pregnant gestating rats, with 16 to 18 days of gestation, whose fetuses were used for other experiments. It is well known the existence of numerous adaptations in the mother’s brain towards the end of pregnancy gestation, especially in the regions containing the “maternal circuitry” that participates in the control of behavior (Numan 2007). Changes, including a reduction in brain size observed in humans (Oatridge et al. 2002), are also found in other regions like the hippocampus, which is highly involved in memory and learning processes (Koehl and Abrous 2011). It has also been established that pregnancy gestation is associated with various types of structural and functional neuronal changes, leading to altered excitability and plasticity, occurring changes in receptor expression mainly in hormonal systems of maternal regions (Hillerer et al. 2014). In the hippocampus, during pregnancy gestation, the dendritic tree of CA3 pyramidal neurons is less complex (Pawluski et al. 2010), and adult neurogenesis does not seem to be relevant (Hillerer et al. 2014). Although these morphological changes are known, their implications in mossy fiber synaptic transmission have not yet been determined. In previous studies performed at these synapses, using the fluorescent zinc indicator Newport Green, we have measured similar KCl-evoked zinc signals from brain slices of either male, non-gestating or pregnant gestating adult rats. This suggests that mossy fiber synaptic activity is no significantly altered during gestation and that the results of this study are independent of the pregnancy gestation condition.

The optical data was obtained from hippocampal slices containing the zinc selective fluorescent probe Newport Green (NG), which has a moderate affinity for zinc ($K_D \approx 1 \mu M$) and is virtually insensitive to calcium ($K_D \approx 100 \mu M$) (Haugland 1996). It was previously shown, in hippocampal slice experiments, that the permeant form of NG reports only postsynaptic zinc changes, because the indicator is hydrolyzed by enzymes in the cytoplasm becoming charged and unable to pass through vesicular membranes (Li et al. 2001). The origin of the signals was confirmed by the fact that they were blocked by the impermeant zinc chelator CaEDTA and by CNQX, an antagonist of the AMPA/Kainate receptor, which blocks synaptic transmission (Li et al. 2001). Thus, since presynaptic zinc is not detected with the permeant form of NG, it is considered that the signals observed with this form of the indicator have a postsynaptic origin (Li et al. 2001). It was observed that the application of SMX (180 µM) caused an increase in the basal zinc signals, as observed in the presence of KCl, that is the opposite of the effect due to TEA, that caused a depression. The antibiotic also interfered with the action of TEA, when the two drugs were applied sequentially.

## Materials and Methods

### Animals and slices

All experiments were carried out in accordance with the Directive 2010/63/EU of the European Parliament and Council. All efforts were made to minimize animal suffering and to use only the number of animals necessary to produce reliable scientific data. The experiments were performed in brain slices from Wistar rats, 10 to 13 weeks old, with 16 to 18 days of pregnancy gestation. The brains were from females animals sacrificed used by other research groups, to prepare primary cultures of cerebrocortical and hippocampal neurons cells from parts of the fetuses. These animals received no pharmaceutical treatment or other intervention before being used in the experiments. Transverse slices, 400 µm thick, were taken from the hippocampus.

### Solutions

The hippocampal slices were maintained and tested in artificial cerebrospinal fluid (ACSF) with the following constituents (in mM): NaCl 124; KCl 3.5; NaHCO$_3$ 24; NaH$_2$PO$_4$ 1.25; MgCl$_2$ 2; CaCl$_2$ 2 and D-glucose 10, pH 7.4. The slices were perfused at 1.5–2 ml/min, at a temperature in the range 30–32°C.

The hippocampal slices were incubated with the permeant form of the Newport Green (NG) indicator (5 µM), during 1 hour, at room temperature, being continuously oxygenated (5% CO$_2$ and 95% O$_2$). The NG solution was obtained dissolving 1 mg NG in 250 µl of DMSO and then diluting 5 µl of this mixture in 5 ml of ACSF containing 5 µl of pluronic acid F-127. The antibiotic SMX was applied at a concentration of 180 µM, corresponding to the EC50 that causes a 50% reduction in bacteria bioluminescence (Dantas et al. 2008). The KCl solution consisted of ACSF with higher concentration of KCl, 20 mM. The TEA medium applied in some experiments consisted of ACSF containing 25 mM of TEA and higher concentrations of CaCl$_2$ and KCl, 10 mM and 5 mM, respectively.

### Zinc measurements

The optical signals were measured in an experimental trans-fluorescence setup based on a microscope (Zeiss Axioskop), including an halogen light source (12 V, 100 W), an excitation narrow band filter (480 nm, BW 10 nm) and a high pass filter (>500 nm). The light was collected by a water immersion lens (40×, N.A. 0.75) from hippocampal CA3 area, and then focused on a photodiode (Hamamatsu, 1 mm$^2$), passing its signal through a current/voltage converter (I/V), with a 1 GΩ feedback resistance. The signals were digitally processed by
means of a 16 bit analog/digital converter, at a frequency of 1.67 Hz, using the Signal Express™ software from National Instruments. The average value of each group of 100 consecutive points was used for the purpose of illustration, after background correction.

Statistics and materials

The statistical treatment was performed using the Mann-Whitney U-test ($p < 0.05$). Drugs used were: NG (Life Technologies Inc.), SMX and TEA (Sigma Aldrich).

Results

Fluorescence data, considered to have a postsynaptic origin, were obtained from the mossy fiber synapses of hippocampal CA3 area, in slices prepared from the brain of gestating rats previously incubated with the permeant form of the zinc indicator NG. In these synapses extracellular application of KCl (20 mM), induces a zinc enhancement that is reduced during washout (Fig. 1A). Thus, it can be seen that after an initial period of 10 min in ACSF, the application of KCl for 30 minutes caused a $16 \pm 1\%$ ($n = 5$) increase of the signals with respect to baseline, in the period 35–40 min (Fig. 1A).

Unlike the results obtained with KCl, the zinc signals are reduced in the TEA (25 mM) medium, recovering, upon its removal, to values above the baseline (Fig. 1B). Thus, they are depressed in the presence of TEA and potentiated following its removal. The TEA medium caused a significant decrease in the NG-zinc signals to a steady level that is $23 \pm 4\%$ ($n = 8$) below the baseline, at 35–40 min. After washout the signals returned to positive values, measuring $7 \pm 2\%$ above the baseline, in the last 5 minutes of the experiment (Fig. 1B).

The application of the antibiotic SMX (180 µM) during a period of 30 minutes caused an increase of the zinc signals, as shown in Fig. 2A. The mean enhancement of the fluorescence signals obtained in the presence of the antibiotic, in

![Figure 1](image1.png)

**Figure 1.** Newport Green zinc signals evoked by depolarizing media. **A.** Zinc changes induced by KCl (20 mM, $n = 5$). **B.** Zinc signals evoked by tetraethylammonium (TEA) (25 mM, $n = 8$). The solutions were applied during the period indicated by the bars. The data points represent the average ± SEM of normalized dye-related fluorescence values.

![Figure 2](image2.png)

**Figure 2.** Effect of sulfamethoxazole (SMX) and tetraethylammonium (TEA) on zinc signals from slices incubated with Newport Green. **A.** Pooled data associated with the application of SMX (180 µM), during the period indicated by the bar (until 40 minutes, $n = 7$, after that $n = 4$). **B.** Zinc changes evoked by SMX followed by the perfusion of the TEA medium ($n = 3$). The solutions were applied at the time intervals indicated by the bars.
the period 35–40 min, was 6 ± 2% of control (n = 7). After the removal of SMX the zinc signals decreased to values that are 4 ± 2% below the baseline, in the last 5 min. We also tested if SMX affected the zinc changes associated with the formation of chemical TEA-LTP. For that purpose, a TEA (25 mM) containing modified ACSF solution was perfused during 30 min. Following the perfusion with SMX, characterized by increasing signals, the TEA-evoked changes had a different behavior than those registered in the experiments without prior application of the antibiotic (Fig. 2B). Thus it can be observed that the signals had a smaller reduction in the presence of TEA (15 ± 4% below baseline, n = 3, p < 0.05) at 65–70 min. In a similar way, after washout of TEA, the potentiation was significantly different from that observed in the experiments performed without previous exposure to SMX (3 ± 2% of control, n = 3, p < 0.05) in the interval 95–100 min. An illustration of the mentioned zinc changes, for the different media, is represented by the bar graphs of Figure 3. They suggest that the application of SMX interferes with the consecutive TEA-evoked zinc changes, since these are smaller following the antibiotic (Fig. 2B).

Discussion

The experiments presented in this work were carried out in brain slices from pregnant gestating animals. For this reason it might be argued that the observed effects are, in total or in part, due to the pregnancy gestation state. Support against this idea comes from studies performed in slices from male Wistar rats, at the same synapses. In these experiments, the application of KCl evoked a significant zinc potentiation (Li et al. 2001), similar to that we observed using pregnant gestating animals (Bastos et al. 2017b). These observations suggest that the results obtained in the present study, in particular those evoked by SMX, are only due to the action of the applied agents.

The first type of zinc changes observed in this work was evoked by KCl (20 mM). In the presence of this solution the resting membrane potential becomes depolarized (Bancila et al. 2004), being calculated approximately as ~50 mV, under our experimental conditions. Cell depolarization causes the release of glutamate, which induces the opening of postsynaptic membrane receptor channels, such as AMPA, kainate and NMDA and also of L- and T-type VDCCs (Fig. 4). The increase of the zinc signals in the presence of exogenous potassium (20 mM) is thus considered to be due to higher co-released zinc followed by zinc entry into the postsynaptic area, through the mentioned types of receptors and VDCCs (Sensi et al. 1997; Ketterman and Li 2008; Takeda et al. 2009).

The application of TEA (25 mM) gave rise to quite different zinc changes. The TEA-evoked depolarization is known to elicit mossy fiber LTP (Suzuki and Okada 2009). This form of chemical LTP is likely due to the simultaneous activation of a multitude of presynaptic mossy fiber boutons, containing high densities of sodium channels and fast inactivating potassium channels (Geiger and Jonas 2000; Bischofberger et al. 2006), that causes intense glutamate and zinc release. The activation of presynaptic K<sub>ATP</sub> channels by zinc leads to the efflux of potassium and the consequent hyperpolarization of the presynaptic region. This in turn leads to the closing of VDCCs and K<sub>ATP</sub> channels, and to a decrease in presynaptic calcium entry and of glutamate and zinc co-release. This sequence of events explains the observed zinc depression, in agreement with previous findings, from combined LTP and zinc experiments (Bancila et al. 2004; Quinta-Ferreira and Matias 2005; Matias et al. 2010). Upon TEA removal the field potentials become po-

![Figure 3](image-url)

**Figure 3.** Bar graphs showing the amplitude of the fluorescence signals evoked by KCl, SMX and TEA. Normalized changes during and after the application of KCl (20 mM, n = 5) and SMX (180 µM, n = 7 during, n = 4 after) (A), TEA (25 mM) without (n = 8) and with (n = 3) prior application of SMX (180 µM) (B). The bars represent the mean ± SEM.
tentiated, forming TEA LTP, due likely to the initial intense TEA evoked glutamate release (Suzuki and Okada 2009), being a related zinc potentiation observed in this work.

As for the effect of SMX the present results show that this antibiotic caused a rise in postsynaptic zinc, in agreement with the idea that it also evokes an enhancement in glutamate release. This was expected knowing that SMX inhibits carbonic anhydrase enzymes (Caspasso and Supuran 2016), leading to presynaptic alkalization and thus, higher transmitter release (Sinning and Hubner 2013). Intense co-release of glutamate and zinc and subsequent zinc permeation through postsynaptic receptors and channels (Sensi et al. 1997; Marin et al. 2000; Frederickson et al. 2005), then explain the observed rise in postsynaptic zinc. After washout the zinc signals decrease to values below the baseline, suggesting that the antibiotic alters mossy fiber synaptic physiology. Furthermore, the TEA-induced zinc depression and the subsequent potentiation, formed upon TEA removal, were both affected by prior application of SMX. The observed amplitude reductions suggest that, at the applied concentration, the TEA-evoked postsynaptic zinc changes are sensitive to this antibiotic.

Our data suggest that, at the concentration used, the effect of SMX is not exactly reversible being thus potentially harmful.

There is a multiplicity of causes of neurotoxic activity in the brain, including the formation of reactive oxygen species (Dringen 2000), excessive activation of synaptic glutamate receptors, such as N-methyl-D-aspartate (NMDA) receptors (Rothman and Olney 1995; Zou and Crews 2005), and influxes to postsynaptic neurons of divalent cations such as calcium (Choi 1998) and zinc (Sensi et al. 1997; Choi and Koh 1998; Marin et al. 2000). Since excessive zinc influx may play a major role in pathological neuronal activity (Sensi et al. 2011), the measurement of zinc signals using fluorescent indicators is a valuable tool to assess potential toxic agents.

Released zinc can bind to multiple sites existing in pre- and postsynaptic proteins including glutamate receptors, VDCCs and KATP channels, enhancing or decreasing their conductance (Smart 1989; Bancila et al. 2004) (Fig. 4). Zinc ions (Zn$^{2+}$) are represented by dark dots, glutamate (Glu) by open diamonds and other ions (Ca$^{2+}$, Mg$^{2+}$, Na$^{+}$ and K$^{+}$) by grey dots.
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