Ammonium induced impairment of axonal growth is prevented through glial creatine.

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Abstract

Hyperammonemia in neonates and infants affects brain development and causes mental retardation. We report that ammonium impaired cholinergic axonal growth and altered localization and phosphorylation of intermediate neurofilament protein (NF-M) in rat reaggregated brain cell primary cultures. This effect was restricted to the phase of early maturation but did not occur after synaptogenesis. Exposure to NH₄Cl decreased intracellular creatine, phosphocreatine and ADP. We demonstrate that creatine co-treatment protected axons from ammonium toxic effects, although this did not restore high energy phosphates. The protection by creatine was glial cell dependent. Our findings suggest that means to efficiently sustain central nervous system (CNS) creatine concentration in hyperammonemic neonates and infants should be assessed to prevent impairment of axonogenesis and irreversible brain damage.
Introduction

Liver failure and genetic defects affecting the urea cycle lead to hyperammonemia with reversible and irreversible neurological damage that might be life-threatening (Podolsky and Isselbacher, 1998; Brusilow and Horwich, 2001). Symptoms of irreversible damage include cognitive impairment, seizures and cerebral palsy (Flint Beal and Martin, 1998). They mainly occur in case of prolonged hyperammonemic crises and/or when blood ammonium reaches levels between 180 and 500 µM, during the two first years of life (Msall et al., 1984; Uchino et al., 1998; Bachmann et al., 2002a, Bachmann et al., 2002b). In the few reported brain autopsies of pediatric patients, Alzheimer type II astrocytes, perivascular spongiosis and cystic necrosis at the junction of cortical grey and white matter have been observed (Harding et al., 1984; Filloux et al., 1986; Dolman et al., 1988). The mechanisms leading to irreversible alterations are not understood.

We have shown previously that mimicking hyperammonemia by adding ammonium in a rat reaggregating brain cell culture model causes a reduction in aggregate size, which could be due to a deficiency or developmental delay of cellular processes in regions where axons are prevalent (Braissant et al., 1999b). There is substantial evidence that creatine (Cr) and the Cr / phosphocreatine (PCr) / creatine kinase (CK) system are involved in neuronal growth cone activity and axonal elongation (Wang et al., 1998). In addition, the central nervous system (CNS) is the main affected target in infants with a creatine-deficiency syndrome due to guanidinoacetate methyltransferase, arginine:glycine amidinotransferase or Cr transporter deficiencies. Such patients exhibit delayed psychomotor development or bilateral myelination delay (Stöckler et al., 1994; Schulze et al., 1997; Salomons et al., 2001; Item et al., 2001). We and others have also shown that the synthesis of arginine, the main precursor of creatine, is
altered by hyperammonemia (Rao et al., 1995; Braissant et al., 1999b). Taken together, these findings suggest that hyperammonemia impairs Cr metabolism and/or transport in the CNS. Cr is used by the CNS Cr/PCr/CK system to buffer and supply ATP, particularly to neurons (Hemmer and Wallimann, 1993). CNS, at least in the rat, seems dependent on its own Cr synthesis (Braissant et al., 2001).

The aim of the present work was to test whether axons were affected by hyperammonemia, and whether this could be due to a Cr loss or decrease in brain cells. We used reaggregated brain cell cultures prepared from the telencephalon of rat embryos, which provide a three-dimensional network of neurons and glial cells that progressively acquire a tissue specific pattern resembling that of the brain (Honegger et al., 1979; Honegger and Monnet-Tschudi, 2001). We analyzed the axonal expression of the intermediate neurofilament protein (NF-M, 160kD) and determined the intracellular levels of Cr, PCr, ATP, ADP and AMP under NH$_4$Cl exposure. We identified the neurons impaired in axonal growth by ammonium exposure. Finally, we tested the protective effect of Cr on the growth of axons exposed to NH$_4$Cl, and the influence of the brain cell developmental stage on the axonal vulnerability to ammonium.
Materials and Methods

Reaggregated brain cell cultures

Reaggregated brain cell primary cultures were prepared from mechanically dissociated telencephalon of 16-day rat embryos as previously described (Honegger and Monnet-Tschudi, 2001). Cultures were grown in serum-free, chemically defined medium consisting of Dulbecco’s modified Eagle’s medium with high glucose (25 mM) supplemented with insulin (0.8 µM), triiodothyronine (30 nM), hydrocortisone-21-phosphate (20 nM), transferrin (1 µg/ml), biotin (4 µM), vitamin B12 (1 µM), linoleate (10 µM), lipoic acid (1 µM), L-carnitine (10 µM), and trace elements (Honegger and Monnet-Tschudi, 2001). Gentamicin sulfate (25 µg/ml) was used as antibiotic. The cultures were maintained under constant gyratory agitation (80 rpm), at 37°C and in an atmosphere of 10% CO2 / 90% humidified air.

Two different kinds of aggregate cultures were grown:

A) Regular mixed-cell aggregates, developing with neurons and glial cells.

B) Neuron-enriched aggregates, which were obtained by the treatment of the regular mixed-cell cultures at days 1 and 2 with cytosine arabinoside (0.4 µM) to eliminate the proliferating glioblasts (Honegger and Monnet-Tschudi, 2001). Previous work has shown that the neuron-enriched cultures contain > 90% neurons, less than 8% astrocytes and very few, if any, oligodendrocytes (Honegger and Pardo, 1999). Neuron-enriched cultures received conditioned medium taken from mixed-cell (neuron-glia) sister cultures and diluted 1:1 with fresh medium.

Aggregates were grown for 13 or 28 days. Aggregates harvested at 13 days were treated from day 5 onwards with NH4Cl and/or Cr, with medium replenishments at days 8 and 11 (5 ml
replaced out of 8 ml total). Aggregates harvested at 28 days were treated from day 20 onwards with NH₄Cl and/or Cr, with medium replenishments at days 22, 24 and 26 (5 ml replaced out of 8 ml total). On the day of harvest, culture media were recovered, immediately centrifuged to remove cell debris, and frozen in liquid nitrogen. Aggregate pellets were washed three times with ice cold PBS, and embedded for histology or frozen in liquid nitrogen. Until analysis, culture media and aggregate pellets were kept at -80°C.

**Histology and immunohistochemistry**

Aggregates were embedded in tissue freezing medium (Jung, Nussloch, Germany), frozen in liquid nitrogen cooled isopentane and kept at -80°C until used. For histological staining, 16 µm thick cryosections were prepared, postfixed 1 hour in 4% paraformaldehyde (PFA) in PBS and stained by hematoxylin coloration or Bielschowsky’s silver impregnation (Cox, 1977). NN-18 (Chemicon International, Temecula CA; Shaw et al., 1986; Harris et al., 1991) and RMO-44 (Zymed Laboratories, San Francisco CA; Lee et al., 1987) monoclonal antibodies were used for immunohistochemistry against NF-M. Axonal growth cones were detected with a monoclonal antibody directed against growth cone associated protein 43 (GAP43 Chemicon International). Astrocytes and oligodendrocytes were labeled using monoclonal antibodies directed against glial fibrillary acidic protein (GFAP, Chemicon International) and myelin basic protein (MBP, Boehringer Ingelheim). Cholinergic, GABA-ergic and glutamatergic neurons were characterized by using monoclonal antibodies against choline acetyltransferase (ChAT, Oncogene, Cambridge MA), glutamic acid decarboxylase (GAD 65/67 kD, Santa Cruz Biotechnology, Santa Cruz CA) and neuronal excitatory amino acid transporter 3 (EAAT3, Santa Cruz Biotechnology) respectively (Furuta et al., 1997; Pardo and Honegger, 1999; He et al., 2000). For immunohistochemistry against NF-M, GAP43, GFAP, GAD and EAAT3, cryosections (16 µM) were fixed for 1 hour in 4% PFA-
PBS at room temperature, washed in PBS (3 x 5 min) and permeabilized for 5 min in 0.1% sodium citrate and 0.1% Triton X-100. For immunohistochemistry against ChAT, cryosections (16 µM) were fixed for 1 min in 5% acetic acid / 95% ethanol at 4°C and washed in PBS (3 x 5 min). For immunohistochemistry against MBP, cryosections (16 µM) were fixed for 1 hour in 4% PFA-PBS at room temperature, washed in PBS (3x 5 min), dehydrated with increasing concentrations of EtOH, delipidated in xylene, rehydrated with decreasing concentrations of EtOH, and finally washed in PBS. Sections were then processed for immunohistochemistry using the Histostain-Plus kit according to the manufacturer’s protocol (Zymed Laboratories). The primary antibody was diluted in 1% bovine serum albumin in PBS and applied to sections. After washing, sections were incubated with a biotinylated anti-mouse IgG secondary antibody followed by a streptavidin-peroxidase conjugate. Peroxidase staining was performed for 10 min using aminoethyl carbazole and H₂O₂ and stopped in distilled water. Sections were mounted under glycerol, observed and photographed on an Olympus BX50 microscope equipped with a DP-10 digital camera (Olympus Opticals).

**Protein dephosphorylation on cryosections**

Cryosections (16 µm thick) were digested 30 min (37°C) with alkaline phosphatase (200 µg/ml, Roche Biochemicals), in Tris 100mM, NaCl 100mM, MgCl₂ 50 mM, pH 9.5. Sections were washed in PBS, fixed 30 min in 4% PFA-PBS, and washed 3 x 5 min in PBS. Immunohistochemistry against NF-M was then performed as described above.

**Quantitative western blot analysis**

Mixed-cell and neuron-enriched aggregates were homogenized in 10 mM Tris/HCl pH 7.5 containing 4 M urea, 0.1 % SDS and protease inhibitors (Complete, Roche, Switzerland). Homogenates were centrifuged at 16'000 g for 10 min and supernatants were recovered.
Supernatant proteins were measured by the bicinchoninic acid assay (Pierce, USA) and diluted at a final concentration of 1 µg/µl in Laemmli sample buffer (Laemmli, 1970). Proteins were separated by SDS-PAGE (9% total acrylamide). After transfer of the proteins to polyvinylidene difluoride membranes (Immobilon, Millipore, USA), blots were probed with NN-18 and RMO-44 anti-NF-M, anti-GAP43 and anti-GFAP monoclonal antibodies. Western blots were revealed by chemiluminescence (ECL, Amersham Biosciences, England). The radiographs (Kodak X-OMAT AR, USA) were scanned with a ImageScanner (Amersham Biosciences, England) and processed by image analysis (ImageMaster 1D, Amersham Biosciences, England).

**Measure of intracellular Cr, PCr, ATP, ADP and AMP**

Frozen aggregate pellets were suspended in ice-cold 25 mM phosphate buffer pH 7.5 containing 1 mM iodoacetate and 0.1 mM diadenosine pentaphosphate. Cells were extracted with 0.4 M perchloric acid, and neutralized with 2 M K₂CO₃ followed by a centrifugation at 16'000 g for 5 min. Supernatants were assayed for adenine nucleotides (ATP, ADP and AMP) and creatine compounds (Cr and PCr) according to the isocratic reverse-phase HPLC method of Seidl et al. (2000) using an HPLC (Agilent, USA) coupled with a dual absorbance detector. The absorbance at 214 nm (for Cr and PCr) and 254 nm (for ATP, ADP and AMP) were monitored simultaneously. The acid precipitates of the aggregates were solubilized in 0.1 M NaOH, 1 % SDS and their protein content was determined by the bicinchoninic acid assay (Pierce, USA). The concentrations of Cr, PCr, ATP, ADP and AMP were expressed as nmol · (mg prot)⁻¹.
Measure of NH₄⁺, glucose and lactate in culture media

Ammonium was measured on a Cobas FARA II automate (Roche), using the UV Enzymatic Ammonium Kit (Biomérieux, N° 61025). Glucose and lactate were measured on a Hitachi 917 automate (Roche), using the Glucose Ecoline 100 kit (Merck, N° 1.14891.0001) and the Lactate kit (Roche, N° 1 822 837).
Results

Net uptake of NH$_4^+$ and glucose, and net release of lactate, by mixed-cell aggregate cultures exposed to NH$_4$Cl are shown in Table 1. During the whole period of NH$_4$Cl exposure, NH$_4^+$ uptake by cultures was dose dependent. Glucose uptake and lactate release showed highly significant increases at the maximal dose of 5 mM NH$_4$Cl. The extent of metabolic changes observed with increasing NH$_4$Cl concentrations paralleled morphological changes in the aggregates, which were absent or barely detectable at 1 mM and 2.5 mM NH$_4$Cl (data not shown) but evident at 5 mM (see below). Therefore, morphological and biochemical effects of hyperammonemia were further analyzed by treating aggregate cultures with 5 mM NH$_4$Cl.

NH$_4$Cl exposure impaired axonal growth in developing mixed-cell aggregates

Control mixed-cell aggregate cultures at day 13 presented a characteristic distribution of cells, including a peripheral zone with a low density of cell bodies (Fig. 1A,B; asterisk) in which fibers were prevalent (Fig. 1C). These fibers were NF-M positive, using the monoclonal anti-NF-M NN-18 antibody which did not stain neuronal soma (Fig. 1D). Aggregates exposed to 5 mM NH$_4$Cl from days 5 to 13 showed more densely packed cell bodies at their periphery (Fig. 1K,L), and the almost complete absence of NF-M positive fibers (Fig. 1M,N). Another monoclonal anti-NF-M antibody, RMO-44, localized NF-M in neuronal cell bodies but not in fibers (Fig. 1E). The somatic expression of NF-M was not altered by 5 mM NH$_4$Cl exposure (Fig. 1O). Western blot analysis of NF-M showed that NH$_4$Cl exposure caused a drastic decrease of NF-M (3-fold using NN-18, Fig. 1U; and 2-fold using RMO-44, Fig. 1V).

The specificity of NN-18 and RMO-44 antibodies for NF-M recognition was further characterized by dephosphorylation of proteins in situ using alkaline phosphatase.
Dephosphorylation abolished most of the NN-18 staining in fibers of the aggregate periphery (Fig. 2A,B) and induced RMO-44 NF-M immunoreactivity in distal fibers located at the aggregate periphery and in proximal fibers connecting neuronal cell bodies to the aggregate periphery (Fig. 2C,D). This indicates that NN-18 predominantly recognized phosphorylated NF-M, whereas RMO-44 predominantly recognized non-phosphorylated NF-M. Thus, the absence of NN-18 anti-NF-M staining after NH₄Cl exposure (Fig. 1N) suggests that ammonium exposure inhibited NF-M phosphorylation. Fibers located at the aggregate periphery and containing phosphorylated NF-M were most probably axons, as suggested by analyzing the expression of GAP43, an axonal growth cone marker. In control day 13 mixed-cell aggregates, GAP43 was highly expressed in the same peripheral zone exhibiting strong NF-M staining by NN-18 (Fig. 3A). Like for NF-M, the GAP43 staining was lost in this region after NH₄Cl exposure (Fig. 3G). Interestingly, a strongly GAP43 immunoreactive zone appeared at the border of the aggregates (Fig. 3G, bracket). Western blotting analysis of GAP43 after NH₄Cl exposure showed a small decrease of the GAP43 signal (-20%, Fig. 3M).

In day 13 mixed-cell aggregates, glial cells were identified by immunohistochemical staining for GFAP (specific for astrocytes, Fig. 3B) and MBP (specific for oligodendrocytes, Fig. 3C). NH₄Cl exposure increased the number of GFAP positive astrocytic processes, particularly at the aggregate border (Fig. 3H, bracket), whereas no significant effect was found for oligodendrocytes (Fig. 3I). Western blotting analysis of GFAP after NH₄Cl exposure showed a 2.2-fold increase of the GFAP signal (Fig. 3N).

**NH₄Cl exposure decreased intracellular Cr, PCr and ADP in developing mixed-cell aggregates**
To test whether energy rich phosphates and/or the Cr/PCr/CK system were involved in ammonium induced axonal growth impairment, we measured intracellular levels of Cr, PCr, ATP, ADP and AMP in mixed-cell aggregates (Table 2). NH$_4$Cl exposure reduced Cr, PCr and ADP significantly (83, 75 and 69 % of controls, respectively), whereas no significant effect was observed for ATP and AMP content.

**Creatine prevented NH$_4$Cl induced axonal growth impairment in developing mixed-cell aggregates**

Because Cr and PCr were decreased in NH$_4$Cl exposed aggregates, we examined whether Cr had a protective effect on axonal growth under NH$_4$Cl exposure. By immunohistochemistry with the NN-18 antibody, we found indeed that 1 mM Cr added to NH$_4$Cl exposed aggregates protected the peripheral axons, their expression of NF-M as well as NF-M phosphorylation (Fig. 1P-S; see controls: A-D; and NH$_4$Cl exposure: K-N), whereas Cr given alone increased the density of NF-M positive peripheral axons (Fig. 1F-I). Immunohistochemistry with RMO-44 did not reveal any difference in NF-M expression between aggregates treated with Cr and controls (Fig. 1J,E) nor between NH$_4$Cl+Cr and NH$_4$Cl exposed aggregates (Fig. 1T,O). Analysis of NF-M by western blotting showed that Cr co-treatment maintained NF-M at control levels in mixed-cell aggregates exposed to NH$_4$Cl (NN-18, Fig. 1U) or showed partial protection (RMO-44, -25% as compared to controls, +50% as compared to NH$_4$Cl exposure, Fig. 1V), whereas Cr exposure alone did not affect the NF-M level as compared to controls (Fig. 1U,V). A higher concentration of Cr (25 mM) was also tested, which did not improve the protection of axonal growth under ammonium exposure, as compared to that obtained with 1 mM Cr (data not shown).

GAP43 expression was partially protected in the peripheral NF-M positive region of NH$_4$Cl exposed aggregates co-treated with Cr (Fig. 3J, asterisk; see control: A; and NH$_4$Cl exposure:
G). However, the border of the aggregates was strongly positive for GAP43, as after exposure to NH₄Cl alone (Fig. 3J,G, bracket). As compared to controls, Cr given alone increased the GAP43 signal in the center of the aggregates (Fig. 3D,A). By western blot analysis, no difference was observed for GAP43 expression between Cr exposed cultures and controls, nor between NH₄Cl+Cr and NH₄Cl exposed cultures (Fig. 3M). Astrocytes and oligodendrocytes presented similar patterns of expression for GFAP and MBP in cultures treated with Cr and in controls (GFAP Fig. 3E,B; MBP Fig. 3F,C) as well as in NH₄Cl+Cr and NH₄Cl treated aggregates (GFAP Fig. 3K,H; MBP Fig. 3L,I).

No restoration of nor increase in intracellular PCr, ATP or ADP could be observed in mixed-cell aggregates co-treated with 5 mM NH₄Cl and 1 mM Cr as compared to cultures exposed to NH₄Cl only (Table 2), whereas intracellular Cr was increased significantly (170% of controls). Cr was efficiently taken up by mixed-cell aggregates exposed to 1 mM Cr only (190% of controls, p < 0.001); however their PCr, ATP or ADP content was not modified. NH₄Cl induced a significant decrease of intracellular Cr in aggregates exposed to 1mM Cr (75% of Cr alone exposure, p < 0.005).

**Creatine did not prevent the NH₄Cl induced axonal growth impairment in developing neuron-enriched aggregates**

Untreated neuron-enriched aggregate cultures also developed a peripheral zone with a low density of cell bodies, but devoid of the glial cell lining found at the border of mixed-cell aggregates (Fig. 4A vs. 1B; compare also staining for GFAP, Fig. 4D vs. 3B; and MBP, Fig. 4E vs. 3C). Like in mixed-cell cultures, the number of axons that developed in the aggregate periphery of day 13 neuron-enriched aggregates (Fig. 4B) was decreased under NH₄Cl exposure, as was NF-M phosphorylation (Fig. 4K,L). In contrast to mixed-cell cultures
however, Cr co-treatment of neuron-enriched cultures did not prevent the NH$_4$Cl induced
axonal growth impairment nor the decrease of NF-M phosphorylation (Fig. 4P,Q; see controls
Fig. 4A,B; and NH$_4$Cl exposure, Fig. 4K,L). In contrast to mixed-cell cultures, Cr given alone
did not change the density of NF-M positive axons (Fig. 4F,G; see controls Fig. 4A,B).
Similar densities of NF-M positive neuronal cell bodies stained with RMO-44 were found in
controls, NH$_4$Cl, and NH$_4$Cl+Cr exposed neuron-enriched aggregates (Fig. 4C,M,R).
However, exposure to Cr alone caused an increased density of neuronal cell bodies, as well as
a less intense staining of the non-phosphorylated NF-M around the cell nucleus (Fig. 4H).
Analysis by western blotting and staining with either NN-18 or RMO-44 showed that NF-M
was quantitatively decreased in NH$_4$Cl exposed neuron-enriched aggregates, and that Cr co-
treatment of NH$_4$Cl exposed cultures did not protect NF-M expression (Fig. 4U,V).

The presence of few remaining astrocytes in neuron-enriched cultures was confirmed by anti-
GFAP labeling (Fig. 4D; see mixed-cell cultures Fig. 3B). NH$_4$Cl exposure (with or without
Cr co-treatment) did not change the density of GFAP positive astrocytic processes, but caused
an increase in the size of astrocyte cell bodies, reminiscent of Alzheimer type II astrocytes
(Fig. 4N,S, arrows; see controls Fig. 4D). Exposure to Cr alone did not modify the GFAP
level, nor the astrocyte soma diameter, nor the astrocytic process density (Fig. 4I; see controls
Fig. 4D). The complete absence of oligodendrocytes in neuron-enriched aggregates was
confirmed by anti-MBP staining in both controls and treated cultures (Fig. 4E,J,O,T).

Creatine and phosphocreatine levels in neuron-enriched cultures were not altered by
NH$_4$Cl exposure
In control conditions, the Cr concentration found in neuron-enriched aggregates was 10-fold lower than in mixed-cell cultures (compare Tables 3 to 2), suggesting that neurons had a limited capacity to synthesize Cr. In contrast to mixed-cell aggregates, NH$_4$Cl exposure of neuron-enriched cultures did not affect intracellular Cr and PCr (Table 3), but ATP, ADP and AMP were significantly lower (67%, 58% and 50% of controls, respectively). Cr co-treatment did not attenuate the NH$_4$Cl induced decrease of adenine nucleotides, nor did it modify the PCr concentration (Table 3). Neuron-enriched cultures were able to take up Cr efficiently, as exposure to 1mM Cr increased intracellular Cr 8.4-fold when compared to control levels (Table 3, p < 0.001; compare to 1.9-fold only in mixed-cell cultures, Table 2), and 3.5-fold when comparing NH$_4$Cl+Cr with NH$_4$Cl exposed cultures (p < 0.01). NH$_4$Cl caused a significant decrease of Cr in neuron-enriched aggregates exposed to 1mM Cr (40% of Cr alone exposure, p < 0.005).

**Neurons affected by NH$_4$Cl exposure in day 13 mixed-cell aggregates are cholinergic**

The neuronal identity of axons affected by NH$_4$Cl was analyzed by immunohistochemistry against ChAT, GAD 65/67 kD and EAAT3 to discriminate between cholinergic, GABA-ergic and glutamatergic neurons respectively. ChAT was expressed in neuronal cell bodies of day 13 aggregates, as well as in punctuate formations localized in the peripheral zone where axons grow (Fig. 5A). Following NH$_4$Cl exposure, ChAT immunostaining decreased in the peripheral axonal region (Figure 5B), but was maintained in aggregates exposed to NH$_4$Cl+Cr (Fig. 5D). Cr alone did not modify the punctuate staining of ChAT in the aggregate periphery, but increased it in neuronal cell bodies (Fig. 5C). GAD 65/67 kD was localized in numerous neuronal cell bodies, and its expression was not modified by NH$_4$Cl (Fig. 5E,F). At day 13, no expression of the glutamate transporter EAAT3 could be detected in the aggregates (data not shown).
NH₄Cl and creatine exposure did not alter axons of mature mixed-cell aggregates

To investigate whether NH₄Cl affected axons also in mature neurons, aggregates were exposed to 5 mM NH₄Cl and/or 1 mM Cr from days 20 to 28, i.e. at a stage where neurons progressively undergo synaptogenesis and myelination (Honegger and Monnet-Tschudi, 2001). At day 28, the peripheral dense fiber zone was clearly visible (Fig. 6A, B) and richer in number of axons expressing NF-M than at day 13 (Fig. 6C, compare to Fig. 1D). NH₄Cl (5 mM) exposure did not affect axons of mature aggregates (Fig. 6D). Aggregates treated with 1 mM Cr (Fig. 6E) or with NH₄Cl+Cr (Fig. 6F) were also comparable to control aggregates (Fig. 6C). Quantitative analysis of NF-M by western blotting, using NN-18, did not reveal any significant difference between controls and treated cultures (Fig. 6G).
Discussion

Ammonium exposure impairs cholinergic axonal growth, decreases NF-M and inhibits NF-M phosphorylation

We provide the first experimental demonstration of neuronal fiber growth impairment under ammonium exposure, together with a decrease in NF-M protein and an inhibition of NF-M phosphorylation. This is in line with the growing number of recently described neurological pathologies showing abnormal expression or phosphorylation of neuronal cytoskeleton proteins such as neurofilaments (NFs) or microtubule associated proteins (Hirokawa and Takeda, 1998; Julien, 1999; Saez et al., 1999; Sanchez et al., 2000). Our results are consistent with clinical findings in hyperammonemic neonates or infants presenting brain lesions compatible with neuronal fiber loss or defects of neurite outgrowth, such as cortical atrophy, ventricular enlargement, demyelination or gray and white matter hypodensities (Msall et al., 1984; Harding et al., 1984; Wakamoto et al., 1999), which can be acquired in utero (Filloux et al., 1986).

Understanding ammonium toxicity to neurons implied to identify impaired NF-M positive fibers. NN-18 and RMO-44 anti-NF-M antibodies are directed against phosphorylation independent epitopes and both equally recognize NF-M in denaturing SDS-PAGE conditions (Lee et al., 1987; Harris et al., 1991; and this work). On cryosections however, we showed that NN-18 stained fibers but not neuronal cell bodies, whereas RMO-44 stained neuronal soma, but not fibers. We showed moreover that protein dephosphorylation on cryosections suppressed the staining of fibers by NN-18 but induced it by RMO-44. Thus, both antibodies recognize non-phosphorylated epitopes that are exposed or not depending on changes of NF-M conformation due to its level of phosphorylation. On histological sections, the epitope
recognized by NN-18 appears to be exposed only when NF-M is phosphorylated, whereas that recognized by RMO-44 seems exposed only when NF-M is non-phosphorylated. Accordingly, fibers at aggregate periphery preferentially contain phosphorylated NF-M, whereas neuronal soma essentially contain non-phosphorylated NF-M. Our findings may explain why NN-18 does not stain neuronal soma except in case of hyperphosphorylated NF-M accumulation (Harris et al., 1991; Nguyen et al., 2001), and why RMO-44 preferentially labels neuronal soma (Lee et al., 1987).

NFs are distributed in neuronal soma, dendrites and axons. Non-phosphorylated NFs are predominantly found in the somato-dendritic compartment, whereas phosphorylated NFs are enriched in axons (Rosenstein and Krum, 1996; Brown, 1998; Ulfig et al., 1998). Moreover, GAP43 and ChAT showed a localization similar to that of NN-18 stained NF-M, and reacted like it under NH$_4$Cl and/or Cr exposure. GAP43 is found in growth cones of axons exclusively (Goslin et al., 1988), and ChAT is recognized as presynaptic axonal marker (Phelps et al., 1985). Taken together, these data suggest that NN-18 stained axons specifically.

Cholinergic, GABA-ergic and glutamatergic neurons differentiate in the brain aggregate culture system (Pardo and Honegger, 1999; Honegger and Monnet-Tschudi, 2001). Data presented here suggest that early in development, axons impaired by NH$_4$Cl belong to cholinergic neurons. This does not exclude that ammonium could also impair axonal growth of glutamatergic and GABA-ergic neurons later-on in development, as glutamatergic neurotransmission and GAD activity are altered by hyperammonemia (Albrecht, 1998; Braissant et al., 1999b). Our findings are in line with data obtained in the spf mouse, a model of hyperammonemia caused by ornithine transcarbamylase deficiency, which shows
cholinergic neuronal loss in cerebral cortex (Ratnakumari et al., 1994). Lack of elongation of cholinergic axons during cortical development may provide a basis for understanding some of the severe cognitive defects caused by hyperammonemia.

**NH₄Cl induced inhibition of axonal growth depends on intracellular Cr**

NH₄Cl exposure of day 13 mixed-cell aggregates impaired axonal growth and decreased intracellular levels of Cr, PCr and ADP. Cr co-treatment under NH₄Cl exposure protected axonal growth, but neither restored nor increased PCr, ATP and ADP levels. This suggests that under NH₄Cl exposure, the rescue of axonal growth by Cr does not depend on high energy phosphates, which remain lowered under NH₄Cl+Cr exposure and are known to decrease in the hyperammonemic CNS (Ratnakumari et al., 1992). We cannot exclude however that concentration changes in specific cell types or in subcellular compartments (e.g. mitochondria) remain undetected. Intracellular Cr was decreased by 60% in neuron-enriched but only by 25% in mixed-cell aggregates exposed to NH₄Cl+Cr as compared to Cr exposure, suggesting that NH₄Cl exposure reduced the capacity to accumulate Cr in neurons preferentially.

**Glial dependency of axonal protection by creatine under NH₄Cl exposure**

Cr added to the culture medium was sufficient to protect axonal growth in NH₄Cl exposed mixed-cell aggregates, but not in neuron-enriched cultures, suggesting that the mechanism of axon protection by Cr depends on glial cells, be it astrocytes or oligodendrocytes. In case of an astrocyte dependent mechanism, the very few astrocytes still present in neuron-enriched cultures might suffice for supporting axonal growth in control conditions, but not for protecting axons exposed to NH₄Cl. The absence of oligodendrocytes in neuron-enriched cultures might also suggest an oligodendrocyte dependent mechanism. Since Cr is taken up
by neuron-enriched cultures without protecting axons exposed to NH₄Cl, it is unlikely that Cr per se is the sole glia-derived axonal growth promoting factor. Our findings preferentially support the hypothesis that a glial factor is needed, which is modified through Cr in glial cells (e.g. by the Cr/PCr/CK system), released and used by neurons to promote axonal growth. This would still allow axonal growth in control neuron-enriched cultures, as they are grown in a culture medium conditioned by mixed-cell cultures (Honegger and Monnet-Tschudi, 2001).

As we have shown that axons do not grow in neuron-enriched aggregates exposed to ammonium, it also implies that NH₄Cl has a direct neuronal inhibitory effect on axonal growth that cannot be prevented in absence of glial cells, but is counteracted by Cr co-treatment in mixed-cell aggregates.

The glial mechanism of axonal growth protection might involve protein phosphorylation, which is directly linked to cell content in high energy phosphates and Cr/PCr/CK system, altered in glial cells under ammonium exposure (Neary et al., 1987; Schliess et al., 2002) and was proposed as a main signaling pathway in axon/glia inter-relationships (Witt and Brady, 2000). Cr dependent modification of glial protein phosphorylation could support axonal growth, which per se is regulated through phosphorylation of axonal cytoskeleton proteins (e.g. NFs) under direct influence of oligodendrocytes (de Waegh et al., 1992). Such a mechanism is supported by our data, namely the NH₄Cl induced inhibition of NF-M phosphorylation that is counteracted by a Cr co-treatment.

**NH₄Cl exposure does not alter axonal morphology in mature aggregates**

We have shown that ammonium impaired axons in developing aggregates, but did not affect them in mature cultures. The absence of NH₄Cl effect on mature axons could be due to fully differentiated astrocytes, which have a higher capacity for ammonium detoxification.
(Butterworth, 1993). Alternatively, protective factors released by target postsynaptic neurons to their presynaptic counterparts could be involved (Keith and Wilson, 2001). This difference in vulnerability of the brain is also found clinically, as hyperammonemia causes irreversible CNS symptoms compatible with axonal loss in neonates and infants but not in adults (Brusilow and Horwich, 2001). Evidence for direct coupling of Cr/PCr/CK system to growth cone activity and axonal growth (Wang et al., 1998) and Cr/PCr age dependency of CNS in the first weeks of postnatal life (Tsuji et al., 1995) also support the difference we found between developing and mature aggregates. It suggests a higher CNS sensitivity to Cr variations in young children than in adulthood.

The brain cell aggregate culture system as a model to study CNS hyperammonemia

Axonal growth was altered in developing brain cell aggregates by 5 mM NH₄Cl exposure. This concentration mimicked the in vivo extracellular ammonium level found in the brain of experimental hyperammonemic rats, which was measured as high as 5mM (Swain et al., 1992). For human hyperammonemic patients, data are lacking on extracellular brain ammonium concentration. However, serum levels of ammonium leading to irreversible damage to the developing CNS can peak as high as 2mM, usually following a chronic hyperammonemia in the range of 200µM (Butterworth, 1998; Bachmann, 2002a). Brain cell aggregate cultures fulfilled the requirements that irreversible ammonium toxicity to CNS should be studied in models mimicking brain complexity (Butterworth, 1998; Braissant et al., 1999a and b) but devoid of confusing variables due to secondary effects of hyperammonemia found in animal studies (Bachmann, 1992). Our findings are in agreement with data showing that hyperammonemia stimulates brain glycolysis with release of lactate into CSF and causes ammonium detoxification through astrocytic glutamine synthesis, which can lead to gliosis and occurrence of Alzheimer type II astrocytes (Butterworth, 1998). Moreover, GAP43, an
exclusive marker of axonal growth cone in physiological conditions, was induced in the GFAP positive border of NH₄Cl exposed aggregates, as in reactive astrocytes (Sensenbrenner et al., 1997).

**Conclusions**

We have shown that glial Cr can protect axonal growth under ammonium exposure. This work exemplifies the importance of neuron-glial interactions in the pathophysiology of hyperammonemia. Future work will aim at identifying those factors modified through glial Cr that support axonal growth and are impaired by hyperammonemia, and at understanding changes in brain Cr metabolism and transport under ammonium exposure. This study also suggests that one should assess means for sustaining the CNS Cr level of hyperammonemic neonates and infants, to prevent irreversible brain damage due to impairment of axonogenesis.
References


Figure legends

Figure 1: Axons are missing and NF-M is decreased in NH₄Cl exposed day 13 mixed-cell aggregates. Creatine, added to NH₄Cl exposure, rescues axons and NF-M.

Cultures grown from days 5 to 13 with or without 5 mM NH₄Cl and/or 1 mM Cr.

Cryosections stained by hematoxylin (Hem.; A,B,F,G,K,L,P,Q), Bielschowsky’s silver impregnation (Silver; C,H,M,R) and immunohistochemistry against NF-M (D,I,N,S: NN-18 antibody; E,J,O,T: RMO-44 antibody). A-E: Controls; F-J: Cr; K-O: NH₄Cl; P-T: NH₄Cl+Cr. Panels B-E, G-J, L-O and Q-T are higher magnifications of the area depicted in A. The dense peripheral axonal zone is indicated by an asterisk. Bar: 100 µm.

U,V: NF-M analysis by western blotting, with NN-18 (U; 20 µg proteins per lane) and RMO44 (V; 10 µg protein per lane) antibodies. Means ± SD of 3 separate cultures.

Figure 2: NN-18 and RMO-44 anti-NF-M antibodies recognize phosphorylated, respectively non-phosphorylated, NF-M in native conformation. Mixed-cell aggregates at 13 days of culture. A,C: Controls; B,D: Dephosphorylation by alkaline phosphatase.

Cryosections were stained by immunohistochemistry against NF-M (A,B: NN-18 antibody; C,D: RMO-44 antibody). Arrow: neuronal cell body; arrowhead: proximal axon, arising from neuronal cell body and growing towards aggregate periphery; asterisk: peripheral zone enriched in axons. Bar: 100 µm.
Figure 3: GAP43, an axonal growth cone marker, is decreased by NH₄Cl exposure, and rescued by Cr co-treatment, in the peripheral axonal zone of the aggregates. Ammonium exposure induces GAP43 and increases GFAP in reactive astrocytes of the aggregate border.

Cultures treated from days 5 to 13 with or without 5mM NH₄Cl and/or 1mM Cr.

Cryosections stained by immunohistochemistry against GAP43 (A,D,G,J), GFAP (B,E,H,K) and MBP (C,F,I,L). A,D,G,J were counterstained by hematoxylin. A-C: Controls; D-F: Cr; G-I: NH₄Cl; J-L: NH₄Cl+Cr. The dense peripheral axonal zone is indicated by an asterisk. Highly GFAP-positive astrocytes in the border of aggregates are shown by brackets. Bar: 100 µm.

M,N: GAP43 (M) and GFAP (N) analysis by western blotting (10 µg proteins per lane). Means ± SD of 3 separate cultures.

Figure 4: Cr, added to NH₄Cl exposed day 13 neuron-enriched aggregates, do not rescue developing axons and NF-M levels.

Cultures treated from days 5 to 13 with or without 5mM NH₄Cl and/or 1mM Cr.

Cryosections stained by hematoxylin (A,F,K,P) and immunohistochemistry against NF-M (B,G,L,Q: NN-18 antibody; C,H,M,R: RMO-44 antibody), GFAP (D,I,N,S) and MBP (E,J,O,T). A-E: Controls; F-J: Cr; K-O: NH₄Cl; P-T: NH₄Cl+Cr. The dense peripheral axonal zone is indicated by an asterisk. Arrows point to GFAP-positive giant astrocytes in NH₄Cl exposed cultures. Bar: 100 µm.

U,V: NF-M analysis by western blotting, with NN-18 (U; 20 µg proteins per lane) and RMO44 (V; 10 µg protein per lane) antibodies. Means ± SD of 3 separate cultures.
**Figure 5: Cholinergic axons of day 13 mixed-cell aggregates are affected by NH₄Cl exposure and rescued by Cr co-treatment.**

Aggregates were treated from day 5 to day 13 with or without 5mM NH₄Cl and/or 1mM creatine.

Immunohistochemistry against ChAT (A-D) and GAD 65/67 kD (E,F). A,E: controls; B,F: NH₄Cl; C: Cr; D: NH₄Cl+Cr. The dense peripheral axonal zone is indicated by an asterisk. Bar: 100 μm.

**Figure 6: NH₄Cl and Cr do not affect axons and NF-M in day 28 mature mixed-cell aggregates.**

Cultures were treated from days 20 to 28 with or without 5mM NH₄Cl and/or 1mM Cr.

A,B: Hematoxylin staining of day 28 control aggregates at low (A) and high (B) magnification. C-F: Immunohistochemistry against NF-M (NN-18 antibody). C: control; D: NH₄Cl; E: Cr; F: NH₄Cl+Cr. Panels B-F are higher magnifications of the area depicted in A. The dense peripheral axonal zone is indicated by an asterisk. Bar: 100 μm.

Table 1: NH₄Cl dose dependent uptake and release of ammonium, glucose and lactate by mixed-cell aggregate cultures.a.

<table>
<thead>
<tr>
<th>NH₄Cl [mM]</th>
<th>Ammonium</th>
<th>Glucose</th>
<th>Lactate</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Days 5 =&gt; 8</td>
<td>Days 8 =&gt; 11</td>
<td>Days 11 =&gt; 13</td>
</tr>
<tr>
<td>0</td>
<td>-26 ± 1</td>
<td>-4 ± 1</td>
<td>-2 ± 1</td>
</tr>
<tr>
<td>1.0</td>
<td>-55 ± 13 &lt;0.001</td>
<td>-31 ± 4 &lt;0.001</td>
<td>-38 ± 1 &lt;0.001</td>
</tr>
<tr>
<td>2.5</td>
<td>-113 ± 19 &lt;0.001</td>
<td>-96 ± 2 &lt;0.001</td>
<td>-114 ± 1 &lt;0.001</td>
</tr>
<tr>
<td>5.0</td>
<td>-176 ± 16 &lt;0.001</td>
<td>-250 ± 2 &lt;0.001</td>
<td>-264 ± 10 &lt;0.001</td>
</tr>
<tr>
<td>0</td>
<td>-399 ± 19</td>
<td>-464 ± 12</td>
<td>-756 ± 4</td>
</tr>
<tr>
<td>1.0</td>
<td>-372 ± 32 NS</td>
<td>-389 ± 22 &lt;0.01</td>
<td>-656 ± 48 &lt;0.05</td>
</tr>
<tr>
<td>2.5</td>
<td>-525 ± 32 &lt;0.01</td>
<td>-468 ± 7 NS</td>
<td>-835 ± 17 &lt;0.01</td>
</tr>
<tr>
<td>5.0</td>
<td>-672 ± 21 &lt;0.001</td>
<td>-623 ± 19 &lt;0.001</td>
<td>-1026 ± 7 &lt;0.001</td>
</tr>
<tr>
<td>0</td>
<td>137 ± 5</td>
<td>166 ± 6</td>
<td>685 ± 26</td>
</tr>
<tr>
<td>1.0</td>
<td>152 ± 18 NS</td>
<td>149 ± 39 NS</td>
<td>549 ± 21 &lt;0.01</td>
</tr>
<tr>
<td>2.5</td>
<td>269 ± 22 &lt;0.001</td>
<td>153 ± 22 NS</td>
<td>696 ± 73 NS</td>
</tr>
<tr>
<td>5.0</td>
<td>431 ± 34 &lt;0.001</td>
<td>370 ± 41 &lt;0.001</td>
<td>1137 ± 22 &lt;0.001</td>
</tr>
</tbody>
</table>

a Net uptake (net decrease in culture medium) and release (net increase in culture medium) by aggregates, calculated from days 5 to 8, 8 to 11 and 11 to 13, and expressed as nmol · h⁻¹ · (mg prot)⁻¹. Measures were taken at days 5 (start of treatment), 8 (before and after medium change), 11 (before and after medium change) and 13 (end of treatment). Proteins were measured at day 13. Data are means ± SD of 3 separate cultures. Statistical p are shown (Student’s t-test) for comparison to controls (NH₄Cl = 0 mM); NS: not significant.
Table 2. Intracellular creatine, phosphocreatine, ATP, ADP and AMP in mixed-cell aggregates at day 13 of culture a.

<table>
<thead>
<tr>
<th>Treatments</th>
<th>Control</th>
<th>Cr</th>
<th>NH₄Cl</th>
<th>NH₄Cl+Cr</th>
<th>Comparisons</th>
<th>NH₄Cl vs. Controls</th>
<th>NH₄Cl+Cr vs. NH₄Cl</th>
</tr>
</thead>
<tbody>
<tr>
<td>NH₄Cl [mM]</td>
<td>0</td>
<td>0</td>
<td>5</td>
<td>5</td>
<td></td>
<td>&lt; 0.01</td>
<td>&lt; 0.01</td>
</tr>
<tr>
<td>Creatine [mM]</td>
<td>178.6 ± 5.8</td>
<td>340.2 ± 3.5</td>
<td>147.7 ± 4.1</td>
<td>255.5 ± 25.6</td>
<td></td>
<td>&lt; 0.01</td>
<td></td>
</tr>
<tr>
<td>Phosphocreatine</td>
<td>13.0 ± 0.4</td>
<td>12.1 ± 0.3</td>
<td>9.7 ± 0.8</td>
<td>9.8 ± 0.5</td>
<td></td>
<td>&lt; 0.05</td>
<td>NS</td>
</tr>
<tr>
<td>ATP</td>
<td>5.5 ± 0.7</td>
<td>7.0 ± 0.9</td>
<td>4.5 ± 2.1</td>
<td>3.1 ± 0.2</td>
<td></td>
<td>NS</td>
<td>NS</td>
</tr>
<tr>
<td>ADP</td>
<td>39.3 ± 2.3</td>
<td>37.6 ± 2.9</td>
<td>27.2 ± 1.5</td>
<td>29.1 ± 3.8</td>
<td></td>
<td>&lt; 0.01</td>
<td>NS</td>
</tr>
<tr>
<td>AMP</td>
<td>17.2 ± 0.4</td>
<td>18.8 ± 1.4</td>
<td>19.6 ± 7.3</td>
<td>15.3 ± 4.2</td>
<td></td>
<td>NS</td>
<td>NS</td>
</tr>
</tbody>
</table>

aData are expressed as nmol · (mg prot)¹ and are means ± SD of 3 separate cultures. Statistical p are shown (Student’s t-test; NS: not significant).
Table 3: Intracellular creatine, phosphocreatine, ATP, ADP and AMP in neuron-enriched aggregates at day 13 of culture.

<table>
<thead>
<tr>
<th>Treatments</th>
<th>Control</th>
<th>Cr</th>
<th>NH₄Cl</th>
<th>NH₄Cl+Cr</th>
<th>Comparisons</th>
<th>Comparisons</th>
</tr>
</thead>
<tbody>
<tr>
<td>NH₄Cl [mM]</td>
<td>0</td>
<td>0</td>
<td>5</td>
<td>5</td>
<td>NH₄Cl vs. Controls</td>
<td>NH₄Cl+Cr vs. NH₄Cl</td>
</tr>
<tr>
<td>Creatine [mM]</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>1</td>
<td>NS</td>
<td>&lt; 0.01</td>
</tr>
<tr>
<td>NH₄Cl+Cr</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>NS</td>
<td></td>
</tr>
<tr>
<td>Phosphocreatine</td>
<td>4.4 ± 1.9</td>
<td>6.8 ± 2.1</td>
<td>4.1 ± 0.7</td>
<td>4.8 ± 1.6</td>
<td>NS</td>
<td></td>
</tr>
<tr>
<td>ATP</td>
<td>6.7 ± 0.9</td>
<td>9.1 ± 0.6</td>
<td>4.5 ± 0.6</td>
<td>4.3 ± 0.8</td>
<td>&lt; 0.05</td>
<td>NS</td>
</tr>
<tr>
<td>ADP</td>
<td>3.3 ± 0.4</td>
<td>3.7 ± 0.3</td>
<td>1.9 ± 0.2</td>
<td>2.0 ± 0.4</td>
<td>&lt; 0.05</td>
<td>NS</td>
</tr>
<tr>
<td>AMP</td>
<td>7.6 ± 1.1</td>
<td>9.4 ± 2.0</td>
<td>3.8 ± 0.4</td>
<td>3.7 ± 0.1</td>
<td>&lt; 0.01</td>
<td>NS</td>
</tr>
</tbody>
</table>

aData are expressed as nmol · (mg prot)⁻¹ and are means ± SD of 3 separate cultures. Statistical p are shown (Student’s t-test; NS: not significant).
Creatine 

\[ \text{ChAT} - \text{NH}_4\text{Cl} + \]

A  

B  

Creatine 

\[ + \]

C  

D  

\[ \text{GAD} - \text{NH}_4\text{Cl} + \]

E  

F  

\[ \text{**} \]
Hematoxylin

A

B

NF-M (NN-18) - NH₄Cl +

C

D

Creatine -

E

F

+ Cr 1 mM -+ -+

NF-M (NN 18) NH₄Cl 5 mM -- + +

G

Volume AU

0 500 1000 1500 2000

NF-M (NN 18)

NH₄Cl 5 mM - - + +

Cr 1 mM - + - +