Introduction

Cellular iron homeostasis is a finely regulated process that prevents cellular damage due to iron accumulation and the formation of free radicals through the Fenton reaction [1]. The iron concentration in the brain increases with age and is much higher in the central nervous system of subjects affected by neurodegenerative diseases [2–5]. An important pathogenic role of iron has been suggested in Alzheimer’s, Parkinson’s and Huntington’s diseases, as significant iron accumulation was found in affected brain regions of patients [6]. The relevance of neuronal cellular damage by increased iron levels was further addressed by in vitro and in vivo studies of iron and 6-hydroxydopamine (6-OHDA)-dependent neurodegeneration, respectively [7]. Increased iron content, correlated with a reduced number of TH-positive neurons, was found in the substantia nigra (SN) of rats that had been overloaded with iron dextran. Significant neuroprotection was produced by deferoxamine (DFO), an iron chelator capable of permeating the blood–brain barrier, and more recent chelators in experimental models of Parkinson’s and Alzheimer’s diseases [8–12], brain ischemia-reperfusion [13,14] and hemorrhage [15].

Iron could be transported into mammalian cells as transferrin (Tf)-bound iron (TBI) via Tf receptor (TIR) mediated endocytosis or through the non-transferrin-bound iron (NTBI) pathway via divalent metal transporter-1 (DMT1). The role of TIR-mediated iron transport in neurodegeneration and ischemia is still controversial. TBI, TBI-binding sites and TIR expression are poorly correlated with the final steady-state distribution of iron [16]. Moreover, the number of TBI-binding sites decreased in dopaminergic neurons of the SN of PD patients [17,18], suggesting that the NTBI pathway is preferentially involved in the iron accumulation of PD brains. Conversely, both TIR and
DMT1 were recently shown to increase in the ischemic cortex of rats subjected to middle cerebral artery occlusion (MCAO) [13]. A significant consensus has emerged about the involvement of the NTBI pathway in neurodegenerative diseases, with iron accumulation mediated by DMT1 in specific brain areas [19]. DMT1 is highly expressed in mammalian neuronal cells [20–22,6] and is present at a relevant concentration in the basal ganglia, caudate-putamen and substantia nigra pars reticulata [23].

The mammalian DMT1 gene family (SLC11A2, Nramp2) is composed of integral membrane proteins with 10–12 putative membrane-spanning domains [24] subjected to alternative splicing. The 5’ alternative splicing generates two isoforms with or without the iron responsive element (IRE) motif in the 3’ UTR, named (+)IRE or (−)IRE isoforms, respectively. These variants give rise to four DMT1 isoforms, all of which are active in ferrous iron transport. The two (+)IRE isoforms are post-transcriptionally regulated by the IRE/Iron Regulatory Protein (IRP) system which stabilize them in the absence of iron, while (−)IRE splice variants are not susceptible to iron regulation [25,26,27]. Further complexity is added by the post-translational glycosylation of DMT1, which produces two different glycosylated molecular components: the immature, partially glycosylated, endo H-sensitive form and the mature, fully glycosylated, PNGase-F-sensitive component, with molecular masses of 60 and 90 kDa, respectively [28].

A broad up-regulation of DMT1 expression was found in the substantia nigra of PD cases as well as in animal models of PD [29,30]. However, the specific expression of (−)IRE DMT1 in neuromelanin-positive dopaminergic neurons suggested that the (−)IRE isoform is more involved in mediating abnormal increases in iron and neuronal cell loss. In line with this hypothesis, both MPP+ treatment in dopaminergic MES23.5 cells [10] and L-DOPA treatment in primary cortical neurons produced an increase of (−)IRE DMT1 expression and increased iron content with consequent cell death. These effects were counteracted by the specific silencing of the (−)IRE isoform [31]. More recently, the role of DMT1 was highlighted as being responsible for the elevated iron content in an ischemic model of MCAO [13].

Hypoxia induces transcription factors HIF-1α and NF-κB, which, in response to various stimuli, have been found to differentially activate the 5’ regulatory regions of 1A–1B/DMT1 [32–36]. NF-κB was found to specifically regulate the expression of both (−)IRE and (+)IRE 1B/DMT1 isoforms in neuronal cells exposed to sodium-nitro-prusside [37,38] and the (−)IRE DMT1 isoform in MES23.5 cells exposed to MPP+ [39]. The only indirect evidence for NF-κB-mediated DMT1 regulation in brain ischemia comes from a study showing that tanshinone II A, a natural compound reported to inhibit NF-κB activity [40], can downregulate DMT1 expression, iron elevation and brain infarct volume in mice exposed to MCAO [13]. However, a clear-cut demonstration of DMT1 regulation by NF-κB and its role in post-ischemic brain damage is still lacking.

We recently demonstrated that post-ischemic neurodegeneration relies on the activation of the NF-κB p50/RelA dimer [41,42] and requires specific RelA acetylation on the Lys310 residue. The acetylation status of RelA is maintained by the coordinated activity of epigenetic regulators such as the CREB binding protein (CBP), endowed with histone acetyl transferase activity, and sirtuin 1, a NAD–dependent histone deacetylase [43,44]. The specific RelA-Lys310 acetylation discriminates neuroprotective activation of p50/RelA during brief preconditioning ischemia from p50/RelA neurotoxic activation induced by prolonged ischemia. We speculated that 1B/(−)IRE DMT1 might be an early target gene for acetylated RelA-Lys310 in the pro-apoptotic cascade activated by ischemic injury. To test this hypothesis, we investigated the expression pattern of DMT1 isoforms during the early phase of neuronal ischemia and its relationship with the acetylation status of RelA and with cell survival. Our study provides new evidence that inhibition of 1B/(−)IRE DMT1 expression, per se or through RelA-Lys310 hypo-acetylation, might be a potential therapeutic approach to counteract post-ischemic neurodegeneration in stroke patients.

Materials and Methods

Cell culture

SK-N-SH cell culture. The human SK-N-SH neuroblastoma cell line was purchased from American Type Culture Collection. Cells were grown as previously reported in the presence of 50 μM retinoic acid (Sigma) for 10–12 days to induce mitotic arrest and differentiation into a neuronal-like phenotype [45].

Primary cultures of mouse cortical neurons. Cortical neurons were prepared from 15-day embryonic mice (C57Bl/6 dams, Charles River) and cultured for 10 days as previously described [42,46].

Cerebral ischemia models

Transient middle cerebral artery occlusion (tMCAO). All animal procedures were approved and carried out in accordance with the guidelines of Institutional Animal Care and Use Committee at University of Brescia, School of Medicine (N.07/2010), Italian Ministry of Health (n. 145/2008 – B. 30/09/2008) and the European Communities Council Directive (86/609/EEC). C57Bl/6 male mice (Harlan, Milan, Italy) were exposed to transient (20 min) MCAO as previously described [47]. Examination of infarct volume was performed in brains frozen in liquid nitrogen to avoid post-mortem changes. To prepare nuclear and cytosolic extracts, mice were killed by decapitation 4 hours after MCAO (n = 3).

Total RNA was extracted after exposure to 1 hour MCAO followed by 1 hour reoxygenation in cortices ipsilateral and contralateral to the ischemic lesion to perform expression analysis of DMT-1 mRNA isoforms by real-time polymerase chain reaction (qRT-PCR) assay (n = 3).

OGD. Primary cortical neurons at 11 DIV were exposed to Oxygen-Glucose-Deprivation (OGD) as previously described [42], for 3 hours. Cellular lysates were prepared at the end of OGD for the Luciferase reporter gene assay. SK-N-SH neuronal cells were exposed to 4 hours OGD for analysis of DMT1 in nuclear and cytosolic extracts. Total cellular RNA was extracted for the analysis of 5’-spliced mRNA isoforms. The cells were also grown on glass-coverslip and immunostained for (−)IRE DMT1. Total cellular lysates were analyzed for atomic iron content. SK-N-SH cells were also exposed to 8 hours-OGD and replaced in fresh DMEM without serum for 15 hours reoxygenation, with or without deferoxamine 50,100 μM. The same OGD protocol was performed at 40 hours expression of human (−)IRE DMT1 siRNA. Neuronal injury was then evaluated by measuring the amount of LDH released (Promega, Madison, WI, USA).

Western blot analyses

Nuclear and cytosolic protein extracts were prepared for the Immunoblot analyses from differentiated SK-N-SH as previously described [42]. The following antibodies were used: anti-pan DMT1, anti-C23 nucleolin, anti-β-actin, anti-GAPDH (Santa
Cruz Biotechnology, Santa Cruz, CA, USA), anti DMT1 (α-IRE DMT1 (Alpha Diagnostic International) and anti-human transferrin receptor (Zymed). Normalization of protein expression was performed by densitometry analysis by Gel Pro.3 analysis software.

Immunostaining

SK-N-SH cells were grown on poly-Lys coated cover glasses at the density of 2x10^5 cells/cm² in 24 wells petri dishes and differentiated as described. After 10 days, the cells were subjected to OGD for 4 hours, then fixed in paraformaldehyde in phosphate-buffered saline and incubated with Triton X-100 (0.2% in phosphate-buffered saline) containing 10% hydrogen peroxide for 15 min. Cells were blocked with Triton X-100 (0.2% in phosphate-buffered saline) containing 2%BSA, 5% goat serum, for 1 hour at room temperature and incubated for 2 hours at room temperature with polyclonal antibodies against (α-IRE DMT1 (ADI, 1:50) or pan DMT1 (Santacruz, 1:50). Biotinylated goat anti-rabbit immunoglobulins (1:500, DAKO) and an ABC kit (DAKO) were used for detection following the manufacturer’s instructions.

Semi-quantitative RT-PCR and real time PCR

Total RNA was isolated with RNAeasy kit (Qiagen) from cultured cell. The first strand of cDNA was obtained by reverse transcription from 2 μg of total RNA with 0,5 μg of oligo(dT)(25) primers (Sigma) with the SuperScript II RT (Invitrogen) according to the manufacturer’s instructions, in the presence of the ribonuclease inhibitor RNasin (Promega). 10% of the reaction was used for PCR amplification by Taq DNA Polymerase (Genespin), performed at 94 °C for 30 sec, 52 °C for 40 sec, and 72 °C for 40 sec for 20–35 cycles for β actin or DMT1 primers, respectively. The obtained cDNA was normalized against β actin by real-time PCR for every sample. The hDMT1 5’ spliced isoforms were amplified with the following primers: 1B/DMT1 For: 5’ GAGGGATCTTCATGAGGTAGTC 3’; 1A/DMT1 For: 5’ GGAGCTGCTTGAGGAAAGTC 3’; 1A/DMT1Rev: 5’ GGAGCTCTCTCATTAAGTAAG 3’; and human β actin For: 5’CTTCTCAAGTGAGCTGGCTG 3’; β actin Rev: 5’GAGGATCTTCATGAGGTAGTC 3’ in order to obtain the expected amplicon size of 222 bp, for 1B/DMT1, 290 bp, for 1A/DMT1, 320 bp for β actin. The PCR products were resolved on ethidium bromide-stained gels and quantified by the NIH ImageJ software for densitometry. DMT1 mRNA levels were determined in triplicate in three independent experiments of 4 hours OGD versus control in SK-N-SH differentiated neuronal cells. In addition, 1A/DMT1 was amplified by RT-PCR from the immortalized human urothelial cell line URO-TSA, as a positive control of 350 bp and 385 bp for 1A- and 1B-DMT1, respectively, and spliced isoforms were amplified with the following primers: 1B/DMT1 For: 5’ GGAGGGATCTTCATGAGGTAGTC 3’ and Rev: 5’ GGAGCTGCTTGAGGAAAGTC 3’ which generated a 150 bp amplicon. Incorporation of SYBR Green dye into PCR products was monitored in real-time with a BIORAD iCycler detection system, allowing determination of the threshold cycle (Ct) at which exponential amplification of PCR products begins. Each reaction was performed in triplicate. The ΔCT values for PCR products were used to calculate the amount of mRNA amplified in each reaction.

Atomic Iron Levels

Total extracts from neuronally differentiated SK-N-SH cells were analysed for Total Reflection X-ray Fluorescence, using Gallium as a reference standard at the concentration of 1 mg/ml. TXRF measurement was performed using a Bruker S2 total-reflection X-ray fluorescence spectrometer (Bruker, Germany). Data from three independent experiments were analyzed for total iron content.

Transfection with small interfering RNAs

Double-stranded small interfering RNAs (siRNA) corresponding to homologous sequences of human (α-IRE DMT1 gene were designed with 3'-hydroxyl and two base overhangs on each strand (Qiagen). The two following gene-specific sequences were successfully used: (α-IRE DMT1 siRNA1: 5'- UCCUGUGCG-CAUUUGUAUCUCA-3', and (α-IRE DMT1 siRNA2: 5’-AGG-CATTGGCGAAGAGCGTAT-3’. As a negative control (non-siRNA) was used the validated ALLstar Negative control (Qiagen). The buffered siRNAs were dissolved in RNase free water to a final concentration of 20 μM. Cell transfection was carried out using HiPerfect Transfection Reagent (Qiagen) as follows: for 10 cm² dishes, 4 μg of (α-IRE DMT1 siRNA or non-siRNA were incubated with 12 μl HiPerfect Transfection Reagent for 15 min at room temperature, according to the manufacturer’s instructions. The transfection complex was added to SK-N-SH cells in 1 ml DMEM without serum and antibiotics. It was replaced with complete DMEM 24 hours later. Time-course studies were performed to assess the selective silencing of (α-IRE DMT1 protein in nuclear extracts. The OGD experiments were performed 40 and 30 hours after transfection for DMT1 siRNA1 and siRNA2, respectively.

Chromatin Immunoprecipitation assay

Chromatin immunoprecipitation (ChIP) assays were performed using a ChIP assay kit (#9003S, Cell Signaling Technology, Massachusetts, USA). Briefly, primary corticles of mouse cortical neurons, exposed to 3 hours OGD followed by 2 hours reoxygenation, were cross-linked with 1% formaldehyde for 10 minutes at 37 °C, the reaction was then stopped with a glycine solution was added for 5 minutes at room temperature. The cells were washed with ice-cold PBS, pooled, pelleted, and incubated on ice for 10 minutes in lysis buffer supplemented with 100 mM phenylmethylsulfonyl fluoride (PMSF), dithiothreitol (DTT), and a protease cocktail inhibitor mix. Nuclei were pelleted and resuspended in buffer supplemented with DTT, digested by micrococcal nuclease, and homogenized on ice. After centrifugation, sheared chromatin was incubated with anti-actetyl Histone H3 [Lys9/18] (#07-593, Upstate-Millipore, Massachusetts, USA), anti-RelA (#sc-372, Santa Cruz Biotechnology, Santa Cruz, CA, USA), or anti-IgG, as negative control, overnight at 4 °C. Then, magnetic-coupled protein G beads were added and the chromatin was incubated for 2 hours in rotation. An aliquot of chromatin not incubated with antibodies was used as the input control sample. Antibody-bound protein/DNA complexes were washed, eluted.
treated with proteinase and subjected to Real-time polymerase chain reaction (qRT-PCR) analyses. Immunoprecipitated DNA (4 μl) was amplified in 25-μl reaction mixture containing SYBR Green master mix (12.5 μl BIORAD). The primer specific for the mouse 1B/DMT1 promoter were as follows: For: 5’ ATGGGGC-GAGCCTCCGTTCC 3’, and Rev: 5’ TCATATCCAG-GAGCCAGC 3’, which generated a 130 bp amplicon. Incorporation of SYBR Green dye into PCR products was monitored in real-time with a BIORAD iCycler detection system, allowing determination of the threshold cycle (Ct) at which exponential amplification of PCR products begins. Each reaction was performed in triplicate. The Ct values for PCR products were used to calculate the amount of chromatin immunoprecipitated in each reaction. Relative enrichment over the no-antibody INPUT was used to calculate the amount of chromatin immunoprecipitated in each reaction. Relative recruitment in OGD was then normalized to the control for each antibody.

**Report gene assay**

In order to evaluate mouse 1B/DMT1 promoter activity during OGD, cortical neurons were transfected at 10 DIV using LF 2000 Reagent with 0.2 mg/well of the DMT1-pG3L3 plasmid, kindly provided by J.A.Roth [38], and 0.8 mg/well of RelA, RelA-K310R or RelA-K310Q mutant constructs or empty expression vector pSG5 as negative control, as previously described [46]. To normalize the transfection efficiency, 0.02 μg/well Renilla luciferase control plasmid (Promega) was used. After 24 hours, neurons were exposed to 3 hour OGD as described above. Cells were then harvested and brightly and Renilla luciferase were measured by using Dual Luciferase Reporter Assay (Promega).

**Expression plasmids, transfection and treatments**

The wild-type RelA plasmid, kindly provided by P.Jalinot [48], was used as template for the substitution of Lys 310 to Arginine or Glutamine in the mutant constructs RelA-K310R, as already described [42], and RelA-K310Q, respectively. The divergent metal transporter-IRE DMT1 pDNA was obtained by PCR amplification from the (−)IRE DMT1 pDEST30 construct, kindly provided by M.Garrick [49] FOR: 5’-CACAATAGATCATCTATGGTTGG- GATCTCAAAAGAAAAGATGCCA-3’; REV: 5’-CGAATT- GATCTTGTAAGCGACACGTAGCTGACCA-3’)

and ligated into the BglII cloning site of pSG3 expression vector (Agilent Technologies, Stratagene products Division, La Jolla, CA, USA). After isolation of the sense clone by restriction analysis, the resulting plasmid was completely sequenced and the derived plasmid named 1B/−(−)IRE DMT1 -pSG5. Transfection of differentiated SK-N-SH cells was carried out according to the manufacturer’s instructions with Lipofectamine 2000 Reagent (LF2000, Invitrogen Corporation, Carlsbad, CA, USA). The day before transfection cultures were changed to normal growth medium containing serum and without antibiotics. Cells were transfected with expression plasmids encoding wild-type RelA, RelA-K310R or RelA-K310Q for 24 hours, before undergoing the OGD experiments, as previously described [42].

Treatment with Dexamethasone (DFO), dissolved in PBS, was performed at the concentration of 100 μM during OGD and overnight recovery. Cells transfected with pSG3 or 1B/−(−)IRE DMT1-pSG3 were then incubated for 1 hour in Hepes buffer at pH 6.0, after washes at pH 7.4, with or without 100 μM ferrous iron, freshly prepared in water and kept on ice [49]. LDH release was determined as neurotoxicity index.

**Statistics**

All results were calculated as the means ± se. The unpaired student’s t test was applied to analyze differences between the groups. All differences were considered statistically significant at the P value < 0.05. Statistical analyses were performed using Graph Pad Prism software (version 4.0).

**Results**

1B/(−)IRE DMT1 is up-regulated during OGD exposure in neuronally differentiated SK-N-SH cells and in the cortex of a mouse model of tMCAO

We examined DMT1 expression in neuronally differentiated human neuroblastoma SK-N-SH cells exposed to OGD, already characterised as a cell-based model of brain ischemia [42], to establish the OGD-dependent modulation of DMT1.

We first investigated whether DMT1 alternative transcripts were up-regulated at the early phase of ischemic damage in neuronal cells. We performed semi-quantitative RT-PCR of the 5’ alternative 1A and 1B/DMT1 mRNAs. We found a significant increase of 1B/DMT1 mRNA at 4 h of OGD with respect to control cells (Fig. 1A) (*, p<0.05). However, 1A/DMT1 mRNA was not detectably expressed in the same samples (Fig. 1B, left panel), in agreement with previous evidence of a highly restricted expression pattern for 1A/DMT1 in the kidney and duodenum [25]. 1A/DMT1 mRNA was amplified in the immortalised human urothelial cell line Uro-TSA, which was used as a control template (Fig. 1B, right panel).

We then investigated DMT1 protein levels during the early OGD phase in neuronal cells. The analysis was performed by western blots of cytoplasmic and nuclear extracts of SK-N-SH cells with a DMT1 antibody directed against epitopes common to the 4 isoforms (pan-DMT1). DMT1 protein was significantly up-regulated in both the nuclear compartment (**p<0.001) and in the cytosolic fraction (**p<0.001) by almost 50% in neurons exposed to 4 h of OGD, compared to control neurons. The pan-DMT1 antibody reactivity (Fig. 1C and D) revealed that both DMT1 components, with molecular masses of 60 and 90 kDa, were up-regulated in the nuclear extracts. These components are known to correspond to 1B/(−)IRE DMT1 [50,51]. Only the predominant 90 kDa component was up-regulated in the cytosolic fractions, where 1B/(−)IRE DMT1 could co-localise [50,51]. The increase of the (−)IRE DMT1 isoform, which is unresponsive to post-transcriptional regulation by intracellular iron levels through the IRE-IRP system, was confirmed by immunocytochemistry in SK-N-SH cells after 4 h of OGD. Fig. 1E clearly showed up-regulation of (−)IRE DMT1 in both nuclear and cytosolic cellular compartments after 4 hours of OGD. When staining was performed using the pan-DMT1 antibody, which did not discriminate between (+)IRE and (−)IRE isoforms, only a minor reactivity was detected (not shown), supporting the predominant up-regulation of (−)IRE isoforms during the early OGD phase.

Data indicated that 1B/(−)IRE DMT1 expression increased at both the mRNA and the protein levels during the early post-ischemic phase. These results were supported by experiments in brains of mice subjected to 20 min MCAO, a condition causing a cortical infarct of 30±28 mm3 in mice evaluated three days after ischemic injury [43]. Examination of mRNA and protein in the ischemic cortical tissue confirmed a relevant DMT1 induction. Real time PCR of cortices exposed to MCAO and 1 hour reperfusion, showed increased expression of the DMT1isoforms 1B- and (−)IRE, when compared to relative contralateral cortices (Fig. 2A). Expression of 1A/DMT1 as well as (+)IRE DMT1 were
much lower and unresponsive to the ischemic insult. Western blot analysis of brain extracts with the pan-DMT1 antibody confirmed a significant up-regulation of IRE DMT1 in both nuclear and cytosolic compartments of ischemic cortices, respect to the relative contralateral hemispheres (Fig. 2B and C).

1B/IRE DMT1 over-expression induces intracellular iron uptake and promotes iron-dependent cell death

To investigate the capability of 1B/IRE DMT1 to uptake ferrous iron in our cell model, we ectopically expressed this isoform in differentiated human neuroblastoma cells. Cells transfected with the 1B/IRE DMT1-pSG5 expression plasmid had increased levels of the DMT1 protein (Fig. 3A) \( *p<0.05 \) vs pSG5 control. When compared to control cells, the 1B/IRE DMT1 over-expressing cells exposed for 1 h to 100 \( \mu \)M ferrous iron elicited a higher iron uptake. This was shown by t-XRF analysis of the cellular iron content (Fig. 3B) \( **p<0.005 \) vs pSG5 ferrous iron treatment). Moreover, the increase of iron uptake was associated with a significant increase of cell death (Fig. 3C) \( ***p<0.0005 \) vs pSG5 ferrous iron treatment).

**Endogenous expression of 1B/IRE DMT1 during OGD promotes iron accumulation and cell death**

To investigate the relevance of iron uptake in OGD-mediated cell death, we measured iron levels by t-XRF in cells exposed to 8 hours OGD. The intracellular iron level appeared to be increased in cellular extracts of differentiated human neuroblastoma cells exposed to 4 hours (data not shown) or 8 hours OGD.
Densitometric analysis of pan-DMT1 reactivity. Values are expressed as mean ± s.e.m of two independent experiments performed in triplicate, *p<0.05 vs. control.

We then investigated the effect of (-)IRE DMT1 silencing in OGD-treated neuronal cells. DMT1 protein expression was strongly reduced both in the nuclear fraction, where only the 1B/(-)IRE isoform is present, and in the cytosolic compartment of cells treated with either the (-)IRE DMT1 siRNA1 or siRNA2 (Fig. 4A-D). (-)IRE DMT1 knockdown significantly decreased intracellular iron levels in cultures treated with both siRNAs compared to non-siRNA treated cells (Fig. 4E, p<0.05 vs non-siRNA control). Furthermore, in the siRNA-treated cultures cell death induced by 8 hours of OGD and 15 hours of reoxygenation dramatically decreased (**p<0.005 siRNA1 vs. non-siRNA OGD; #p<0.05 siRNA2 vs. non-siRNA OGD) (Fig. 4F). These findings demonstrate that both reduced intracellular iron accumulation and upstream down-regulation of (-)IRE DMT1 expression can reduce OGD-mediated neuronal cell loss.

OGD induced RelA binding and H3 histone acetylation on 1B/DMT1 promoter: effect of acetylated RelA at Lys310 on 1B/DMT1 transactivation

Since DMT1 has been proposed to be a NF-kB target gene [37,38], we investigated the relationship between its expression and NF-kB activation during OGD. To establish the effect of OGD on endogenous NF-kB/RelA interaction with 1B-DMT1 promoter and the associated histone acetylation, chromatin immunoprecipitation analysis was performed using anti-RelA and anti-acetylated histone H3 (H3K9/18ac) antibodies. Results illustrated in Fig. 5A show that RelA binding to the NF-kB activating element on 1B-DMT1 promoter increased in cortical neurons exposed to 3 hours OGD and 2 hours reoxygenation. Concomitantly, the acetylation of H3 histone associated to 1B/DMT1 promoter increased, in line with the increased expression of 1B/DMT1 during OGD.

Similar experimental conditions were used to analyse the NF-kB-dependent activation of a mouse 1B/DMT1 promoter-luciferase construct (m1B/DMT1-pGL3) [38]. Experiments were performed in mouse cortical neurons co-expressing plasmids coding for wild-type RelA or RelA carrying a mutation from lysine 310 to arginine (RelA-K310R) or the acetyl-mimic mutant with glutamine substitution of Lysine310 (RelA-K310Q). Since arginine has the same polar side chain and charge as lysine but cannot be acetylated, RelA-K310R can be used as a molecular determinant of hypo-acetylated RelA, while the glutamine residue, with uncharged, polar side chain can reproduce lysine 310 acetylation [52]. We found that OGD significantly induced 1B/DMT1 promoter activity in cells over-expressing the wild-type RelA and the RelA-K310Q, but not in cells expressing the RelA-K310R mutant plasmid (*, p<0.05 vs. relative control; #, p<0.01 vs. RelA OGD) (Fig. 5B). These results highlight that NF-kB-mediated 1B/DMT1 transcription during OGD is dependent on Lys310 acetylation of RelA.

Acetylation of RelA on Lys310 regulated DMT1 in brain ischemia and NF-kB/RelA Acetylation

We already showed that the vulnerability of neurally differentiated SK-N-SH cells to OGD depends on NF-kB/RelA activation and acetylation at Lys310 [42,43]. The cultures were transiently transfected with expression plasmids coding for wild-type RelA, the hypo-acetylated mutant RelA-K310R or the hyper-acetylated RelA-K310Q mutant. As previously reported [43], expression of RelA did not modify per se the cell survival, but be dependent on iron uptake as it was prevented by contextual application of 100 µM DFO (Fig. 3E, ***p<0.0001 vs. control; ###p<0.0001 vs. control OGD).

Figure 2. 1B/(-)IRE DMT1 is up-regulated in ischemic cortices of mice exposed to tMCAO. (A) Gene expression analysis of the differentially spliced DMT1 isoforms by qRT-PCR in ischemic brain cortices of mice exposed to 1 hour transient MCAO followed by 1 hour reoxygenation compared with contralateral hemispheres (n=3 per group). 1B/DMT1 and (-)IRE isoforms were significantly induced after MCAO. Values are expressed as fold change relative to 1A/DMT1 amplified in control (contralateral) hemisphere after normalization against β-actin cycles. Error bars represent ΔCt means ± s.e.m of two independent experiments performed in triplicate, *p<0.05, **p<0.01. (B) Representative image of pan-DMT1 immunoreactivity in nuclear and cytosolic extracts from brain cortices of mice exposed to 20 min of transient MCAO or from contralateral hemisphere (n=3 per group). Nuclear extracts were prepared 4 h after the end of the experimental condition. The pan-DMT1 reactivity increased in brain extracts of mice exposed to MCAO, both in the nuclear and in the cytosolic fractions. (C) Densitometric analysis of pan-DMT1 reactivity. Values are expressed as ratios relative to β-actin levels. Error bars represent means ± s.e.m. of three separate experiments, ***p<0.007 vs. corresponding control values.

doi:10.1371/journal.pone.0038019.g002

(Fig. 3D). The co-exposure to 100 µM DFO totally abolished the iron uptake (Fig. 3E) *p<0.05 vs.control; **p<0.05 vs. control OGD). Moreover, cell death occurring in cultures exposed to 8 hours OGD and 15 hours of re-oxygenation, also appeared to
greatly enhanced the cell vulnerability to OGD when compared to cells expressing the pSG5 empty vector. A similar cell death enhancement was produced by the hyper-acetylated RelA-K310Q mutant (Table 1). Conversely, the susceptibility of cultured cells to OGD was abolished by the expression of the RelA-K310R, as summarised in Table 1. Twenty-four hours after transfection, 1B/(-)IRE DMT1 up-regulation was examined by immunoblot analysis of nuclear extracts in cells exposed to OGD for 4 hours. The nuclear translocation of RelA and its acetylation on Lys310 were also verified by immunoblot analysis for all of the constructs tested (data not shown). In the cells over-expressing wild-type RelA, OGD promoted a significant up-regulation of both the 60 and 90 kDa components of nuclear DMT1, known to correspond to the 1B/(-)IRE isoform (Fig. 6A and B), as well as in cells over-expressing RelA-K310Q mutant (data not shown) (***p<0.0001 vs relative control). On the contrary, expression of RelA-K310R down-regulated the immature, partially glycosylated, 60 kDa DMT1 isoform to a level lower than basal value and failed to up-regulate the fully glycosylated 90 kDa DMT1 component (Fig. 6A and B). This presumably is a result of the rapid metabolic response of the endo H sensible-component [28], here associated with the regulated expression.

In the same experimental conditions, Transferrin Receptor, the other major receptor responsible for intracellular iron transport known to be induced during the late phase of post-ischemic reperfusion in the liver [53], was not involved in the early phase of OGD (data not shown) and RelA activation. Immunoblot analysis of the cytosolic extracts of neuronal cells transfected with wild-type RelA or RelA-K310R showed no change in the relative TfR component (Fig. 6C and D). Thus, the NTBI pathway through DMT1 seems to play a major role in the model of ischemia-associated neurodegeneration.
Discussion

Iron has a conserved role in evolution as a finely coordinated iron and oxygen equilibrium is essential for aerobic cellular life. Iron was selected in molecular evolution because it is critical for oxygen transport, electron transfer and, in the central nervous system, neurotransmitter biosynthesis [1]. A finely regulated iron...
homeostasis is crucial for most brain activity and conditions affecting this equilibrium may cause neurodegenerative and cognitive dysfunction [54, 52]. However, despite the recognized importance of iron delivery to the brain, the mechanisms responsible for cerebral iron uptake and their regulation in the pathogenesis of ischemia remain to be established.

This study analyses the role of DMT1, responsible for non-transferrin bound iron transport (NTBI), in cell-base and animal models of post-ischemic neurodegeneration. Our main finding was that 1B/(−)IRE DMT1 is a target gene for the Lys310-acetylated form of RelA during brain ischemia. The neuronal cell death induced by acetylation of RelA in brain ischemia was prevented by DMT1 silencing and iron chelation. The involvement of NTBI iron transport via DMT1 in neurodegenerative diseases has already been highlighted by studies that showed increased levels of iron and the DMT1 protein in SN dopaminergic neurons of both PD patients and PD animal models [29]. These studies showed that microcytic mice (mk/mk) and a Belgrade rat model carrying a G185R mutation of DMT1, which impairs iron transport, were more resistant to MPTP- and 6-hydroxydopamine-induced cell death where detected along with the up-regulation of 1B/(−)IRE DMT1 is a target gene for the Lys310-acetylated form of RelA during brain ischemia. The neuronal cell death induced by acetylation of RelA in brain ischemia was prevented by DMT1 silencing and iron chelation. The involvement of NTBI iron transport via DMT1 in neurodegenerative diseases has already been highlighted by studies that showed increased levels of iron and the DMT1 protein in SN dopaminergic neurons of both PD patients and PD animal models [29]. These studies showed that microcytic mice (mk/mk) and a Belgrade rat model carrying a G185R mutation of DMT1, which impairs iron transport, were more resistant to MPTP- and 6-hydroxydopamine-induced neurotoxicity, respectively. This evidence shows that neurotixin-induced cell death is influenced by DMT1 iron transport and for this reason modulation of iron overload through DMT1 could be a new therapeutic strategy for PD treatment.

More recently, increased DMT1 expression, with consequent increased iron content in the brain, was shown in a cerebral ischemia rat model [55]. Pharmacological neuroprotection by tanshinone IIa, previously reported to be a natural NF-κB inhibitor [40], was associated with DMT1 down-regulation [13]. A pivotal role of DMT1 in brain ischemia is evident, as DMT1 is a proton co-transporter with an optimum ferrous iron uptake at pH 5.5 [49], and the ischemia-associated extracellular acidosis may very well contribute to exacerbated iron uptake via DMT1. Analysis of 1B/ and 1A/DMT1 transcripts in SK-N-SH cells revealed a selective expression of the 1B/DMT1 isoform that was significantly up-regulated during OGD exposure. This agrees with evidence that there is little 1A/DMT1 expression in the brain and when there is, it is restricted to the kidney and duodenum [25]. The up-regulation of 1B/(−)IRE DMT1 was evident even at the protein level during OGD.

Using immunoblot analysis with the pan-DMT1 antibody, we recognised two components with molecular masses of 60 and 90 kDa corresponding to the partially and fully glycosylated mature forms of DMT1, respectively [28]. After OGD, we found a specific increase of the protein in the nuclear compartment, where only the 1B/(−)IRE DMT1 isoform has been reported to localise [49,50,51]. A different reactivity was present in the cytosolic fraction, where both (−)IRE and (+)IRE isoforms could contribute to the single 90 kDa band detected here. The predominant contribution of (−)IRE DMT1 in the early phase of OGD was confirmed by immunocytochemical staining, using an antibody selective for (−)IRE DMT1 isoform. After 4 hours of OGD, the up-regulated (−)IRE protein displayed a nuclear and cytosolic distribution. This does not exclude the involvement of the (+)IRE isoform in the later phase of OGD. Indeed, immunocytochemistry with the pan-DMT1 antibody at 4 hours of OGD did not show the same relevant staining as the (−)IRE antibody, but displayed a stronger reactivity in the whole cell after 10 hours of OGD, when (−)IRE reactivity decreased (data not shown).

We then evaluated the changes of DMT1 expression in ischemic cortices of mice subjected to transient MCAO compared to the contralateral hemispheres. We found significant up-regulation of the 1B/(−)IRE DMT1 mRNA, but not of 1A/DMT1 isoform. Also, (+)IRE DMT1 expression appeared unchanged, possibly as a consequence of its negative post-transcriptional regulation by iron increase [25–27]. Along with 1B/(−)IRE DMT1 mRNA expression, 1B/(−)IRE DMT1 protein level raised in ischemic cortices. The DMT1 nuclear and cytosolic reactivity seemed to be able to be superimposed to that observed in the cellular model. This would suggest that 1B/(−)IRE DMT1 can contribute to the increase of cerebral iron content observed in mice models of brain ischemia [13]. Indeed, increases in intracellular iron content and cell death where detected along with the up-regulation of 1B/(−)IRE DMT1 in cultures exposed to OGD as well as in cells over-expressing 1B/(−)IRE DMT1 and exposed to ferrous iron.

The application of iron chelator DFO during the OGD and reoxygenation phases, reduced both iron uptake and cell death, in accordance with previous reports showing the ability of DFO to suppress neuronal injury in experimental models of brain ischemia and subarachnoid hemorrhage [14,56–60]. The DFO-induced reduction of intracellular iron uptake during OGD confirms the relevant role of iron chelation in the neuroprotection elicited by DFO. A possible stabilization of HIF-1α protein, regulated via the
Fe²⁺- and O₂-dependent enzyme prolyl hydroxylase (PDH), may contribute to DFO neuroprotection [61].

To interfere with the upstream pathway of intracellular iron transport, we performed (²)IRE DMT1 silencing. The increase of intracellular iron levels and cell death after OGD were reduced in siRNA treated cells, compared to non-siRNA treated controls. These results further confirm that (²)IRE DMT1 up-regulation and the subsequent increase of iron influx, greatly contributing to the development of acute ischemic neurotoxicity. Consistent with our results, previous findings showed that (²)IRE DMT1 silencing protected cortical neurons from L-DOPA-mediated cell death [31]. DMT1-dependent iron uptake and toxicity was specifically addressed in a recent study of DMT1 over-expression in the human neuroblastoma SH-SY-5Y stable cell lines, which demonstrated significant enhancement of iron uptake and associated cell death [62]. Furthermore, the up-regulation of (²)IRE DMT1, but not (+)IRE, was shown to induce iron accumulation in MPP⁷-treated MES23.5 dopaminergic cells [10,39,63] as a mechanism downstream to NF-κB activation.

A recent study has demonstrated that during brain ischemia, the aberrant activation of NF-κB/p50/RelA driving the pro-apoptotic transcription of the Bim gene, relies on both p50/RelA nuclear translocation and RelA site-specific acetylation on Lys310 residue. Molecular and pharmacological deacetylation of RelA-Lys310, the latter mediated by the sirtuin-1 activator resveratrol, represses NF-κB-dependent activation of the Bim promoter as well as neuronal cell loss [43]. We therefore hypothesise that 1B/(⁻)IRE DMT1 up-regulation during OGD may depend on transactivation mediated by acetyl-RelA Lys310. The over-expression of wild-type RelA, as well as the acetyl-mimic construct RelA-K310Q, increased the 1B/DMT1 promoter activity, while expression of the acetylation-resistant RelA-K310R downregulated both. The peculiar role of endogenous NF-κB/RelA in the activation of 1B/DMT1 promoter through its proximal cis-acting element was established by chromatin immunoprecipitation analysis in cortical

Figure 6. 1B/(⁻)IRE DMT1 mediates the NTBI iron transport during OGD through Lys310-acetylated RelA. (A) Pan-DMT1 immunoreactivity in nuclear extracts from neuronal SK-N-SH cells transfected with wild-type RelA or RelA-K310R plasmids for 24 hours and further exposed to 4 hours of OGD. OGD produced a significant up-regulation of (⁻)IRE DMT1 protein in wild-type RelA overexpressing cells, with basal levels of DMT1 analogous to the empty vector (pSG5) transfected cells. Expression of RelA-K310R down-regulated the immature, partially glycosylated 60 kDa DMT1 isoform and failed to up-regulate the fully glycosylated 90 kDa DMT1 component. (B) Densitometric analysis of a representative pan-DMT1 immunoblot relative to nucleolin levels. Data from densitometric analysis of anti-panDMT1 antibody are expressed as a ratio relative to Nucleolin levels. Bars are the mean ± s.e.m. of three separate experiments (***p<0.0001 vs corresponding control value). (C) Immunoblot with anti-TfR antibody of cytosolic extracts from neuronal SK-N-SH cells transfected with wild-type RelA and RelA-K310R plasmids for 24 hours and exposed to 4 hours of OGD. The TfR protein level was not significantly modified in the early OGD phase in either RelA or RelA-K310R overexpressing cells. (D) Data from densitometric analysis of anti-TfR antibody immunoblots are expressed as a ratio relative to β-actin levels. Bars are means ± s.e.m. of three separate experiments.

doi:10.1371/journal.pone.0038019.g006
neurons exposed to OGD. The ischemic injury significantly increased the endogenous NF-κB/RelA binding to the promoter and the acetylation of the promoter-associated histone H3, supporting for a mechanism of chromatin regulation responsible for the RelA-dependent transactivation of 1B/DMT1. The same mechanisms, however, did not affect the expression of the transferrin receptor. The TIR was not responsive in early OGD phases, even after RelA over-expression, agreeing with previous reports linking TIR activation to the late phase of ischemia and reperfusion [53].

In conclusion, this study shows that site-specific acetylation of RelA at Lys310 switches on the NF-κB/IRE DMT1 during the early phase of brain ischemia. Further studies will be addressed to establish whether 1B/IRE DMT1 may become a druggable target for treatment of post-ischemic brain injury, likewise the Lys310-acetylated RelA [43]. Interfering with 1B/IRE DMT1 function may be the strategy of choice to selectively block the iron-dependent onset of neurodegeneration without perturbing iron uptake mediated by other mechanisms at the basis of the physiological aerobic homeostasis.

Acknowledgments
We wish to thank Sandra Sigala for the kind permission to use the immortalized human uterotelial cell line; Jerom A. Roth who kindly provided the mouse DMT1 promoter construct for LacZ Fraser reporter gene assay and Michael D. Gerrick for the construct (−)IRE DMT1-pDEST30 and for the extensive and helpful discussion.

Author Contributions
Conceived and designed the experiments: RI MP. Performed the experiments: RI AI IS MB F. Blasi AC LB F. Bilo LD. Analyzed the data: RI MP. Wrote the paper: RI MP. Advice and data discussion: PFS MP.

Table 1. SK-N-SH cell death after OGD depends on RelA-Lys310 acetylation.

| Relative LDH release | C | OGD1h |
|----------------------|---|------|
| pSG5                | 0.05±0.005 | 0.27±0.02 |
| RelA                | 0.04±0.003 | 0.45±0.04* |
| RelA-K310R          | 0.06±0.002 | 0.13±0.01* |
| RelA-K310Q          | 0.04±0.003 | 0.4±0.04* |

Data show the cell survival of SK-N-SH cells over-expressing a control vector, pSG5, or wild-type RelA or RelA-K310R or RelA-K310Q constructs and exposed to 15 hours of OGD, compared to controls. Wild-type RelA and RelA-K310Q over-expression significantly enhanced toxicity due to OGD, while RelA-K310R over-expression prevented cell loss. Data are mean±SEM of three separate experiments run in triplicate and represent the amount of LDH released by each well, relative to total releasable LDH. *p<0.0005 RelA or RelA-K310Q OGD vs relative pSG5; #p<0.0005 RelA-K310R OGD vs relative RelA. doi:10.1371/journal.pone.0038019.t001

References
1. Connor JR, Ghio AJ (2009) The impact of host iron homeostasis on disease. Biochim Biophys Acta 1790(7): 581–2.
2. Zecca L, Youdim MB, Riederer P, Connor JR, Crichton RR (2004) Iron, brain ageing and neurodegenerative disorders. Nat Rev Neurosci 5(1): 863–73.
3. Youdim MB, Stephen son G, Ben Shachar D (2008) Ironing iron out in Parkinson’s disease and other neurodegenerative diseases with iron chelators: a lesson from 6-hydroxydopamine and iron chelators, deferoxamine and VK-28. Ann NY Acad Sci 1012: 306–325.
4. Moos T, Rosengren Nielsen T, Skjørringe T, Morgan EH (2007) Iron trafficking inside the brain. J Neurochem 103(3): 1730–40.
5. Snyder AM, Connor JR (2009) Iron, the substantia nigra and related neurological disorders. Biochim Biophys Acta 1790(7): 606–14.
6. Moos T, Morgan EH (2004) The methabolism of neuronal iron and its pathogenic role in neuronal disease. Ann NY Acad Sci 1012: 14–26.
7. Youdim MB, Grunblatt E, Mandel S (1999) The pivotal role of Iron in NF-κB activation and nigrostriatal Dopaminergic Neurondegeneration. Ann NY Acad Sci 890: 7–25.
8. Jiang H, Luan Z, Wang J, Xie J (2006) Neuroprotective effects of iron chelator Deferoxamine on dopaminergic neurons in the substantia nigra of rats with iron overload. Neurochem Int 48(6): 605–9.
9. Mandel S, Amit T, Bar-Am O, Youdim MB (2007) Iron dysregulation in Alzheimer’s disease: multimodal brain permeable iron chelating drugs, possessing neuroprotective- neurorescue and amyloid precursor protein-processing regulatory activities as therapeutic agents. Prog Neurobiol 82: 348–360.
10. Zhang S, Wang J, Song X, Xie J, Jiang H (2009) Up-regulation of divalent metal transporter 1 is involved in 1-methyl-4-phenylpyridinium (MPP+) induced apoptosis in MES23.5 cells. Neurochem Aging 30: 1466–76.
11. Kupershmidt L, Weinreb O, Amit T, Mandel S, Bar-Am O, et al. (2011) Novel molecular targets of the multi-functional brain-permeable iron chelating drug M30 in mouse brain. Neuroscience 189: 345–54.
12. Gassen M, Youdim MB (1997) The potential role of iron chelators in the treatment of Parkinson’s disease and related neurological disorders. Pharmacol Toxicol. 80(4): 159–66.
13. Yang L, Zhang B, Yin L, Cai B, Shan H, et al. (2011) Tanshinone II A prevented brain iron dyshomeostasis in cerebral ischemic rats. Cell Physiol Biochem 27(1): 23–30.
14. Siddig A, Aminova LR, Troy CM, Suh K, Messer Z, et al. (2009) Selective inhibition of hypoxia-inducible factor (HIF) prolyl-hydroxylase 1 mediates neuroprotection against normoxic oxidative death via HIF- and CREB-independent pathways. J Neurosci 29(7): 2820–30.
15. Okauchi M, Hua Y, Keep RF, Morgenstern LB, Schallert T, et al. (2010) Deferoxamine treatment for intracerebral hemorrhage in aged rats: therapeutic time window and optimal duration. Stroke 41(2): 375–82.
32. Liu A, Paradkar PN, Singleton S, Kao HC, Garrick MD, et al. (2005) Hypoxia induces changes in expression of isoforms of the divalent metal transporter (DMT1) in rat pheochromocytoma (PC12) cells. Biochem Pharmacol 69(11): 1647–53.
33. Li Z, Lai Z, Ya K, Fang D, Ho YW, et al. (2008) Correlation between the expression of divalent metal transporter 1 and the content of hypoxia-inducible factor-1 in hypoxic HepG2 cells. J Cell Mol Med 12(2): 569–79.
34. Shah, YM, Matsubara T, Ito S, Yam SH, Gonzalez FJ (2009) Intestinal hypoxia-inducible transcription factors are essential for iron absorption following iron deficiency. Cell Metabolism 9: 152–164.
35. Wang D, Wang LH, Zhao Y, Lu YP, Zhu L (2010) Hypoxia regulates the ferrous iron uptake and reactive oxygen species level via divalent metal transporter 1 (DMT1) Exon1B by hypoxia-inducible factor-1. J Biochem 394: 173–183.
36. Qian ZM, Wu XM, Fan M, Yang L, Du F, et al. (2011) Divalent metal transporter 1 is a hypoxia-inducible gene. J Cell Physiol 226(6): 1596–603.
37. Paradkar PN, Roth JA (2006) Post-translational and transcriptional regulation of DMT1 during P19 embryonic carcinoma cell differentiation by Retinoic acid. Biochemical J 394: 173–183.
38. Shah, YM, Matsubara T, Ito S, Yam SH, Gonzalez FJ (2009) Intestinal hypoxia-inducible transcription factors are essential for iron absorption following iron deficiency. Cell Metabolism 9: 152–164.
39. Faraco G, Blasi F, Min W, Wang ZQ, Moroni F, et al. (2010) Brain ischemic injury. J Neurosci 26;50): 12896–903.
40. Inta I, Paxian S, Maegele I, Zhang W, Pizzi M, et al. (2006) Bim and Noxa are candidates to mediate the deleterious effect of the NF-kappa B subunit RelA in neuronally differentiated SK-N-SH human cell line. Eur J Neurosci 16(12): 2342–50.
41. Pizzi M, Boroni F, Bianchetti A, Moraitis C, Sarnico I, et al. (2002) Expression of functional NR1/NR2B-type NMDA receptors in neuronally differentiated SK-N-SH human cell line. Eur J Neurosci 16(2): 2342–50.
42. Chen LF, Mu Y, Greene WC (2002) Acetylation of p65/RelA at discrete sites regulates distinct nuclear functions of NF-kB. EMBO J 21: 6359–6360.
43. Tacchini L, Fusar Poldi D, Bernelli-Zazzera A, Cairo G (2002) Transferrin receptor gene expression and transferrin-bound iron uptake are increased during post-ischemic rat liver reperfusion. Hepatology 36(1): 103–11.
44. Parada, PN, Roth JA (2006) Nitric oxide transcriptionally down-regulates specific isoforms of divalent metal transporter (DMT1) via NF-kappaB. J Neurochem 96(6): 1768–77.
45. Jang SI, Kim HJ, Kim YJ, Jeong SI, You YO (2006) Tanshinone IIA inhibits ferrous iron uptake and reactive oxygen species level via divalent metal transporter 1 (DMT1) Exon1B by hypoxia-inducible factor-1. J Biochem 394: 173–183.
46. Pizzi M, Boroni F, Bianchetti A, Moraitis C, Sarnico I, et al. (2002) Expression of functional NR1/NR2B-type NMDA receptors in neuronally differentiated SK-N-SH human cell line. Eur J Neurosci 16(12): 2342–50.
47. Lis A, Paradkar PN, Singleton S, Kuo HC, Garrick MD, et al. (2005) Hypoxia regulates distinct nuclear functions of NF-κB. J Neurosci 26(50): 12896–903.
48. Chen LF, Mu Y, Greene WC (2002) Acetylation of p65/RelA at discrete sites regulates distinct nuclear functions of NF-kB. EMBO J 21: 6359–6360.
49. Sarnico I, Lanzillotta A, Boroni F, Benarese M, Alghisi M, et al. (2009) NF-kappaB p50/RelA and c-Rel-containing dimers: opposite regulators of neuron vulnerability to ischemia. J Neurochem 101(2): 475–85.
50. Pizzi M, Spera P (2006) Distinct roles of diverse nuclear factor-kappaB complexes in neuropathological mechanisms. Eur J Pharmacol 545(1): 22–4.
51. Jang SI, Kim HJ, Kim YJ, Jeong SI, You YO (2006) Tanshinone IIA inhibits LPS-induced NF-kappaB activation in RAW 264.7 cells: possible involvement of the NIK-IKK, ERK1/2, p38 and JNK pathways. Eur J Pharmacol 542(1-3): 1–7.
52. Pizzi M, Spera P (2006) Distinct roles of diverse nuclear factor-kappaB complexes in neuropathological mechanisms. Eur J Pharmacol 545(1): 22–4.
53. Tacchini L, Fusar Poldi D, Bernelli-Zazzera A, Cairo G (2002) Transferrin receptor gene expression and transferrin-bound iron uptake are increased during post-ischemic rat liver reperfusion. Hepatology 36(1): 103–11.
54. Parada PN, Roth JA (2006) Nitric oxide transcriptionally down-regulates specific isoforms of divalent metal transporter (DMT1) via NF-kappaB. J Neurochem 96(6): 1768–77.
55. Jang SI, Kim HJ, Kim YJ, Jeong SI, You YO (2006) Tanshinone IIA inhibits ferrous iron uptake and reactive oxygen species level via divalent metal transporter 1 (DMT1) Exon1B by hypoxia-inducible factor-1. J Biochem 394: 173–183.
56. Lis A, Paradkar PN, Singleton S, Kuo HC, Garrick MD, et al. (2005) Hypoxia regulates distinct nuclear functions of NF-κB. J Neurosci 26(50): 12896–903.
57. Parada PN, Roth JA (2006) Nitric oxide transcriptionally down-regulates specific isoforms of divalent metal transporter (DMT1) via NF-kappaB. J Neurochem 96(6): 1768–77.
58. Xu H, Jiang H, Wang J, Xie J (2010) Rgl1 protects the MPP+/treated MES23.5 cells via attenuating DMT1 up-regulation and cellular iron uptake. Neuropharmacology 58(2): 488–94.
59. Sarnico I, Lanzillotta A, Boroni F, Benarese M, Alghisi M, et al. (2009) NF-kappaB p50/RelA and c-Rel-containing dimers: opposite regulators of neuron vulnerability to ischemia. J Neurochem 101(2): 475–85.
60. Lanzillotta A, Sarnico I, Inggrassia R, Boroni F, Branca C, et al. (2010) The acetylation of RelA in Lys310 dictates the NF-kB-dependent response in post-ischemic injury. Cell Death Dis 1(11): e96.
61. Yang XD, Takedashir E, Chen LF (2010) Functional interplay between acetylation and methylation of the RelA subunit of NF-kB. Mol Cell Biol 30(9): 2170–80.
62. Pizzi M, Boroni F, Bianchetti A, Moraitis C, Sarnico I, et al. (2002) Expression of functional NR1/NR2B-type NMDA receptors in neuronegally differentiated SK-N-SH human cell line. Eur J Neurosci 16(2): 2342–50.
63. Timiriazeva B, Tsakanov N, Tsakanov A, Tsakanov S, et al. (2009) Brain ischemic injury. J Cereb Blood Flow Metab 30(5): 1245–50.
64. Ben-Shachar D, Eshel G, Finberg JP, Youdim MB (1991) The iron chelator deferoxamine (Desferal) retards 6-hydroxydopamine-induced degeneration of nigrostriatal dopamine neurons. J Neurochem 56: 1411–1414.
65. Chawla V, Sharma R, Jain R, et al. (2011) Enhanced autophagy from chronic toxicity of iron and mutant A53T α-synuclein: implications from neuronal cell death in Parkinson disease. J Biol Chem 286(38): 33380–9.
66. Xie J, Jiang H, Wang J, Xie J (2010) Rgl1 protects iron-induced neurotoxicity through antioxidant and iron regulatory proteins in 6-OHDA-treated MES23.5 cells. J Cell Biochem 111(8): 1537–45.