Role of lipocalin 2 in intraventricular haemoglobin-induced brain injury

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ABSTRACT

Objective: Our recent studies have shown that blood components, including haemoglobin and iron, contribute to hydrocephalus development and brain injury after intraventricular haemorrhage (IVH). The current study investigated the role of lipocalin 2 (LCN2), a protein involved in iron handling, in the ventricular dilation and neuroinflammation caused by brain injury in a mouse model of IVH.

Design: Female wild-type (WT) C57BL/6 mice and LCN2-deficient (LCN2^{-/-}) mice had an intraventricular injection of haemoglobin, and control mice received an equivalent amount of saline. MRI was performed presurgery and postsurgery to measure ventricular volume and the brains were used for either immunohistochemistry or western blot.

Results: Ventricular dilation was observed in WT mice at 24 h after haemoglobin (25 mg/mL, 20 µL) injection (12.5±2.4 vs 8.6±1.5 mm³ in the control, p<0.01). Western blotting showed that LCN2 was significantly upregulated in the periventricular area (p<0.01). LCN2 was mainly expressed in astrocytes, whereas the LCN2 receptor was detected in astrocytes, microglia/macrophages and neurons. Haemoglobin-induced ventricle dilation and glia activation were less in LCN2^{-/-} mice (p<0.01). Injection of high-dose haemoglobin (50 mg/mL) resulted in lower mortality in LCN2^{-/-} mice (27% vs 86% in WT; p<0.05).

Conclusions: Intraventricular haemoglobin caused LCN2 upregulation and ventricular dilation. Haemoglobin resulted in lower mortality and less ventricular dilation in LCN2^{-/-} mice. These results suggest that LCN2 has a role in haemoglobin-induced brain injury and may be a therapeutic target for IVH.

INTRODUCTION

Acute hydrocephalus is a common complication following haemorrhagic stroke. Almost 50% of intracerebral haemorrhages cause the development of hydrocephalus,¹ and the incidence of acute hydrocephalus within 3 days of subarachnoid haemorrhage ranges from 20% to 63%.^{2–3} It has been reported that the presence of intraventricular haemorrhage (IVH) is the major factor related to hydrocephalus.⁴ Our previous study demonstrated that IVH causes lateral ventricle dilation and that iron plays a role in

IVH-induced brain injury and ventricular enlargement in rats.⁵

Lipocalin 2 (LCN2) is a siderophore-binding protein that has a role in endogenous iron chelation. It has a bacteriostatic effect by competing with bacteria for iron loading.⁶ In the central nervous system, LCN2 is an acute phase protein⁷ and is expressed following spinal traumatic injury,⁸ ischaemic stroke,⁹ intracerebral haemorrhage^{10–11} and neuroinflammation.¹² Diverse functions of LCN2 have been reported. Thus, it has been associated with neuronal cell death,¹³ reactive astrogliosis¹⁴ and activated microglia.¹⁵ However, little is known about the role of LCN2 in IVH-induced hydrocephalus.

The present study addressed the hypothesis that LCN2 may have a potential role in acute ventricular dilation after IVH in mice after haemoglobin injection.

MATERIALS AND METHODS

Animal preparation and haemoglobin injection into the ventricle

Animal-use protocols were approved by the University of Michigan Committee on the Use and Care of Animals. Animals were housed under standard 12:12 light–dark conditions and allowed free water and food. A total of 31 adult female wild-type (WT) C57BL/6 mice (19–24 g; Charles River Laboratories) and 21 adult female LCN2-deficient (LCN2^{-/-}) mice (19–27 g; University of Michigan Breeding Core, gift from Dr Xiaoli Chen, University of Minnesota) were used in this study. Animals were anaesthetised with ketamine (90 mg/kg) and xylazine (5 mg/kg) intraperitoneally and core body temperature was kept at 37.0°C with a controlled heating pad. Mice were positioned in a stereotaxic frame (Kopf Instruments, Tujunga, California, USA). A midline scalp incision was performed to expose the central areas of the skull between the coronal and lambdoid sutures. A cranial burr hole was drilled and a 26-gauge needle inserted stereotaxically into

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the right lateral ventricle (coordinates: 0.5 mm posterior, 1.1 mm lateral to the bregma and 2.6 mm ventral). Either haemoglobin or saline was injected over 5 min using a microinfusion pump (World Precision Instruments, Sarasota, Florida, USA). After injection, the needle was removed, the burr hole filled with bone wax and the skin incision was closed with sutures.

Experimental groups

This study included three parts. First, WT mice received a 20 μ L injection of saline only ($n=6$) or haemoglobin ($n=8$, bovine haemoglobin; 25 mg/mL, ICN Biomedical Inc, Aurora, Ohio, USA) in saline into the right lateral ventricle. MRI was performed 24 h later and the mice were then euthanised for immunohistochemistry and western blot analysis. Second, WT and LCN2^{-/-} mice had an intraventricular injection of haemoglobin (25 mg/mL in 20 μ L saline) or saline only ($n=5$ for each). They underwent MRI presurgery and postsurgery at 24 h. Mice were then euthanised for immunohistochemistry. Third, WT ($n=7$) and LCN2^{-/-} mice ($n=11$) received an intraventricular injection of a high dose of haemoglobin (50 mg/mL, 20 μ L). The mortality rate was determined at 24 h.

MRI for ventricular volume measurement

MRI was performed in a 7.0-T Varian MR scanner (Varian) with acquisition of T2 fast spin-echo sequences, using a field of view of 20×20 mm, matrix of 256×256 mm and 25 coronal slices (0.5 mm thick). Mice were anaesthetised with 1.5% isoflurane throughout the MRI examination. Ventricular volume was measured as described previously.⁵ Briefly, ventricles from the frontal horn of the lateral ventricles to the foramen of Luschka were outlined and measured. Ventricular volume was calculated by totalling the ventricular areas over all slices and multiplying by section thickness. All image analysis was performed using ImageJ software, by an observer blinded to the experimental group. For the comparison between genotypes, ventricular volumes were measured before and after haemoglobin and saline injection, and ventricular dilation was determined as a per cent increase compared with presurgical values.

Immunohistochemistry

Brains were postfixed in 4% formaldehyde and immersed in 30% sucrose in 0.1 M cold phosphate-buffered saline (pH 7.4) for 2–3 days at 4°C. They were then embedded and sliced into 18 μ m-thick coronal sections on a cryostat. Immunohistochemical staining was performed using the avidin–biotin complex technique. Sections were incubated in either 1:10 rabbit or goat serum (Vector Laboratories) for 30 min and then incubated overnight with primary antibody. The antibodies were: polyclonal goat anti-LCN2 (1:200 dilution, B&D System), polyclonal rabbit antiglial fibrillary acidic protein (GFAP; 1:1000 dilution, Millipore) and polyclonal goat anti-Iba-1 (1:400 dilution, Abcam). The

secondary antibodies were rabbit anti-goat IgG (Bio-Rad; 1:1000) or goat anti-rabbit IgG (Bio-Rad; 1:1000). Negative immunohistochemical control procedures included omission of the primary antibody.

Immunofluorescent double labelling

For immunofluorescent double labelling, sections were incubated with 15% normal donkey serum at room temperature for 30 min and then incubated overnight at 4°C with primary antibodies. After washing, sections were incubated with secondary antibodies for 2 h. The primary antibodies were polyclonal goat anti-LCN2 (1:200 dilution, R&D system), polyclonal rabbit anti-SLC22A17 (marker for LCN2 receptor, 1:50 dilution, Abcam), polyclonal mouse anti-GFAP (1:400 dilution, Millipore), monoclonal mouse anti-neuronal nuclei (NeuN; 1:100 dilution, Millipore) and polyclonal goat anti-Iba-1 (1:500 dilution, Abcam) or rabbit anti-Iba-1 (1:500 dilution, Wako). The secondary antibodies were donkey anti-goat IgG(H+L) Alexa Fluor 594 (1:500 dilution, Invitrogen), donkey anti-rabbit IgG(H+L) Alexa Fluor 594 (1:500 dilution, Invitrogen), donkey anti-mouse IgG(H+L) Alexa Fluor 488 (1:500 dilution, Invitrogen), donkey anti-rabbit IgG(H+L) Alexa Fluor 488 (1:500 dilution, Invitrogen) and donkey anti-goat IgG(H+L) Alexa Fluor 488 (1:500 dilution, Invitrogen). The double labelling was analysed using a fluorescence microscope.

Western blot analysis

A 3 mm thick coronal brain slice was cut 3 mm from the frontal pole. The ipsilateral cortex and periventricular area including white matter were sampled. Samples were sonicated in western blot sample buffer. Protein concentrations were determined by Bio-Rad protein assay kit and 10 μ g protein samples separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis, and transferred to a Hybond-C pure nitrocellulose membrane (Amersham). Membranes were probed with primary antibody: polyclonal goat anti-LCN2 (1:200 dilution, R&D system). The secondary antibody was rabbit anti-goat IgG (1:1000 dilution, Bio-Rad). Image analysis was performed using ImageJ software by a blinded observer.

Cell counting

The numbers of GFAP and Iba-1 immunopositive cells were measured on high-power images (×40 magnification) taken in the periventricular area by a digital camera, using sections 0.5 mm anterior from the bregma. The numbers of activated astrocytes and microglia were counted by a blinded observer and statistically analysed as previously described,^{9 16} using the National Institutes of Health (NIH) imageJ programme. Three regions of interest were examined per section and all measurements counted on three sections, and the mean values were used.

Statistical analysis

Values are presented as the means $\pm$ SD. Statistical differences among groups were analysed using Student's t test or one-way analysis of variance with a Tukey's all pair's comparison. Statistical significance was set at $p<0.05$.

RESULTS

Development of ventricular dilation

No mice died after saline or 25 mg/mL haemoglobin injection. Ventricular volumes 24 h after haemoglobin injection (12.5 ± 2.4 mm³; $n=8$) were significantly larger than those in mice receiving saline injection (8.6 ± 1.5 mm³; $n=6$; $p<0.01$, figure 1).

Expression of LCN2 after haemoglobin injection

Using immunohistochemistry, we found that periventricular and cortical LCN2 immunoreactivity was very weak in the saline injection group (figure 2A). In contrast, there was marked LCN2 expression of LCN2 in both areas 24 h after haemoglobin injection (figure 2B). By western blot, haemoglobin increased LCN2 protein

levels ~20-fold compared with saline injection at 24 h in both the periventricular area ($26\,362\pm 4165$ vs 2780 ± 579 pixels in the control, $n=3-5$, $p<0.01$, figure 2C) and cortex ($12\,352\pm 2812$ vs 687 ± 215 pixels in the control, $n=3-5$, $p<0.01$, figure 2D).

LCN2 was mainly expressed in astrocytes

Double-labelling immunofluorescence was used to determine which cell types expressed LCN2 after haemoglobin injection. LCN2 immunoreactivity predominantly co-localised with GFAP-positive cells in the periventricular area. LCN2 immunoreactivity did not co-localise with Iba-1-positive cells and only a few LCN2-positive cells in the cortex were also NeuN positive (figure 3).

LCN2 receptor was expressed in astrocytes, microglia and neurons

Double-labelling immunofluorescence was also used to determine which cell types expressed the LCN2 receptor (SLC22A17) after haemoglobin injection. Co-expression of SLC22A17 immunoreactivity with GFAP-positive cells was observed in the periventricular area. Co-expression of SLC22A17 immunoreactivity and Iba-1-positive cells was also detected in the periventricular area and ependymal layer. Co-expression of SLC22A17 and NeuN immunoreactivity was observed in the cortex (figure 4).

LCN2 and haemoglobin-induced ventricular dilation

WT and LCN2^{-/-} mice each received an intraventricular injection of haemoglobin (25 mg/mL) or saline only ($n=5$ for each). In the saline injection group, the increase in ventricular volume compared with preinjection MRI was not significantly different between WT and LCN2^{-/-} mice ($10\pm 5\%$ vs $7\pm 4\%$, respectively, $n=5$ for each, $p>0.05$). However, the increase in ventricular volume induced by haemoglobin (25 mg/mL) injection was significantly lower in LCN2^{-/-} mice ($7\pm 8\%$ vs $45\pm 19\%$ in saline controls, $p<0.01$, figure 5).

Glial and microglial activation

In the saline injection groups, the number of GFAP and Iba-1-positive cells in periventricular area was not significantly different between WT and LCN2^{-/-} mice ($p>0.05$; figure 6). At 24 h after haemoglobin injection, the numbers of both GFAP-positive cells and Iba-1-positive cells were significantly increased in the periventricular area of WT mice (figure 6). Those increases were significantly lower in LCN2^{-/-} mice ($p<0.01$, figure 6).

Mortality after high-dose haemoglobin injection

Intraventricular injection of a higher dose of haemoglobin (50 mg/mL, 20 μ L) resulted in high mortality in WT mice by 24 h (86%, 6/7). In contrast, there was much lower mortality in LCN2^{-/-} mice (27%, 3/11; $p<0.05$).

Figure 1 Intraventricular injection of haemoglobin caused ventricular dilation. T2-weighted MRI and frozen coronal brain sections (A) and the measurement of ventricular volume with T2-weighted MRI (B) from mice 24 h after injection of 20 μ L of saline or haemoglobin (25 mg/mL) into the right lateral ventricle. The data points represent means $\pm$ SD, $n=6-8$, ** $p<0.01$ versus saline group.

Figure 2 Lipocalin 2 (LCN2) protein was markedly upregulated in the brain 24 h after intraventricular haemoglobin injection compared with saline controls. Immunohistochemistry staining demonstrated that LCN2 immunoreactivity was expressed in the periventricular area and cortex 24 h after intraventricular haemoglobin injection (A) compared with saline controls (B). Boxes show areas examined at higher magnification in the higher power micrographs on the right (scale bars=100 and 20 μ m). LCN2 protein levels were upregulated in the periventricular area (C) and cortex (D) 24 h after intraventricular haemoglobin injection, as shown by western blot. The data points represent means $\pm$ SD, $n=3-5$, ** $p<0.01$ versus saline group.

DISCUSSION

The present study has several major findings: (1) intraventricular haemoglobin injection caused ventricular dilation and neuroinflammation in the periventricular area; (2) LCN2 expression was markedly increased in the periventricular area and cortex after haemoglobin injection; (3) LCN2^{-/-} mice had less haemoglobin-induced ventricular dilation with less glial activation and (4) in a high-dose haemoglobin injection model, there was lower mortality rate in LCN2^{-/-} mice compared with WT controls.

Posthaemorrhagic hydrocephalus is traditionally thought to involve: (1) arachnoid obstruction by blood components; (2) subarachnoid fibrosis and (3) ventricular wall damage. Consequently, impaired cerebrospinal fluid (CSF) flow, excess CSF production and lack of CSF reabsorption may result in the development of hydrocephalus.¹⁷ Normal ependymal cells are associated with the homeostasis of CSF in the brain and ependymal dysfunction may result in the development of

hydrocephalus.¹⁷ Other reports have indicated that development of ventriculomegaly is associated with a reaction of periventricular astrocytes.^{18, 19} However, the precise mechanisms of IVH-induced hydrocephalus are still not fully understood. Our previous studies have shown that intraventricular injection of autologous arterial blood,⁵ lysed red blood cells²⁰ and haemoglobin²¹ result in marked ventricular enlargement in rats, and that iron accumulation is associated with IVH-induced hydrocephalus.⁵ In addition, iron is involved in LCN2 regulation with deferoxamine, an iron chelator, reducing LCN2 upregulation in a rat model of intracerebral haemorrhage.²¹ In the present study, we demonstrated that intraventricular haemoglobin injection induced acute ventricular dilation in the mouse, and that LCN2 is also upregulated in the periventricular area and cortex. LCN2 expression can be strongly enhanced by kainate,¹² lipopolysaccharide²² and iron.¹⁰ Haemoglobin consists of haem and globin. Haem is degraded into carbon monoxide, biliverdin and iron in the brain,

Figure 3 Immunoreactivity for lipocalin 2 (LCN2), glial fibrillary acidic protein (GFAP), Iba-1 and neuronal nuclei (NeuN) 24 h after intraventricular haemoglobin injection. Immunofluorescent double labelling showed that LCN2-positive cells were mainly astrocytes. Scale bar=50 μ m.

Figure 4 Immunoreactivity for the lipocalin 2 (LCN2) receptor (SLC22A17), glial fibrillary acidic protein (GFAP), Iba-1 and neuronal nuclei (NeuN) 24 h after intraventricular haemoglobin injection. Co-localisation of the LCN2 receptor with GFAP, Iba-1 and NeuN by double labelling is shown. Scale bar=50 μ m.

Figure 5 Ventricular volumes were measured before and after (24 h) the injection of haemoglobin (25 mg/mL) or saline into wild-type (WT) and in lipocalin 2-deficient (LCN2^{-/-}) mice. Ventricular dilation was calculated as a per cent of presurgery values. Haemoglobin caused marked ventricular dilation in WT mice compared with saline controls, but not LCN2^{-/-} mice. The data represent means±SD, n=5 for each, **p<0.01 versus WT saline group, ##p<0.01 versus LCN2^{-/-} haemoglobin group.

resulting in iron overload in the brain after haemorrhagic stroke. A previous study has suggested that LCN2 can bind to iron and deliver it to the cells through a process of endocytosis.²³ Increasing evidence indicates that iron released from haem may contribute to acute hydrocephalus.^{21 24} These results suggest that LCN2

induced by haemoglobin may take part in the development of ventricular dilation by participating in iron overload.

We assessed which cell types express LCN2 and the LCN2 receptor after haemoglobin injection by immunofluorescent double labelling. We found that LCN2 was mainly expressed in astrocytes and that the LCN2 receptor was expressed in astrocytes, microglia and neurons. We hypothesise that after an IVH, the release of haemoglobin induces reactive astrocytes to secrete LCN2, which then influences neuronal and glia function via the LCN2 receptor.

The role of LCN2 in the central nervous system has not been well studied. The present study revealed that LCN2^{-/-} mice have less ventricle dilation, and lower numbers of activated astrocytes and amoeboid microglia after haemoglobin injection. Previous studies have found that LCN2 causes increased GFAP expression¹⁴ and promotes astrocyte and microglia activation.²⁵ Another recent study indicated that LCN2 deficiency attenuates neuroinflammation by reducing glial and microglial activation in a systemic inflammation model.²⁶

The present study found the intraventricular injection of a high dose of haemoglobin caused less mortality in LCN2^{-/-} mice compared with WT mice. A previous study also demonstrated a lower mortality rate in LCN2^{-/-} mice than in WT mice in a brain ischaemia/reperfusion model.⁹ In that study, LCN2^{-/-} mice also had reduced glial activation. The results suggest that LCN2 secreted by reactive astrocytes may act to promote neuroinflammation, resulting in the development of brain injury, acute ventricular dilation and mortality. Further studies should determine the pathways by which LCN2 taking participates in brain injury following IVH.

Figure 6 In wild-type (WT) mice, intraventricular injection of haemoglobin (25 mg/mL) resulted in periventricular activation of astrocytes (glial fibrillary acidic protein (GFAP) staining) and microglia (amoeboid Iba-1 staining) at 24 h compared with saline-injected controls (A). The activation by haemoglobin was attenuated in lipocalin 2-deficient (LCN2^{-/-}) mice (A). The numbers of periventricular cells expressing GFAP (B) and Iba-1 (C) were counted in the WT and LCN2^{-/-} mice. The data points represent the mean±SD, scale bar=50 µm, n=5 per group; **p<0.01 versus WT saline group, ##p<0.01 versus LCN2^{-/-} haemoglobin group.

CONCLUSION

The present study demonstrates that intraventricular haemoglobin injection causes marked periventricular upregulation of LCN2 and acute ventricular dilation, and can cause death in mice. The adverse effects of haemoglobin were ameliorated in LCN2^{-/-} mice, suggesting that LCN2 may have a role in the development of hydrocephalus and brain injury after IVH.

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