Delayed post-ischaemic neuroprotection following systemic neural stem cell transplantation involves multiple mechanisms

Marco Bacigaluppi,1,2,3,*† Stefano Pluchino,2,3,*† Luca Peruzzotti Jametti,2,3,† Ertugrul Kilic,1 Ülkan Kilic,1 Giuliana Salani,2,† Elena Brambilla,2 Mark J. West,4 Giancarlo Comi,2,3 Gianvito Martino2,3,* and Dirk M. Hermann1,5,*

1 Department of Neurology, University Hospital Zurich, Frauenklinikstrasse 26, CH-8091 Zurich, Switzerland
2 Neuroimmunology Unit-DIBIT2 and Institute of Experimental Neurology, San Raffaele Scientific Institute, Via Olgettina 58, I-20132 Milan, Italy
3 Department of Neurology and Neurophysiology, San Raffaele Scientific Institute, Via Olgettina 58, I-20132 Milan, Italy
4 Institute of Anatomy, University Park, DK-8000 Aarhus, Denmark
5 Department of Neurology, University Hospital, University of Duisburg-Essen, Hufelandstrasse 55, D-45122 Essen, Germany

*These authors contributed equally to this work.
†Present address: CNS Repair Unit-DIBIT2, and Institute of Experimental Neurology, San Raffaele Scientific Institute, Via Olgettina 58, I-20132 Milan, Italy

Correspondence to: Prof. Dr Dirk M. Hermann, Chair of Vascular Neurology, Dementia and Ageing Disorders, Department of Neurology, University Hospital Essen, Hufelandstr. 55, D-45122 Essen, Germany
E-mail: dirk.hermann@uk-essen.de

Recent evidence suggests that neural stem/precursor cells (NPCs) promote recovery in animal models with delayed neuronal death via a number of indirect bystander effects. A comprehensive knowledge of how transplanted NPCs exert their therapeutic effects is still lacking. Here, we investigated the effects of a delayed transplantation of adult syngenic NPCs—injected intravenously 72 h after transient middle cerebral artery occlusion—on neurological recovery, histopathology and gene expression. NPC-transplanted mice showed a significantly improved recovery from 18 days post-transplantation (dpt) onwards, which persisted throughout the study. A small percentage of injected NPCs accumulated in the brain, integrating mainly in the infarct boundary zone, where most of the NPCs remained undifferentiated up to 30 dpt. Histopathological analysis revealed a hitherto unreported very delayed neuroprotective effect of NPCs, becoming evident at 10 and 30 dpt. Tissue survival was associated with downregulation of markers of inflammation, glial scar formation and neuronal apoptotic death at both mRNA and protein levels. Our data highlight the relevance of very delayed degenerative processes in the stroke brain that are intimately associated with inflammatory and glial responses. These processes may efficaciously be antagonized by (stem) cell-based strategies at time-points far beyond established therapeutic windows for pharmacological neuroprotection.
Introduction

According to current pathophysiological concepts, the structural histological injury following stroke evolves within several hours to up to 3–4 days, depending on the duration and severity of the ischaemic event (Namura et al., 1998; Hata et al., 2000; Hermann et al., 2001). During that time, secondary energy failure develops in the ischaemic penumbra via multiple mechanisms including excitotoxicity, lactacidosis and peri-infarct depolarizations (Hossmann, 2006). In the ischaemic border zone surrounding the evolving infarct, inflammatory responses and apoptotic programs are activated in this phase that further contribute to injury development (Dirmagl et al., 1999; Hermann et al., 2001; Hossmann, 2006). Following the acute stroke phase, the tissue undergoes substantial matrix remodelling that results in (astro)glial scar formation, which is widely regarded as a major hampering factor for tissue repair and regeneration (Fawcett and Asher, 1999; Nedergaard and Dirmagl, 2005; Yiu and He, 2006).

Several lines of evidence suggest that the structural and functional changes in the ischaemic boundary zone are of key importance for the final stroke outcome (Witte, 1998; Nudo, 1999). This might imply that therapies favouring endogenous tissue repair and/or remodelling may have realistic chances of success also when applied beyond established time-windows of tissue protection, which in the case of thrombolytics in humans are in the range of 3–6 h (Hacke et al., 2008). The pathophysiology of this second phase is highly heterogeneous in time and space, and the underlying mechanisms are still poorly understood (Lee et al., 2008). There is common sense that classical pharmacological strategies aiming at improving neuronal survival are no more useful at this latter stage (Zivin, 1998).

Neural stem/precursor cells (NPCs) display remarkable therapeutic plasticity upon transplantation in experimental conditions mimicking central nervous system (CNS) diseases, by adapting their fate and function(s) to specific environments under pathological conditions (Martino and Pluchino, 2006). Key functions such as the replacement of neural cells (Pluchino et al., 2003; Karimi-Abdolrezaee et al., 2006) and the delivery of therapeutic gene(s) (Lee et al., 2007) have been recently challenged by intrinsic bystander capacities, mainly exerted by undifferentiated NPCs, releasing at the site of tissue damage, a milieu of immune-regulatory molecules that is temporally and spatially orchestrated by specific environmental needs (Pluchino et al., 2005). Furthermore, NPCs synergize with local immune (e.g. T cells and microglia) (Ziv et al., 2006) and CNS resident (e.g. endothelial cells, astrocytes) cells (Pluchino et al., 2005), modulating the focal release of stem cell regulators and—in turn—promoting functional recovery from CNS injuries. The molecular and cellular mechanisms mediating such bystander effects still remain to be characterized.

The brain repair potential of NPCs acutely transplanted in stroke-like conditions—both ischaemic and haemorrhagic—has solid pre-clinical evidence (Bliss et al., 2007; Lee et al., 2008). Thus, the issue as whether (and how) NPCs also modulate delayed cerebral responses after stroke has important therapeutic relevance that should be clarified in order to correctly adjust cell-based therapies to tissue needs. Here, we report the capacity of the post-acute transplantation of syngenic adult NPCs injected intravenously (i.v.) at 72 h post-ischaeemia to induce functional neurological recovery in mice with transient middle cerebral artery occlusion (MCAO). Histological analysis revealed a so far unreported very delayed neuroprotective effect that began to evolve at 10 days post-transplantation (dpt) and was even more pronounced at 30 dpt. Molecular biological and histochemical studies revealed profound anti-inflammatory, glial scar-inhibitory and anti-apoptotic effects of NPCs, which were responsible for the neuroprotection and the functional improvements.

Materials and Methods

Intraluminal MCAO

Details on the study design are presented in Supplementary Fig. 1. Forty-five minutes of MCAO were induced in adult male 8- to 10-week old (weight 20–25 g) C57Bl/6 mice (Harlan Nossan, Netherlands), as previously described (Hata et al., 2000; Hermann et al., 2001). Experiments were performed at the University Hospital Zurich according to the National Institutes of Health guidelines for the care and use of laboratory animals with approval of local government authorities (Cantonal Veterinary Office, ZH 169/2005). During the experiments, up to 15 min after reperfusion, laser Doppler flow (LDF) was monitored above the core of the middle cerebral artery territory. Further information is provided in the Supplementary materials and Methods section.

NPC preparation and transplantation

Adult neurospheres were generated from the subventricular zone of 8-month-old C57Bl/6 mice, as described (Pluchino et al., 2005), and i.v. delivered at 72 h after reperfusion via the tail vein. Further information is provided in the Supplementary Materials and Methods section.

Behavioural analysis

Modified neurological severity score (mNSS) was evaluated at baseline, on the day of transplantation prior to cell delivery, on a daily basis between 1 and 11 dpt, and then every second to fourth day up to 30 dpt. Grip strength test was performed at baseline, at 3 days after MCAO, immediately before cell transplantation, and at 3, 10, 20 and 30 dpt. Behavioural tests were performed during the light phase of the circadian cycle beginning 4 h after lights on. Further
information is provided in the Supplementary Materials and Methods section.

**Tissue pathology**

At sacrifice, mice were transcardially perfused with 4% paraformaldehyde and brain, liver, spleen, kidneys and lungs were removed and processed for tissue histopathology as frozen tissue samples (Pluchino et al., 2005). Further information is provided in the Supplementary Materials and Methods section.

**Stereological analysis**

In the striatum, cell numbers for NeuN, Darp-32 and D2R were quantified according to the optical fractionator method with the assistance of the Stereo Investigator v 3.0 software (MicroBrightField, Inc., Colchester, VT, USA) (West et al., 1991).

Estimates of the length of capillaries (threshold smaller than 8 \( \mu \text{m} \)) were made throughout the striatum (ischaemic and contralateral) on CD 31 stained sections with the use of virtual isotropic hemispherical probes (radius = 22 \( \mu \text{m} \)) on the Stereo Investigator v 3.0 software (MicroBrightField) (Mouton et al., 2002). Further information is provided in the Supplementary Materials and Methods section.

**Real-time PCR**

Real-time quantitative PCR was performed using pre-developed Taqman\textsuperscript{TM} Assay Reagents on an ABI Prism\textsuperscript{TM} 7700 Sequence Detection System (Applied Biosystems, Carlsbad, California, USA) according to the manufacturer’s protocol. Further information is provided in the Supplementary Materials and Methods section.

**Statistical analysis**

For statistical analyses, we used a standard software package (GraphPad Prism version 4.00). To test the treatment effect on each of the behaviour scores, behavioural tests were evaluated by means of repeated measurement analysis of variance (ANOVA). Whenever a treatment by time interaction or treatment effect was present at the 0.05 level, a post hoc analysis was done by Bonferroni test. Histological data were evaluated by unpaired two-tailed \( t \)-tests. Gene expression analyses were evaluated by two-tailed \( t \)-tests and by two-way ANOVA followed by Bonferroni test.

**Results**

**Progressive improvement of neurological deficits after NPC transplantation**

Mice subjected to transient MCAO (Hata et al., 2000) were injected i.v. with syngenic subventricular zone-derived green fluorescent protein (GFP)\textsuperscript{+} adult NPCs (1.0 \( \times \) 10\textsuperscript{6} cells per mouse) at 72 h after reperfusion, following a delayed (post-acute) cell transplantation scheme (Pluchino et al., 2003, 2005). Post-randomization analysis of the LDF measurements, weight loss and neurological deficit scores (mNSS and grip strength test) both prior to the induction of cerebral ischaemia (−3 dpt) and before treatment (Day 0) did not show any differences between the groups (Fig. 1A−D).

Neurological performance was monitored all along the follow-up until 30 dpt. Indeed, remarkable development of neurological deficits was observed upon MCAO in both groups of mice. NPC-transplanted MCAO mice showed a progressively enhanced recovery of motor skills in the mNSS—starting to be significant from 18 dpt onwards (\( P \leq 0.05 \))—as compared with sham-treated controls. This improvement was also reflected by stronger grip strength in the right paretic forepaw (Fig. 1 C–D). The grip strength in the left non-paretic forepaw did not differ between the groups (data not shown).

**Accumulation and long-term persistence of transplanted NPCs in the ischaemic boundary zone**

To assess the distribution and the identity of i.v.-injected NPCs in the ischaemic brain, a detailed morphometric analysis was carried out at 3, 10 and 30 dpt in the brain and in peripheral tissues such as spleen, liver, kidneys and lungs. Only 0.09% of injected NPCs were detected in the brain by 3 dpt and the cells accumulated in both hemispheres, ipsilateral and contralateral to the stroke (3.47 ± 0.96 and 4.25 ± 1.6 cells/mm\(^3\), respectively). At later time-points, NPCs were detected exclusively in the ischaemic hemisphere. As such, the number of NPCs increased by 5.5-fold at 10 dpt (0.23% of transplanted NPCs, 19.4 ± 4.87 cells/mm\(^3\), \( P \leq 0.05 \) compared with 3 dpt), the majority GFP\textsuperscript{+} NPCs accumulating in the ischaemic boundary zone. At 30 dpt numbers of NPCs remained similarly high (0.28% of transplanted NPCs, 20.5 ± 4.2 cells/mm\(^3\), \( P \leq 0.05 \) compared with 3 dpt). Sham-treated ischaemic mice did not display any immunoreactivity for GFP, neither in the brain nor in peripheral tissues (Fig. 2A and B).

To characterize the proliferation and differentiation capacity of transplanted NPCs, different immunohistochemical stainings were performed. Twenty-five percent of GFP\textsuperscript{+} cells at 3 dpt expressed the Ki67 antigen, a molecular marker that is expressed exclusively in proliferating cells during the late G\(_1\), S, G\(_2\) or M phase of the cell cycle. The great majority of NPCs within the brain at 3 and 10 dpt did not express any of the major antigens of the neural lineage, such as glial fibrillary acidic protein, doublecortin (Dcx), microtubule-associated protein (MAP)-2 and the oligodendroglial transcription factor (Olig 2) (data not shown). The percentage of Ki67\textsuperscript{+} NPCs decreased at later time-points (6.9% at 10 dpt, 0.8% at 30 dpt), whereas the majority of cells still remained undifferentiated. Even at 30 dpt, only 4.4% and 0.8% of i.v.-injected NPCs at 30 dpt expressed Olig2 and doublecortin, respectively (Fig. 2C–F).

Similarly to what others and we have shown in experimental autoimmune encephalomyelitis, NPCs detected in the ischaemic boundary zone established anatomical interaction(s) with von Willebrand factor\textsuperscript{+} endothelial cells, CD45\textsuperscript{+} blood-derived leucocytes and f4/80\textsuperscript{+} phagocytes. Occasionally, f4/80\textsuperscript{+} phagocytes being immunoreactive also for GFP were identified, thus
suggesting that only very low numbers of i.v.-injected NPCs might have been phagocytosed in vivo (Fig. 2G–H).

Few scattered GFP+ NPCs were found in the spleen, liver, kidneys and lungs up to 30 dpt (Supplementary Fig. 2). No pathological signs suggestive of overt toxic effects (e.g. inflammation, necrosis, cell degeneration) were found in any of these organs.

Protection from delayed ischaemic injury and enhancement of neuronal survival

To understand how NPCs delivery improved functional recovery after stroke, we analysed the treatment effect on histopathological brain injury. Cresyl violet-based morphometrical analysis of brain swelling or atrophy showed no difference, when NPC-treated MCAO mice were compared with sham-treated controls at both 0, 3 and 10 dpt (Fig. 3A and C). In contrast, significant reduction of the lesion volume by 21.5% was detected at 30 dpt in NPC-treated mice (P < 0.05) (Fig. 3B–E). The regional analysis of the surviving tissue indicated that NPC effect on lesion size was due to a protection of residual striatal brain tissue (total striatal volume in the ischaemic hemisphere: sham treated, 1.99 ± 0.20 mm3; NPC treated, 2.59 ± 0.20 mm3, P = 0.05; 23% lesion volume reduction). The morphometric analysis of the surviving cortical tissue showed no significant difference between the groups.

The total lesion volume was found to be inversely correlated with grip strength at 30 dpt (r = −0.877, r² = 0.769; P = 0.0002), but not to the mNSS.

Modulation of inflammation-, astrogliosis- and neuronal survival-related genes at mRNA level

In order to elucidate the molecular mechanism(s) underlying the observed NPC-mediated protection from brain ischaemic injury progression, we next performed a wide TaqMan®-based semi-quantitative gene expression profiling in the two brain hemispheres (both ipsi- and contralateral to the stroke) from sham-treated and NPC-treated MCAO mice at 0, 3, 10 and 30 dpt.

Forty-five different mRNA species involved in angiogenesis, astrogliosis, inflammation and neuronal survival were measured in the brain. Significant and homogeneous downregulation of several mRNA species was observed between 3 and 30 dpt in NPC-treated MCAO mice, as shown by progressive decrease of the regression curve slope values (m) at 3 (m = 1.15), 10 (m = 0.67) and 30 dpt (m = 0.57), respectively. Further correlation index analysis on scatter plots revealed that differences in gene expression were mostly seen in the ischaemic hemispheres at
10 dpt ($r^2 = 0.82$), when compared with 3 and 30 dpt ($r^2 = 0.96$ and 0.93, respectively) (Supplementary Fig. 3), which is notably significant before group differences in brain injury became evident. Interestingly, 42.2% (19/45) of the genes did not change at any time-point. Only 2.2% (1/45) of the genes were upregulated in NPC-treated mice at 3 dpt, among which we identified DARPP-32 (1.7-fold) (Fig. 4A). On the other hand, 55.6% (25/45) of the genes were downregulated in NPC-treated mice. Among these, the most prominent were (i) inflammatory regulators [e.g. interferon (IFN)-γ at 3 dpt (0.4-fold, $P \leq 0.05$) and at 10 dpt (0.6-fold), tumour necrosis factor (TNF)-α (0.5-fold, $P \leq 0.05$) at 10 dpt, interleukin (IL)-1β (0.5-fold, $P \leq 0.05$), IL-6 (0.5-fold, $P \leq 0.05$) and leptin receptor (0.7-fold, $P = 0.07$) at 30 dpt]; (ii) regulators of (astro)glial proliferation and reactivity [e.g. fibroblast growth factor (FGF)-II (0.7-fold, $P = 0.01$) at 10 dpt, the marker of reactive astrocytes vimentin (0.4-fold, $P \leq 0.05$) (Galou et al., 1996) at 30 dpt] and (iii) neuronal death and plasticity [e.g. the executioner caspase-3 (0.7-fold, $P \leq 0.05$) at 10 dpt, the growth-associated protein (GAP)-43 (0.6-fold, $P \leq 0.01$) and the chondroitin sulphate proteoglycan versican
(0.7-fold, $P = 0.059$) at 30 dpt (Fig. 4). Genes involved in the angiogenic response to ischaemia (Hif-1α, KDR, EPO, Flt-1, VEGF-α) in the perilesional area changed only slightly after ischaemia and only subtle changes were observed between the two treatment groups. At 30 dpt, stereological-based estimates of the total density and length of capillaries in sham- and NPC-treated mice were not significantly different (total capillary density: 6.751 ± 3.978 vs. 6.682 ± 2.846, $P = 0.890$; total length of capillaries: 1.732 ± 0.85 × 10^6 vs. 2.20 ± 0.85 × 10^6, $P = 0.163$, in sham and NPC-treated mice, respectively). In the contralateral non-ischaemic hemisphere, the large majority of mRNA species (29/45, 64.4%) remained unchanged. A comprehensive description of the gene expression data is shown in the Supplementary Table 1. These data indicate that the delayed i.v. injection of NPCs potently regulates the expression of several mRNA species in the ischaemic brain.

**Attenuation of the inflammatory response**

To further characterize the effects of NPC transplantation on the local inflammatory response, we analysed morphometrically the total brain area stained with the common leucocyte antigen CD45 as well as the density of microglial cells in the ischaemic boundary zone both at 3, 10 and 30 dpt. While NPC-treated mice showed a significant reduction of both CD45+ and ionized calcium binding adaptor molecule 1 (IBA-1+) cells at 30 dpt only (both $P < 0.05$, when compared with sham-treated controls) (Fig. 5A–C and G), a significant reduction of IBA-1+/major histocompatibility complex class II molecule (MHC-II+) cells was observed at both 3 and 10 dpt (both $P < 0.05$) (Fig. 5D–F).

**Inhibition of (astro)glial scar formation**

Gliotic scar formation is regarded as a major factor hampering recovery from stroke (Li et al., 2005; Yiu and He, 2006). Thus, we next measured the area and thickness of the Glial fibrillary acidic protein (GFAP+) glial scar wall in the infarct border zone (Fig. 6A). Again, NPC-treated mice showed a smaller scar area at 30 dpt ($P < 0.05$). When corrected for the border length profile irregularities, this also resulted in a significantly thinner scar ($P < 0.01$) (Fig. 6B–E), the difference among the two treatment groups
groups of mice was particularly evident at the rostrocaudal level of the bregma.

**Promotion of neuronal survival**

Further analysis showed that NPC-treated MCAO mice had higher densities of surviving neuronal nuclear antigen (NeuN) + neurons in the striatum ipsilateral to the stroke starting at 3 and 10 dpt, getting significant at 30 dpt, compared with sham-treated controls (Fig. 7A–C and Supplementary Fig. 4). No differences were seen in other structures, namely in the cortex, hippocampus and thalamus (data not shown). These findings were also substantiated with unbiased stereological counts (Fig. 7D–G). Furthermore, by applying the optical-fractionator methodology at tissues obtained at 30 dpt only, we observed significantly more Darpp-32 + medium spiny neurons, as well as D2R + neurons (a subpopulation of medium spiny neurons) (Kawaguchi, 1997) (Fig. 7F and E, \( P \leq 0.05 \) and \( P \leq 0.01 \), respectively) in the striatum of NPC-treated ischaemic mice, when compared with sham-treated controls. Finally, though NPC-transplanted mice displayed a slightly higher number of ChAT + interneurons (a subpopulation of interneurons particular resistant to ischaemia) (Andsberg et al., 2001), the difference to sham-treated animals was not statistically significant (\( P = 0.33 \)) (Fig. 7G).

The enhanced neuronal survival was also reflected by a significant reduction (\( P \leq 0.05 \)) of the total number of apoptotic cells (in particular neurons) displaying either cleavage of caspase 3 (3 dpt) or DNA fragmentation (3 and 10 dpt) in the ischaemic hemisphere of NPC-treated MCAO mice, when compared with sham-treated controls (Fig. 8A and B, Supplementary Fig. 4).

**Prevention of corpus callosum atrophy by NPCs**

There is substantial evidence that ischaemic lesions may induce degeneration of long distance, interhemispheric projections (Napieralski et al., 1996), such as those crossing the corpus callosum (CC). Indeed, the quantification of the medial CC ipsilateral to the stroke revealed that NPC-treated ischaemic mice had a significantly thicker CC at 30 dpt, when compared with sham-treated controls (\( P \leq 0.05 \)) (Fig. 8C). These latter data point towards a more successful remodelling of interhemispheric projections in NPC—as compared with sham-treated MCAO mice.

**Discussion**

In ischaemic stroke the time-window, in which survival promoting therapies are efficacious, is extremely limited. As such, recanalizing drugs (i.e. thrombolytics) (Busch et al., 1998; Kilic et al., 2000; Hacke et al., 2008) as well as neuroprotective compounds [e.g. NMDA receptor antagonists (Ma et al., 1998), free radical scavengers (Yang et al., 2000), growth factors (Cerami, 2001) or caspase-3 inhibitors (Endres et al., 1998)] protect brain tissue only when delivered in the first 1–3 and in some conditions up to 6 h after stroke. That the therapeutic window is so narrow has been explained by the fact that brain infarct evolves very rapidly, typically within a few hours up to 3–4 days after stroke, depending on the duration and severity of ischaemia (Garcia et al., 1993; Namura et al., 1998; Hata et al., 2000; Li et al., 2000; Hermann et al., 2001). In case of ischaemias exhibiting a severity that is close to the threshold at which brain infarcts develop, some
continued injury has also been reported outside that time-window period, i.e. up to 1 week after the ischaemia (Du et al., 1996; Snider et al., 1999; Hermann et al., 2001; Zheng et al., 2003; Guadagno et al., 2008). Whether this delayed injury is relevant for the final functional outcome of stroke remained unclear.

Using a broad set of behavioural neurological, histopathological, immunohistochemical and molecular biological techniques, we have characterized the effects of adult syngenic NPCs i.v.-administered 72 h after reperfusion (i.e. in delayed delivery approach) in MCAO mice. We show that NPCs are capable of integrating into the stroke brain—although at a low percentage (up to 0.28%)—where they are able to survive at least for 30 dpt. As such, the large majority of transplanted cells remained mainly in an undifferentiated state throughout the observation period. This resulted in a hitherto unreported very delayed structural neuroprotective effect, which we attribute to a protracted anti-inflammatory effect, occurring soon after cell transplantation but persisting up to 30 dpt, paralleled by a glial scar-inhibitory effect. Therefore, our data exemplify that neural differentiation is not a pre-requisite for adult NPCs to exert their restorative action thus representing an attractive tool for stroke treatment.

That NPCs integrate into the stroke brain has been shown previously (Bliss et al., 2007; Bacigaluppi et al., 2008). In contrast to the healthy brain, in which intraparenchymally transplanted NPCs migrate only over small distances and in small numbers, the ischaemic brain tissue fosters a selective NPC migration towards the stroke lesion where they eventually differentiate into functional neural cells (Kelly et al., 2004; Buhnemann et al., 2006; Darsalia et al., 2007). Whether robust cell replacement is...
Indeed responsible for the functional improvement induced by NPCs was recently challenged by studies using i.v. transplantation strategies. Early evidence showed a substantial integration and differentiation of i.v.-injected human- or embryonic-neural stem cells into neurons and astrocytes in rodents with experimental intracerebral haemorrhage or focal cerebral ischaemia (Jeong et al., 2003; Jin et al., 2005). Indeed, more recent studies have identified an additional bystander immune-modulatory mechanism, sustained and mediated by i.v.-injected NPC, that is capable of ameliorating clinically-pathological signs of experimental bacterial collagenase VII-induced intracerebral haemorrhage (Lee et al., 2008). This latter mechanism is considered one of the main mechanisms by which systemically injected NPCs exert their therapeutic effects in chronic and acute inflammatory CNS diseases (the concept of ‘therapeutic plasticity’) (Martino and Pluchino, 2006).

In our study, anti-inflammatory and glial scar-inhibitory effects might be common mechanistic pathways through which transplanted NPCs protected the ischaemic brain from delayed injury. The anti-inflammatory mechanism in NPC-transplanted mice is supported by the observed reduction of mRNA levels of pro-inflammatory mediators (i.e. IFN-γ, TNF-α, IL-1β and IL-6) (Barone et al., 1997; Hallenbeck, 2002; Allan et al., 2005) that contribute to the detrimental post-ischaemic brain inflammation in the lesioned areas of transplanted mice and also by the decreased number of activated microglial cells. This latter evidence, recorded soon after cell transplantation, was further reinforced by the reduction of blood-borne CNS-infiltrating inflammatory cells (CD45) at 30 dpt. Further, a glial scar-inhibitory effect in NPC-transplanted mice takes place as suggested by the downregulation of genes promoting astrogliosis (e.g. FGF-II, transforming growth factor-beta (TGF-β), vimentin) in the lesioned area that was accompanied by a statistically significant reduction of the perilesional glial scar thickness at 30 dpt. Although we did observe a slight but non-significant increase of the total length of capillaries, vessel density was not altered after NPC treatment. We favour this as the in vivo proof that NPC treatment did not induce an angiogenic response. The larger volume of spared striatal in NPC-treated animals can, in fact, explain the increase total length values.

Figure 6 Reduction of astroglial scar formation. Quantitative analysis of the GFAP* astroglial scar area (A) and thickness (B) along the ischaemic border zone. White bars refer to sham-treated MCAO mice, while black bars refer to NPC-treated MCAO mice. Data in B and C are expressed as mean values and have been obtained from a total of six regions of interest, as shown by the white boxes in A. GFAP* astroglial scar thickness (red double arrow) in the hemisphere ipsilateral to the stroke in representative sham- (D) and NPC- (E) treated MCAO mice at 30 dpt. The white-dashed line outlines the astroglial scar. The green staining in A, D and E is for GFAP. Scale bars: A, 1 mm, D and E, 20 μm. Data are means ± SE. *P ≤ 0.05, compared with sham-treated mice.
Finally, we cannot exclude that the anti-inflammatory effects exerted in MCAO mice transplanted with NPCs is in part exerted in the body periphery, as recently shown after brain haemorrhage (Lee et al., 2008). However, we found that soon after MCAO (e.g. 3 dpt) a comparable number of blood-borne inflammatory cells infiltrated the CNS, thus indicating that the recruitment of ‘effector’ immune cells from the periphery was not impaired by the cell treatment and, again, suggesting that the protective effect may have been induced within the CNS.

By providing a detailed temporospatial analysis, our data show that focal cerebral ischaemia is followed by delayed neurodegenerative process that continues over >10 dpt, i.e. 13 days after the stroke, resulting in continuous neuronal loss in those parts of the striatum that survived the ischaemic insult, leading to secondary lesion growth. In our study, post-acute NPC treatment provided a delayed structural neuroprotective effect—particularly evident on striatal medium spiny neurons—despite the fact that treatment was initiated far beyond established time-windows.
for neuroprotection. We tend to attribute this late effect to the early anti-inflammatory effect we observed.

A significant downregulation of TNF-α, a known pro-inflammatory cytokine exerting a profound pro-apoptotic effect (Kaushal and Schlichter, 2008), was observed in the striatum of NPC-treated mice. Downregulation of IFN-γ is also worth mentioning. It has been shown that IFN-γ is able to induce apoptosis by increasing expression of death receptor on target cells (Furlan et al., 2001; Hallenbeck, 2002). Considering that neuronal cells do express death receptors such as tumour necrosis factor receptor-1 (TNFR1) (Haase et al., 2008), it is tempting to speculate that the reduction of IFN-γ secretion within the lesioned area might contribute per se to the pro-survival neuronal effect. Further, the importance of both TNF-α and IFN-γ is emphasized by the demonstration that antagonization of the production of either of these inflammatory cytokines significantly prevented secondary infarct growth after stroke (Liesz et al., 2009). The reduction of expression of genes coding for cell death molecules (e.g. caspase-3) in transplanted mice would further support this view.

This immune-modulatory, rather than immune-suppressive action exerted by NPCs may be of pivotal interest as stroke is accompanied by an early peripheral immunodeficiency phase (Meisel et al., 2005; Offner et al., 2006; Offner et al., 2009) that limits the application of systemic immunosuppressive therapies.

Our data exemplify that delayed structural injury indeed exists after ischaemic stroke, and that this injury can be partially reversed by NPC transplantation leading to clinically relevant functional neurological improvements. As these structural changes were observed in a time-window in which remodelling and plasticity processes are very active in the tissue, our data question whether delayed degeneration and remodelling should indeed be regarded as distinct phenomena. Indeed, it has previously been shown that successful remodelling is accompanied by structural changes both in ischaemic tissue and remote from it. As such, remodelling of axonal fibre bundles (Jiang et al., 2006) associated with an increase in the thickness of the CC (Shen et al., 2006) has been reported in stroke animals exhibiting enhanced motor recovery after treatment with bone marrow cells. Yet in this study, the structural changes seen were interpreted as consequence of delayed neuroprotection rather than post-acute plasticity.

In conclusion, our data support the concept that transplanted NPCs, remaining undifferentiated, might promote recovery from ischaemic injury via a multifaceted therapeutic response. This is preferentially exerted through an anti-inflammatory effect that leads to decreased glia scar formation and protects the brain from delayed injury. Based on our here-presented results, cell-based strategies may particularly be attractive for this kind of tissue injury.

Supplementary material

Supplementary material is available at Brain online.
Acknowledgements

We acknowledge the assistance and help of Miriam Ascagni, Cesare Covino, Angela Fendel, Filippo Martinelli Boneschi and Annett Spudich.

Funding

NCCR “Neural plasticity” (to D.M.H.); COST Short Term Scientific Mission (COST-STSM-BM0603-2843); the Swiss National Science Foundation (3200B0-112056/1, to D.M.H.); the BMW Italy Group (to G.M.); Banca Agricola Popolare di Ragusa (to S.P.).

References

Yang Y, Li Q, Shuaib A. Neuroprotection by 2-h postischemia administration of two free radical scavengers, alpha-phenyl-n-tert-butyl-nitrone (PBN) and N-tert-butyl-(2-sulfophenyl)-nitrone (S-PBN), in rats subjected to focal embolic cerebral ischemia. Exp Neurol 2000; 163: 39–45.