
DR YA HUA (Orcid ID : 0000-0003-0100-2289)

DR GUOHUA XI (Orcid ID : 0000-0003-2846-5937)

Article type : Special Issue Article

Activation of epiplexus macrophages in hydrocephalus caused by subarachnoid hemorrhage and thrombin

Yingfeng Wan^{1,2}, Ya Hua¹, Hugh J. L. Garton¹, Nemanja Novakovic¹,
Richard F. Keep¹, and Guohua Xi¹

¹Department of Neurosurgery, University of Michigan, Ann Arbor, Michigan, USA

²Department of Neurosurgery, Sir Run Run Shaw Hospital, Zhejiang University, Hangzhou,
China

Correspondence and Reprint Requests:

Guohua Xi, M.D.

R5018 Biomedical Science Research Building

University of Michigan

109 Zina Pitcher Place, Ann Arbor, Michigan 48109-2200

Telephone: (734) 764-1207

Fax: (734) 764-7322

E-mail: guohuaxi@umich.edu

This is the author manuscript accepted for publication and has undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the [Version of Record](#). Please cite this article as [doi: 10.1111/CNS.13203](https://doi.org/10.1111/CNS.13203)

This article is protected by copyright. All rights reserved

Running Head: Epiplexus macrophages and hydrocephalus in SAH

Summary

Aims: We have found that hydrocephalus development in spontaneously hypertensive rats was associated with activation of epiplexus cells. The current study examined whether epiplexus cell activation occurs in a rat subarachnoid hemorrhage (SAH), whether activation would be greater in a subset of rats that developed hydrocephalus and the potential role of thrombin in epiplexus cell activation.

Methods: There were two parts to this study. First, an endovascular perforation was performed in rats to induce SAH. Second, rats received an intraventricular infusion of either thrombin or saline. Magnetic resonance imaging was used to measure the ventricular volumes.

Immunofluorescence and immunohistochemistry were used to study epiplexus cell activation.

Results: Iba-1, OX-6, and CD68 were expressed in the epiplexus cells of the choroid plexus in sham-operated rats. SAH increased Iba-1 and CD68 immunoreactivity in epiplexus cells in addition to an increase in Iba-1 positive cell soma size. Those effects were greater in rats that developed hydrocephalus. Intraventricular thrombin mimicked the effects of SAH on epiplexus cell activation and hydrocephalus.

Conclusion: This study supports the concept that epiplexus cell activation is associated with hydrocephalus development. Epiplexus cell activation may be in response to thrombin production after hemorrhage and it may be a therapeutic target.

Keywords: epiplexus cells, hydrocephalus, subarachnoid hemorrhage, thrombin

1. Introduction

There is a well-established association between an intraventricular hemorrhage (IVH) and hydrocephalus development. However, the precise mechanisms remain unclear. Understanding the mechanisms may lead to changes in how we manage patients with hydrocephalus, especially because we currently experience poor clinical outcomes in patients that develop hydrocephalus after IVH.¹ Additionally, it is known that hydrocephalus and IVH are both independent predictors of poor outcome in spontaneous ICH.² Patients with concurrent IVH and ICH

development had a favorable outcome of 15% compared to 31% with ICH alone. Hydrocephalus, which develops in 55% of patients with IVH, further decreased the chance of a favorable outcome to 11.5%.³ Another independent prognostic factor for unfavorable outcomes is IVH in patients with subarachnoid hemorrhage (SAH). 20-30% of such patients develop acute hydrocephalus.⁴ The increased intracranial pressure associated with hydrocephalus correlates with poorer prognosis in patients with aneurysmal SAH.^{5,6}

Hydrocephalus development after IVH is not only a major complication in adult patients with ICH and SAH, but also in premature infants. More than 12,000 premature infants in the United States develop hydrocephalus after IVH every year. IVH is an independent poor prognostic predictor for neonates after a germinal matrix hemorrhage.⁷

The choroid plexus is a hyper-vascularized tissue in the brain ventricle system which predominantly functions to secrete CSF. Epiplexus cells, also named “Kolmer cells,” were initially detailed in 1921 by Kolmer. These cells have primarily been examined using immunohistochemistry and electron microscopy.⁸ They exhibit diverse morphologies including round, polar, and stellate, are macrophages of monocytic origin in the brain ventricle system considered to have an immunologic role due to their involvement with antigen presentation, nitric oxide (NO) production, phagocytosis of various foreign bodies, and iron accumulation.⁹

It is generally accepted that inflammation has a role in hydrocephalus development.¹⁰ Recent evidence indicates that inflammatory signaling at the choroid plexus is involved in CSF hypersecretion and hydrocephalus development after cerebral hemorrhage.¹¹ We recently found that epiplexus cell activation at the choroid plexus occurs during hydrocephalus development that naturally occurs in spontaneous hypertensive rats.¹² Therefore, the current study was undertaken to examine whether epiplexus cell activation occurs in a rat model of SAH, whether the degree of activation would be greater in those rats that develop hydrocephalus after SAH, and whether the effects of SAH could be mimicked by intraventricular thrombin. Thrombin is produced during clot formation, it is known to activate microglia/macrophages¹³ and we have recently shown it can induce hydrocephalus in rats¹⁴.

2 Materials and Methods

2.1 Animal Preparation

All animal protocols were approved by the University of Michigan Committee on the Use and Care of Animals. Standard 12:12 light-dark conditions were used to house the rats and they were allowed free access to water and food. The experiments used a total of 70 male adult Sprague-Dawley rats (2-3 month old, Charles River Laboratories, Portage, MI).

2.2 Experiment Groups

The study had two parts. In the first part, rats were randomly assigned into two groups: a sham operation (n=12) or SAH induction (n=25). Eight rats died within 24 hours after SAH induction. Surviving rats were sacrificed at 24 hours after the operations. The SAH group was further divided into two groups based on MRI scanning for ventricular volume: SAH with hydrocephalus (SAH with HC, n=15), and SAH without hydrocephalus (SAH w/o HC, n=10). In the second part, 25 rats were injected intracerebroventricularly with 50ul (3U) of rat thrombin (Sigma-Aldrich, St Louis, MO, USA) in saline (n=11) or 50 µl saline alone (n=14) over 7 minutes. They were euthanized at 24 hours after receiving an MRI scan. All brains were used for immunohistochemistry and immunofluorescence staining.

2.3 Subarachnoid Hemorrhage Model

Endovascular perforation was used to induce SAH as previously described.^{15,16} Rats were anesthetized using 5% isoflurane (VetOne Fluriso; MWI, Boise, ID), intubated and placed on mechanical ventilation. Isoflurane was then titrated between 2.5 and 3%. Core body temperature was maintained at 37.5 °C using a heating pad with a feedback-control mechanism. While in the supine position, the left external carotid artery was transected distally and reflected caudally in reference to the internal carotid artery. A 3-0-nylon monofilament suture was inserted into the stump of external carotid artery and advanced distally into the intracranial internal carotid artery until resistance was felt. At that point, the artery at the bifurcation of intracranial internal carotid was perforated to induce SAH. The suture was then removed into the external carotid artery.

2.4 Intraventricular Injection

Rats received were anesthetized with pentobarbital (50 mg/kg; IP) and the core body temperature kept at 37.5 °C using a heating pad. Rats were then positioned in a stereotaxic frame (Kopf Instruments, Tujunga, CA, USA), a 1 mm cranial burr hole drilled, and a 26-gauge needle stereotaxically inserted into the right lateral ventricle using the following coordinates: 4.5 mm ventral, 0.6 mm posterior, and 1.6 mm lateral to the bregma. Rats received either saline or thrombin injected using a microinfusion pump (World Precision Instruments, Sarasota, FL,

USA). The needle was then removed, bone wax was used to fill the burr hole and the skin incision sutured closed.

2.5 Magnetic Resonance Imaging (MRI) and Ventricle Volume Measurement

Rats were anesthetized with 2% isoflurane throughout an MRI using a 7.0-T Varian MR scanner (Varian, Palo Alto, CA, USA) with acquisition of T2 fast spin-echo and T2* gradient-echo sequences using 25 coronal slices (0.5 mm thick) with a field of view of 20x20 mm and matrix of 256x256 mm. Ventricular volume was measured from the frontal horns of the lateral ventricles to the foramen of Luschka using the sum of the ventricular area over all slices and multiplying by section thickness. Image analysis was performed using Image J by a blinded observer. Hydrocephalus was defined as a ventricular volume over +3 standard deviations (SD) from the mean of the sham animals.

2.6 Immunohistochemistry and Immunofluorescence Staining

Immunohistochemistry was performed as described previously.¹⁷ In brief, rats were anesthetized using pentobarbital (100mg/Kg; IP) and then underwent transcardiac perfusion with 4% paraformaldehyde in 0.1 mol/L phosphate-buffered saline (pH 7.4). Brains were then harvested and kept in 4% paraformaldehyde for 24 hours. Next, they were immersed in 30% sucrose for 3 to 4 days at 4°C prior to embedding in a mixture of 30% sucrose and optimal cutting temperature compound (Sakura Finetek Inc, Torrance, CA; 1:2 ratio) and sectioning on a cryostat (18µm). Immunohistochemistry was performed using the avidin-biotin complex technique. The primary antibodies were polyclonal goat anti-Iba-1 (ionized calcium-binding adapter molecule 1, 1:400 dilution; Abcam), monoclonal mouse anti-MHC II RT 1B (**clone number:OX-6**, 1:200 dilution; AbD Serotec), and monoclonal mouse anti-CD68 (**clone number:ED1**, 1:100 dilution; Abcam).

Double labeling immunofluorescence was performed on paraffin-embedded sections. Primary antibodies were added and incubated overnight at 4°C. Sections were then rinsed in PBS prior to incubation at room temperature for 2 hours with the secondary antibodies. The primary antibodies were polyclonal goat anti-Iba-1 (ionized calcium-binding adapter molecule 1, 1:400 dilution; Abcam), monoclonal mouse anti-MHC II RT 1B (OX-6, 1:200 dilution; AbD Serotec), and monoclonal mouse anti-CD68 (ED1, 1:100 dilution; Abcam). Secondary antibodies were Alexa Fluor 594 donkey anti-mouse IgG (1:500, Invitrogen) and Alexa Fluor 488 donkey anti-goat IgG (1:500, Invitrogen). Fluoroshield™ with DAPI (F6057) was used for nuclear labeling.

2.7 Cell Counting and Soma Size Measurement

Immuno-positive cells were counted on high-power images (X40 magnification) from a digital camera. Both lateral ventricle choroid plexuses were counted. The number of epipterus cells was calculated as a percentage of total choroid plexus epithelial cell number. Soma size measurements were determined on high-power images (X100 magnification). Ten immuno-positive cells from one sample were randomly chosen to have their soma sizes measured. Cell counts and soma size measurements were performed using Image J. Mean values are reported from measurements that were repeated three times by a blinded observer.

2.8 Statistical Analysis

Data are expressed as mean \pm standard deviation (SD), and comparisons between groups were performed using Student *t* test or one-way ANOVA with a Tukey's post-hoc test for multiple comparisons. $p < 0.05$ was considered to be statistically significant.

3 Results

3.1 Iba-1, OX-6 and CD68 are expressed in choroid plexus epipterus cells

Epipterus cells are macrophage-like cells that reside on the apical surface of the choroid plexuses. Iba-1, OX-6 and CD68 (three macrophage/microglia markers) immunoreactivity was noted in epipterus cells at 24 hours in sham rat group (Fig. 1A). These cells displayed diverse morphologies ranging from round to stellate. Immunofluorescence double staining revealed that almost all CD68-positive epipterus cells also stained for Iba-1. In contrast, most of OX-6 positive epipterus cells did not stain for Iba-1 positive (Fig. 1B).

3.2 Iba-1 positive cells increased and enlarged after SAH

Examples of MRIs of rats with and without (w/o) hydrocephalus (HC) after SAH are shown in Fig. 2A. In the rats w/o HC, there was an increase in the number of Iba-1 positive cells at 24 hours in SAH ($9.3 \pm 0.6\%$; $n=10$) compared to sham-operated rats ($6.5 \pm 0.4\%$; $n=12$, $p < 0.05$, Fig. 2B). This was further magnified in the SAH with HC group ($13.2 \pm 0.9\%$; $n=15$; $p < 0.01$ vs. w/o HC group; Fig. 2B). Additionally, macrophage soma size in the SAH w/o HC group ($17.4 \pm 0.7 \mu\text{m}^2$; $n=10$) was significantly larger in the sham group ($14.4 \pm 0.5 \mu\text{m}^2$; $n=12$, $p < 0.05$, Fig. 2C). Again, this effect was larger in SAH with HC group ($20.5 \pm 0.8 \mu\text{m}^2$; $n=15$, $p < 0.01$ vs. w/o HC group; Fig. 2C).

3.3 CD68 expression was increased after SAH

The number of CD68 positive cells was also significantly increased at 24 hours in SAH w/o HC group ($3.4 \pm 0.3\%$, $n=8$) compared with sham-operated rats ($0.7 \pm 0.1\%$; $n=7$, $p < 0.01$; Fig. 3). Again, this effect was magnified in the SAH with HC group ($6.0 \pm 0.3\%$, $n=9$, $p < 0.01$ vs. w/o HC; Fig. 3).

3.4 Iba-1 positive cells were upregulated and activated after intraventricular thrombin injection

Intraventricular injection of thrombin induced hydrocephalus in rats as identified on MRI at 24 hours (Fig. 4A). It also increased the number of Iba-1 positive epileptus cells ($13.2 \pm 0.8\%$, $n=14$ vs. $7.9 \pm 0.7\%$ in saline injected group, $n=11$, $p < 0.01$, Fig. 4B). In addition, the size of those cells was bigger ($17.8 \pm 0.5 \mu\text{m}^2$; $n=14$) than in saline injected rats ($12.6 \pm 0.4 \mu\text{m}^2$; $n=11$, $p < 0.01$, Fig. 4C).

3.5 CD68 positive cells increased after intraventricular thrombin injection

Intraventricular injection of thrombin also increased the number of CD68 positive epileptus cells at 24 hours ($5.1 \pm 0.4\%$; $n=13$ vs. $1.1 \pm 0.2\%$ in the saline injected group, $n=11$; $p < 0.01$, Fig. 5A). In addition, the percentage of Iba-1 positive cells that were also CD68 positive was increased after thrombin injection ($45 \pm 3\%$; $n=6$) compared to saline controls ($15 \pm 2\%$, $n=6$; $p < 0.01$, Fig. 5B).

4 Discussion

The main findings of this study are: (1) Iba-1, CD68 and OX-6 are expressed in epileptus cells in sham-operated rats; (2) induction of SAH increased the numbers of CD68 and Iba-1 positive cells, as well as Iba-1 positive cell soma size at 24 hours in rats; (3) Those effects were greater in rats that developed hydrocephalus after SAH; (4) Intraventricular injection of thrombin in rats also increased the number of Iba-1 positive and CD68 positive cells at 24 hours.

The epileptus, free-floating, and supraependymal cells are all considered intraventricular macrophages. Epileptus cells, also named "Kolmer cells," were initially detailed in 1921 by Kolmer, residing on the apical surface (CSF-facing) of the choroid plexus. These cells have been

examined by immunohistochemistry and electron microscopy.⁸ There have been several theories regarding the origin of epiplexus cells. However, the most accepted theory is that circulating monocytes become tissue macrophages after reaching connective tissue spaces by migrating from the thin-walled choroid blood vessels. These macrophages become epiplexus cells after they pass through the choroid plexus epithelium.¹⁸ These cells are in close proximity to the surface of the choroid plexus epithelium and show different morphologies which range from round to polar and stellate.¹⁹ Epiplexus cells are mainly associated with antigen presentation, nitric oxide (NO) production, phagocytosis of various foreign bodies, and iron accumulation, and are thus considered to have an immunological role in the brain ventricles.⁹ A previous study found that epiplexus cells express OX-42, which recognizes complement type 3 receptors (CR3), and OX-18, which recognizes major histocompatibility complex (MHC) antigens in rats of different ages.²⁰ In our study, immunohistochemistry and immunofluorescence staining indicated that Iba-1, OX-6 and CD68 were expressed in epiplexus cells of sham-operated rats. OX-6 recognizes major histocompatibility complex II (MHC II) antigens as an activated macrophage marker in rat brains, while Iba-1 is a macrophage marker in many sites in brain, including the choroid plexus in rat.^{21,22} An interesting phenomenon was that most OX-6 positive epiplexus cells were different from Iba-1 positive epiplexus cells, with only a small portion co-localizing on immunofluorescence staining. The role of these subsets of epiplexus cells requires further investigation.

Subarachnoid hemorrhage resulted in an increase in the number of Iba-1 positive and CD68 positive epiplexus cells on the choroid plexus, as well as an increase in soma size in Iba-1 positive cells. This suggests that SAH results in epiplexus cell activation. One potential factor that might participate in such activation is thrombin that is generated during clot formation. Thrombin can activate microglia/macrophages¹³ and we found that intraventricular injection of thrombin mimicked the effects of SAH, increasing the number of Iba-1 positive and CD68 positive epiplexus cells, and increasing soma size. This suggests that thrombin may play an important role in epiplexus cell activation after SAH.

We previously found that hydrocephalus occurs in about 44% of rats with SAH induced by endovascular perforation¹⁶ and that thrombin directly contributes to hydrocephalus development after intraventricular hemorrhage.¹⁴ Several research studies were interested in the macrophages of the cerebrospinal fluid, cerebral cortex, and the basal ganglia after SAH.²³⁻²⁵ Another study

revealed that intraventricular macrophages were activated by inducing hydrocephalus in prenatal rats by injecting pregnant rats with 6-aminonicotinamide.²⁶ Mature microglia/macrophage have highly ramified processes, adopting an “amoeboid” morphology with larger, rounder cell bodies during activation.²⁷ Interestingly, our data indicates that compared to the SAH without hydrocephalus group, the SAH with hydrocephalus rats have increased epiplexus cell numbers and size. The effect of epiplexus cell activation requires further investigation. These cells become remarkably larger with ruffles, and may participate in phagocytosis of erythrocytes in SAH. The cells stained with Iba-1 and OX-6 antibodies had enhanced immunoreactivity, suggesting immunoregulatory engagement due to upregulation of phagocytic activity. Their location on the apical surface of the choroid plexus epithelium, also suggests they may modify choroid plexus, the site of the blood-CSF barrier, function.

CD68 is a classic M1-like (proinflammatory) phenotype of microglia/macrophage activated in response to cerebral hemorrhage and other neurological conditions.^{28,29} In our study, CD68 expression was very low in both the sham and intraventricular injection of saline groups. However, it significantly increased after SAH and intraventricular injection of thrombin in rats. Iba-1 and CD68 are two widely used markers to identify microglia/macrophage in human and animal brains.^{30,31} In the present study, the percentage of Iba-1 positive epiplexus cells that were CD68 positive was significantly increased by thrombin. This phenomenon suggests that M1 polarization becomes a more predominant response, thus causing more inflammatory changes, eventually leading to hydrocephalus development.

In summary, this study reveals that SAH induces choroid plexus epiplexus cell activation, an effect mimicked by intraventricular thrombin. There was a positive correlation between such activation and hydrocephalus development. Whether epiplexus cell activation participates in, or is a result of, the hydrocephalus development needs further investigation.

Acknowledgements:

GX, YH and RFK were supported by grants NS-073595, NS-090925, NS-096917, NS-106746 and NS-112394 from the National Institutes of Health.

Conflict of interest statement: We declare that we have no conflict of interest.

References

1. Strahle J, Garton HJL, Maher CO, Muraszko KM, Keep RF, Xi GH. Mechanisms of Hydrocephalus After Neonatal and Adult Intraventricular Hemorrhage. *Transl Stroke Res.* 2012;3:S25-S38.
2. Bhattathiri PS, Gregson B, Prasad KS, Mendelow AD, Investigators S. Intraventricular hemorrhage and hydrocephalus after spontaneous intracerebral hemorrhage: results from the STICH trial. *Acta Neurochir Suppl.* 2006;96:65-68.
3. Hanley DF. Intraventricular hemorrhage: severity factor and treatment target in spontaneous intracerebral hemorrhage. *Stroke.* 2009;40(4):1533-1538.
4. Rosen DS, Macdonald RL, Huo D, et al. Intraventricular hemorrhage from ruptured aneurysm: clinical characteristics, complications, and outcomes in a large, prospective, multicenter study population. *J Neurosurg.* 2007;107(2):261-265.
5. Hasan D, Vermeulen M, Wijdicks EF, Hijdra A, van Gijn J. Management problems in acute hydrocephalus after subarachnoid hemorrhage. *Stroke.* 1989;20(6):747-753.
6. van Asch CJ, van der Schaaf IC, Rinkel GJ. Acute hydrocephalus and cerebral perfusion after aneurysmal subarachnoid hemorrhage. *AJNR Am J Neuroradiol.* 2010;31(1):67-70.
7. Strahle JM, Garton T, Bazzi AA, et al. Role of hemoglobin and iron in hydrocephalus after neonatal intraventricular hemorrhage. *Neurosurgery.* 2014;75(6):696-705; discussion 706.
8. Ling EA, Kaur C, Lu J. Origin, nature, and some functional considerations of intraventricular macrophages, with special reference to the ependymal cells. *Microsc Res Tech.* 1998;41(1):43-56.
9. Maslieieva V, Thompson RJ. A critical role for pannexin-1 in activation of innate immune cells of the choroid plexus. *Channels (Austin).* 2014;8(2):131-141.
10. Chen S, Luo J, Reis C, Manaenko A, Zhang J. Hydrocephalus after Subarachnoid Hemorrhage: Pathophysiology, Diagnosis, and Treatment. *Biomed Res Int.* 2017;2017:8584753.
11. Karimy JK, Zhang J, Kurland DB, et al. Inflammation-dependent cerebrospinal fluid hypersecretion by the choroid plexus epithelium in posthemorrhagic hydrocephalus. *Nat Med.* 2017;23(8):997-1003.

-
12. Gu C, Hao X, Li J, Hua Y, Keep RF, Xi G. Effects of minocycline on epiplexus macrophage activation, choroid plexus injury and hydrocephalus development in spontaneous hypertensive rats. *J Cereb Blood Flow Metab.* 2019, in press.
 13. Wan S, Cheng Y, Jin H, et al. Microglia Activation and Polarization After Intracerebral Hemorrhage in Mice: the Role of Protease-Activated Receptor-1. *Transl Stroke Res.* 2016;7:478-487.
 14. Gao F, Liu F, Chen Z, Hua Y, Keep RF, Xi G. Hydrocephalus after intraventricular hemorrhage: the role of thrombin. *J Cereb Blood Flow Metab.* 2014;34(3):489-494.
 15. Lee JY, Keep RF, He Y, Sagher O, Hua Y, Xi G. Hemoglobin and iron handling in brain after subarachnoid hemorrhage and the effect of deferoxamine on early brain injury. *J Cereb Blood Flow Metab.* 2010;30(11):1793-1803.
 16. Okubo S, Strahle J, Keep RF, Hua Y, Xi G. Subarachnoid hemorrhage-induced hydrocephalus in rats. *Stroke.* 2013;44(2):547-550.
 17. Liu R, Cao S, Hua Y, Keep RF, Huang Y, Xi G. CD163 Expression in Neurons After Experimental Intracerebral Hemorrhage. *Stroke.* 2017;48(5):1369-1375.
 18. Carpenter SJ, McCarthy LE, Borison HL. Electron microscopic study of the epiplexus (Kolmer) cells of the cat choroid plexus. *Z Zellforsch Mikrosk Anat.* 1970;110(4):471-486.
 19. Joukal M, Klusakova I, Solar P, Kuklova A, Dubovy P. Cellular reactions of the choroid plexus induced by peripheral nerve injury. *Neurosci Lett.* 2016;628:73-77.
 20. Lu J, Kaur C, Ling EA. Immunophenotypic features of epiplexus cells and their response to interferon gamma injected intraperitoneally in postnatal rats. *J Anat.* 1994;185 (Pt 1):75-84.
 21. Li QY, Barres BA. Microglia and macrophages in brain homeostasis and disease. *Nat Rev Immunol.* 2018;18(4):225-242.
 22. Kirik OV, Sukhorukova EG, Korzhevskii DE. [Calcium-binding Iba-1/AIF-1 protein in rat brain cells]. *Morfologiya.* 2010;137(2):5-8.
 23. Zhang H, Okubo S, Hua Y, Keep RF, Xi G. Basal Ganglia Damage in Experimental Subarachnoid Hemorrhage. *Acta Neurochir Suppl.* 2016;121:141-144.

-
24. Jing C, Zhang H, Shishido H, Keep RF, Hua Y. Association of Brain CD163 Expression and Brain Injury/Hydrocephalus Development in a Rat Model of Subarachnoid Hemorrhage. *Front Neurosci.* 2018;12:313.
 25. Thomas AJ, Ogilvy CS, Griessenauer CJ, Hanafy KA. Macrophage CD163 expression in cerebrospinal fluid: association with subarachnoid hemorrhage outcome. *J Neurosurg.* 2018;1-7.
 26. Ling EA, Tseng CY, Wong WC. An electron microscopical study of epiplexus and supraependymal cells in the prenatal rat brain following a maternal injection of 6-aminonicotinamide. *J Anat.* 1985;140 (Pt 1):119-129.
 27. Zusso M, Methot L, Lo R, Greenhalgh AD, David S, Stifani S. Regulation of postnatal forebrain amoeboid microglial cell proliferation and development by the transcription factor Runx1. *J Neurosci.* 2012;32(33):11285-11298.
 28. Lan X, Han X, Li Q, Yang QW, Wang J. Modulators of microglial activation and polarization after intracerebral haemorrhage. *Nat Rev Neurol.* 2017;13(7):420-433.
 29. Hendrickx DAE, van Eden CG, Schuurman KG, Hamann J, Huitinga I. Staining of HLA-DR, Iba1 and CD68 in human microglia reveals partially overlapping expression depending on cellular morphology and pathology. *J Neuroimmunol.* 2017;309:12-22.
 30. Walker DG, Lue LF. Immune phenotypes of microglia in human neurodegenerative disease: challenges to detecting microglial polarization in human brains. *Alzheimers Res Ther.* 2015;7(1):56.
 31. Hopperton KE, Mohammad D, Trepanier MO, Giuliano V, Bazinet RP. Markers of microglia in post-mortem brain samples from patients with Alzheimer's disease: a systematic review. *Mol Psychiatry.* 2018;23(2):177-198.

Figure Legends

Figure 1. (A) Iba-1, OX-6 and CD68 immunoreactivity in epiplexus cells of rats at 24 hours after sham operation. Scale bar=100 μm in first column, 20 μm in second column and 10 μm for third column. (B) Immunofluorescence staining of Iba-1 and the double labeling of Iba-1 with OX-6 or CD68 in epiplexus cells at 24 hours in sham operated rats. Scale bar=20 μm .

Figure 2. (A) Representative examples of T2-weighted magnetic resonance images of rats in sham, SAH without (w/o) hydrocephalus (HC), and SAH with HC groups at 24 hours after procedure. (B) Iba-1 immunoreactivity in the ependymal cells of rats at 24 hours in sham, SAH w/o HC, and SAH with HC groups. Scale bars=100 μm (upper row) and 20 μm (lower row). The number of Iba-1 positive cells was quantified relative to the number of choroid plexus epithelial cells. The values are means \pm SD; **n=10 in sham group, n=12 in SAH w/o HC group and n=15 in SAH with HC group**, * $p < 0.05$ vs. sham group, # $p < 0.01$ vs. sham group and SAH w/o HC group by one-way ANOVA. (C) Iba-1 positive cell soma size in the ependymal cells of rats at 24 hours in sham, SAH w/o HC, and SAH with HC groups. Values are means \pm SD; **n=10 in sham group, n=12 in SAH w/o HC group and n=15 in SAH with HC group**, * $p < 0.05$ vs. sham group, # $p < 0.01$ vs. sham group and SAH w/o HC group by one-way ANOVA. Scale bars=100 μm (upper row) and 10 μm (lower row).

Figure 3. CD68 immunoreactivity in ependymal cells in sham, SAH without (w/o) hydrocephalus (HC), and SAH with HC groups. The number of CD68 positive cells was quantified relative to the number of choroid plexus epithelial cells. Values are mean \pm SD; **n=7 in sham group, n=8 in SAH w/o HC group and n=9 in SAH with HC groups**, # $p < 0.01$ vs. sham group and SAH with HC group, *# $p < 0.01$ vs. sham group by one-way ANOVA. Scale bar=100 μm (upper row) and 20 μm (lower row).

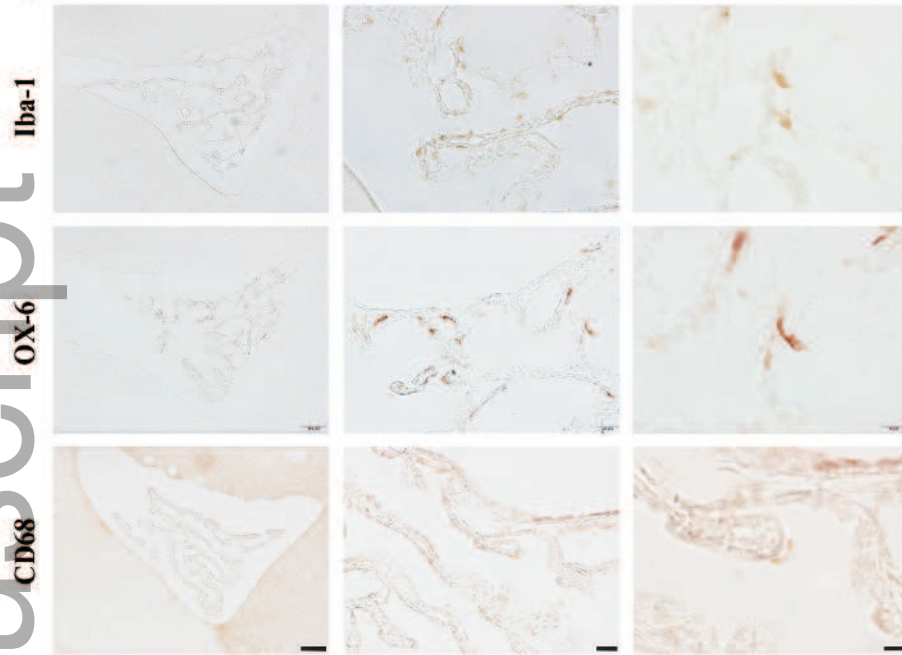
Figure 4. (A) Examples of T2-weighted MRI scan 24 hours after intraventricular injection of saline or thrombin. Note the dilated ventricles in the thrombin injected rats. Ventricular volume was quantified (bar graph). Values are mean \pm SD; **n=11 in saline group and n=14 in thrombin group**, # $p < 0.01$ vs. injected saline group by Student *t* test. (B) Iba-1 immunoreactivity in ependymal cells 24 hours after intraventricular injections of saline or thrombin (3U). The number of Iba-1 positive cells was calculated and expressed relative to the number of choroid plexus epithelial cells. Values are means \pm SD; **n=11 in saline group and n=14 in thrombin group**, # $p < 0.01$ vs. injected saline group by Student *t* test. Scale bar=100 μm (upper row) and 20 μm (lower row). (C) Iba-1 positive cell soma size at 24 hours after injection of 50 μl of saline or thrombin (3U) in saline. Values are means \pm SD; **n=11 in saline group and n=14 in thrombin**

group, # $p < 0.01$ vs. injected saline group by Student t test. Scale bar=100 μm (upper row) and 10 μm (lower row).

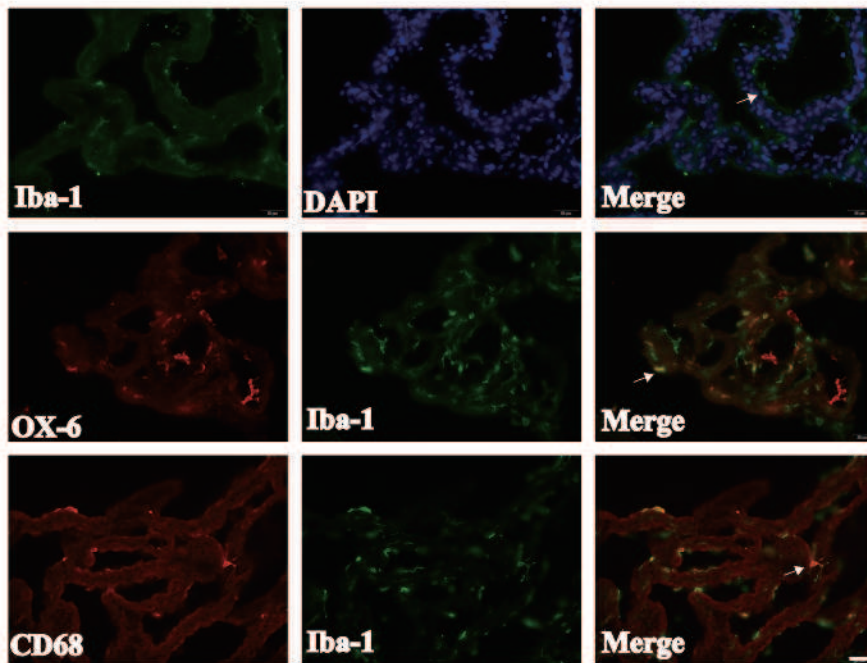
Figure 5. (A) CD68 immunoreactivity in epileptus cells 24 hours after intraventricular injection of saline or thrombin (3U). The number of Cd68 positive cells was quantified and expressed relative to the number of choroid plexus epithelial cells. Values are means \pm SD; **n=11 in saline group and n=14 in thrombin group**, # $p < 0.01$ vs. saline group by Student t test. Scale bars=100 μm (upper row), 20 μm (lower row). (B) Double-staining of CD68 and Iba-1 in epileptus cells in saline and thrombin groups at 24 hours. The percentage of Iba-1 positive epileptus cells that were also CD68 positive also increased with thrombin injection. Values are means \pm SD; n=6, # $p < 0.01$ vs. saline group by Student t test; scale bar= 20 μm .

Author Manuscript

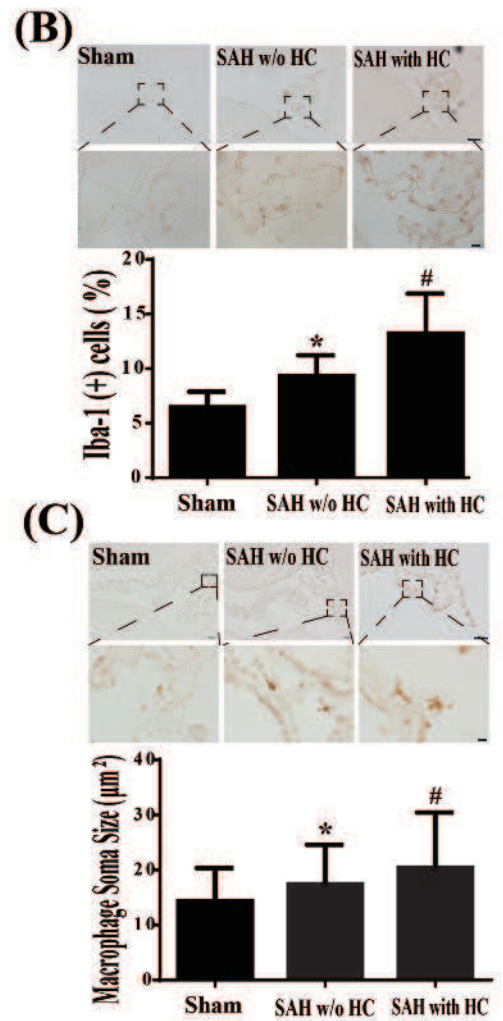
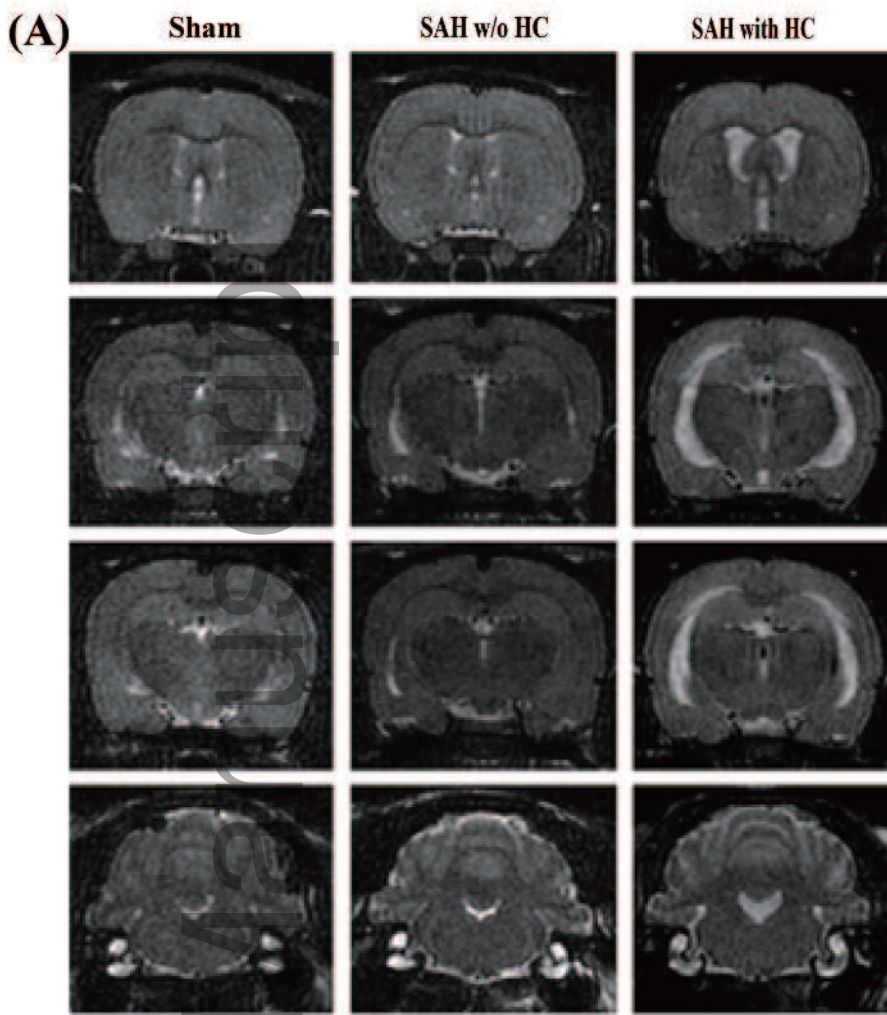
(A)



(B)



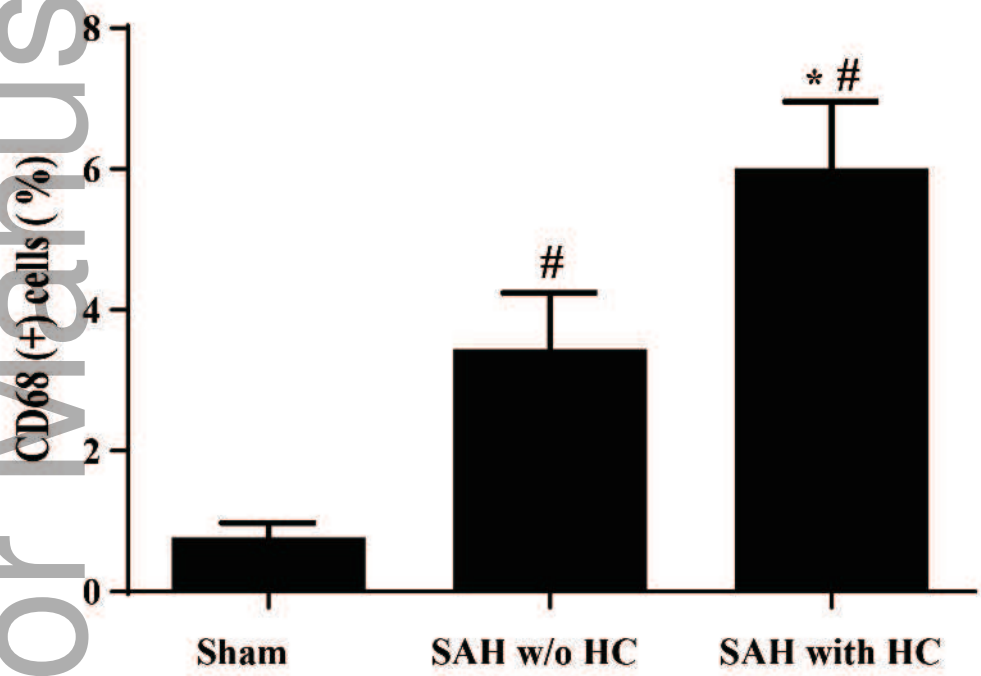
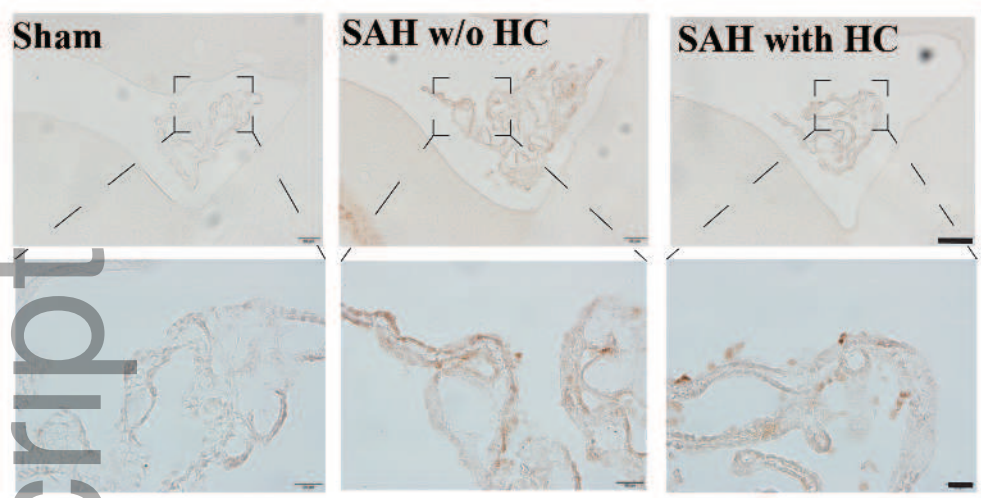
cns_13203_f1.tif



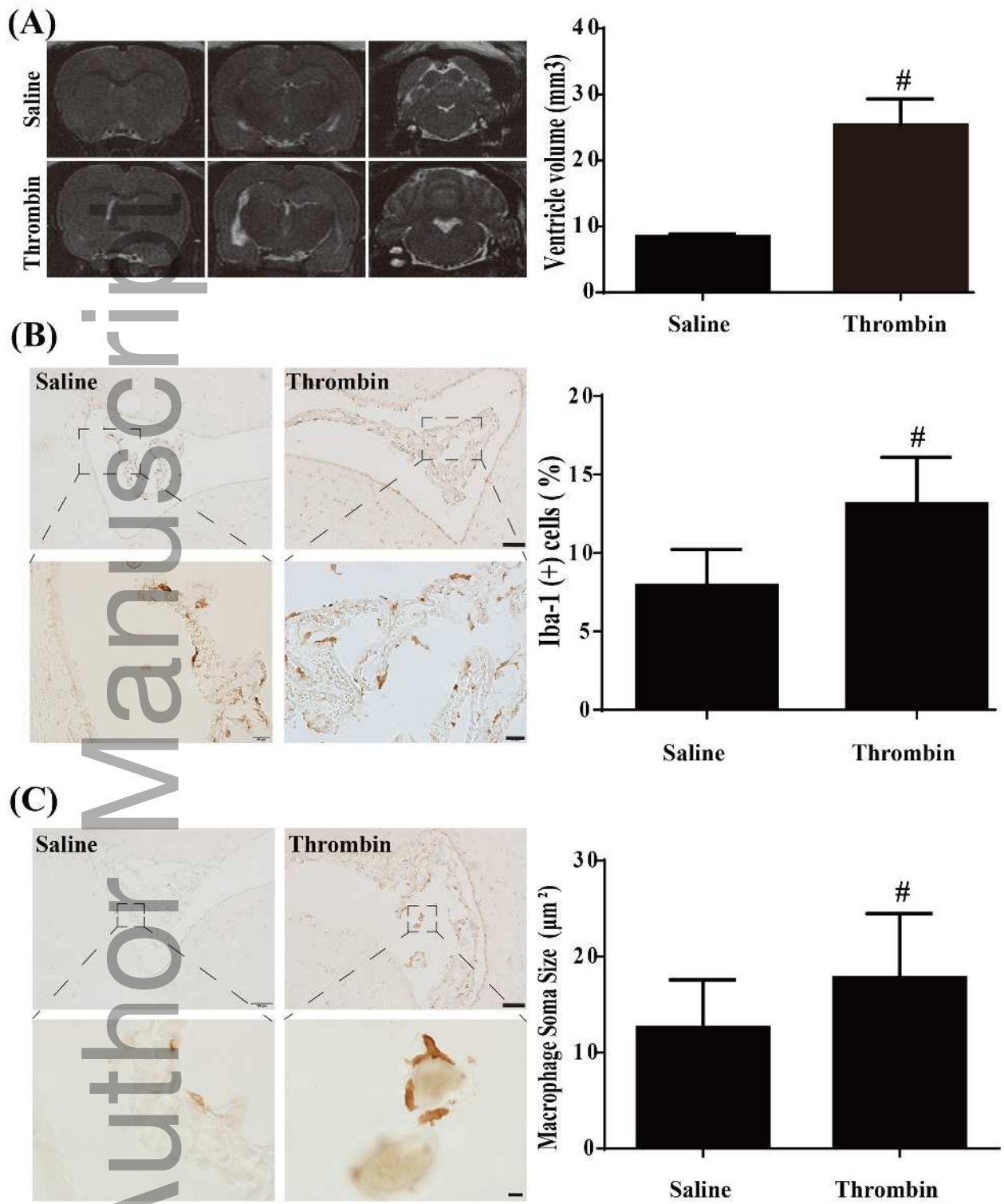
cns_13203_f2.tif

Author

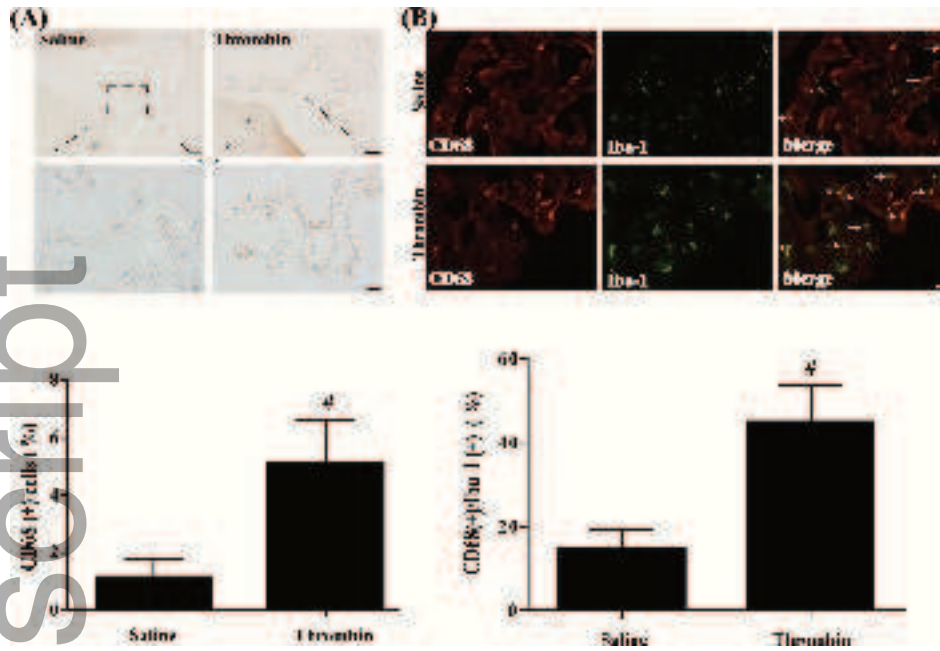
Author Manuscript



cns_13203_f3.tif



cns_13203_f4.tif



cns_13203_f5.tif