

● REVIEW

Fast-tracking regenerative medicine for traumatic brain injury

Brooke Bonsack[‡], Matt Heyck[‡], Chase Kingsbury[‡], Blaise Cozene, Nadia Sadanandan, Jea-Young Lee, Cesar V. Borlongan^{*}

Center of Excellence for Aging and Brain Repair, University of South Florida College of Medicine, Tampa, FL, USA

Funding: CVB was funded by National Institutes of Health (NIH) R01NS071956, NIH R01NS090962, NIH R21NS089851, NIH R21NS094087, and Veterans Affairs Merit Review 101 BX001407.

Abstract

Traumatic brain injury remains a global health crisis that spans all demographics, yet there exist limited treatment options that may effectively curtail its lingering symptoms. Traumatic brain injury pathology entails a progression from primary injury to inflammation-mediated secondary cell death. Sequestering this inflammation as a means of ameliorating the greater symptomology of traumatic brain injury has emerged as an attractive treatment prospect. In this review, we recapitulate and evaluate the important developments relating to regulating traumatic brain injury-induced neuroinflammation, edema, and blood-brain barrier disintegration through pharmacotherapy and stem cell transplants. Although these studies of stand-alone treatments have yielded some positive results, more therapeutic outcomes have been documented from the promising area of combined drug and stem cell therapy. Harnessing the facilitatory properties of certain pharmaceuticals with the anti-inflammatory and regenerative effects of stem cell transplants creates a synergistic effect greater than the sum of its parts. The burgeoning evidence in favor of combined drug and stem cell therapies warrants more elaborate preclinical studies on this topic in order to pave the way for later clinical trials.

Key Words: clinical trials; combined therapy; inflammatory cascade; neuroinflammation; neuroprotection; neurotrauma; pharmacotherapy; preclinical studies; secondary cell death; stem cells

*Correspondence to:

Cesar V. Borlongan, PhD,
cborlong@health.usf.edu.

#These authors contributed equally to this work.

orcid:

0000-0002-2966-9782
(Cesar V. Borlongan)

doi: 10.4103/1673-5374.270294

Received: August 16, 2019

Peer review started: August 22, 2019

Accepted: September 26, 2019

Published online: January 6, 2020

Inflammation Represents an Attractive Target for Treating Traumatic Brain Injury

Traumatic brain injury (TBI) refers to brain damage and dysfunction resulting from an external blow or jolt to the head, such as a puncture, blunt impact, or blast (Maas et al., 2008). Representing a considerable source of health and economic costs, TBI influences approximately 1.7 million lives per year (Acosta et al., 2015a). Additionally, the frequency of TBI-related military casualties has increased markedly due to escalated usage of explosives in modern warfare (Okie, 2005). In humans, TBI severity—categorized as mild, moderate, or severe—can be evaluated using medical imaging techniques and by assessing a patient's level of consciousness with the Glasgow Coma Scale (Lozano et al., 2015). Symptoms of mild TBI, including but not limited to headache, dizziness, fatigue, and nausea, may be temporary and clear up in the short-term; however, severe TBI may lead to chronic neurological problems akin to the degenerative pathological hallmarks of Alzheimer's disease, Parkinson's disease, and dementia pugilistica (Colmano and Gross, 1971; Lozano et al., 2015). According to epidemiological data, suffering a TBI also elevates an individual's risk for developing Alzheimer's disease and Parkinson's disease later in life, but the mechanisms underlying these processes remain unclear (Lozano et al., 2015).

Although TBI was once exclusively deemed an acute injury, its pathology is now divided into the acute and chronic temporal phases. The relatively brief acute phase comprises the immediate neurostructural damage resulting from TBI, whereas the chronic phase is distinguished by rampant

inflammation, secondary cell loss, and impaired neural function which may last for years and is finally irreversible (Werner and Engelhard, 2007; Acosta et al., 2015a). Harmful byproducts arising from the primary injury site, such as pro-inflammatory molecules and reactive species, become dispersed throughout the brain and cause cyclic cascades of neural cell damage (Zhao et al., 2005; Acosta et al., 2015a). The accumulation of such damage forms the penumbra—a region of dead and vulnerable brain tissue surrounding the primary injury core—which chiefly contributes to the risk for chronic evolution of cognitive symptoms (Zhao et al., 2005; Acosta et al., 2015a). However, due to the potentially broad window for effective treatment, alleviating this progressive penumbral cell loss represents a promising approach to improve long-term outcomes for TBI patients (Zhao et al., 2005; Acosta et al., 2015a).

The peri-impact area is a common pathological characteristic of chronic TBI which manifests as the tissue in the vicinity of the focal injury site displaying a myriad of intricate metabolic, immune, and cellular responses (Kumar and Loane, 2012; Lozano et al., 2015). In contrast, the specific pathological progression of these responses, such as their possible spread to remote brain regions, varies based on the location and severity of the TBI (Kumar and Loane, 2012; Lozano et al., 2015; Pabon et al., 2016). Intervention to mitigate the primary damage of a TBI—including necrosis, excitotoxicity, and the mangling of local neuronal, microglial, and vascular structure—is largely impractical, and use of safety equipment such as helmets and seatbelts constitutes

the only feasible means of prevention (Hirschenfang et al., 1968; Kumar and Loane, 2012; Lozano et al., 2015). Thus, recent laboratory investigations have concentrated on the chronic symptoms of TBI in order to suppress secondary cell death mechanisms following the acute phase (Lozano et al., 2015). Due to the relatively delayed onset of inflammatory processes compared to other secondary cell death pathways within the injured brain, targeting post-TBI neuroinflammation has emerged as an especially attractive approach to reduce secondary cell loss and limit neurological deficits (Matsumoto et al., 1986; Offit et al., 1986; Lozano et al., 2015). While inflammation actually protects the injured brain during the acute phase of TBI, a prolonged inflammatory condition throughout the chronic phase significantly exacerbates neural cell death (Lozano et al., 2015). Indeed, an extensive body of evidence demonstrates that attenuating chronic inflammation confers neuroprotective effects (Lozano et al., 2015). Inflammation represents the net result of the competitive interaction between numerous pro-inflammatory (e.g., tumor necrosis factor alpha (TNF- α) and interleukin (IL)-1 β) and anti-inflammatory (e.g., IL-10 and transforming growth factor beta (TGF- β)) molecules, and the shift from a protective to degenerative state following TBI further displays the complex pathology underlying neuroinflammation (Lozano et al., 2015).

After TBI, inflammation-mediated neuronal loss and edema are likely intensified due to disruption of the blood-brain barrier (BBB) (Shlosberg et al., 2010; Neuwelt et al., 2011). The BBB, a highly selective layer of endothelium surrounding cerebral vasculature, regulates the passage of blood-borne compounds into the brain, and the neuroprotective capacity of the BBB hinges upon its structural integrity and the health of its endothelial cells, both of which are maintained by astrocytes (Neuwelt et al., 2011). The physical trauma of TBI impairs BBB structure and function, heightening its permeability and permitting peripheral immune cells, along with exogenous proteins such as albumin, fibrinogen, and thrombin, to infiltrate the parenchymal tissue of the brain (Shlosberg et al., 2010; Neuwelt et al., 2011; Lozano et al., 2015). This foreign influx precipitates microglial activation, which, although initially functioning as a protective immune response, will continue without restriction and may become self-perpetuating unless BBB integrity is restored (Shlosberg et al., 2010; Neuwelt et al., 2011; Lozano et al., 2015). Moreover, pro-inflammatory substances such as cytokines, chemokines, prostaglandins, and free radicals are secreted by the intruding peripheral immune cells, amplifying the inflammatory cascade and further destabilizing the BBB (Lozano et al., 2015). Subsequently, large molecules and cells more readily penetrate the biological barricade, generating an osmotic gradient in the brain which leads to the formation of cerebral edema and the buildup of intracranial pressure (Neuwelt et al., 2011; Lozano et al., 2015). Furthermore, elevated levels of matrix metalloproteinase-9 (MMP-9), vascular endothelial growth factor (VEGF), and other molecules damage tight junctions, continuously inhibiting repair of the BBB (Guo et al., 1989; Lozano et al., 2015).

Within the injured brain, immunological processes become even more complex due to heterogeneous microglial activation (Lozano et al., 2015). More specifically, in response to their local microenvironment, activated microglia, which serve as the brain's resident macrophages under normal conditions, may differentiate into a pro-inflammatory M1 phenotype or an anti-inflammatory M2 phenotype which suppresses the activity of M1 microglia (Lozano et al., 2015), although it must be noted that this terminology has been disputed recently (Ransohoff, 2016). Therefore, in light of the diverse and complicated inflammatory pathophysiology of chronic TBI, techniques devised to downregulate pro-inflammatory mediators, upregulate anti-inflammatory mediators, and stimulate BBB recovery display significant therapeutic potential in laboratory and clinical settings (Lozano et al., 2015). Previously, we have evaluated the role of inflammation in TBI pathology and the potential of stand-alone pharmacotherapy and stem cell-based therapeutic options (Mashkouri et al., 2016). Here, we expand and update this review, culminating in advancing the potential application of combined drug and stem cell therapies.

This review was compiled using PubMed with sources within the last ten years, with an emphasis on the most recent, novel, and comprehensive papers. If the topic did not have relevant information within the last ten years, we used the most recent paper.

Pharmacotherapeutic Options for Reducing Inflammation

With the goal of reducing secondary neural cell death following TBI, a variety of anti-inflammatory drug therapies have been explored in recent laboratory investigations (**Table 1**). To this end, statins represent a class of drugs with a well-established safety record and discrete, manageable side effects. Traditionally considered a high cholesterol treatment, statins display anti-inflammatory and neuroprotective properties that may expand their applicability to TBI. A murine subarachnoid hemorrhage model demonstrates that pretreatment with rosuvastatin, a type of statin, may reverse the increases in proinflammatory TNF- α , MMP-9, and cyclooxygenase-2 usually observed after TBI (Uekawa et al., 2014). Adding to this, simvastatin reduces IL-1 β in *in vitro* cultures of astrocytes (Wu et al., 2010). The mechanism behind the downregulation of these inflammatory mediators may be statins' inhibition of microgliosis and astrogliosis (Wu et al., 2010; Uekawa et al., 2014). Moreover, laboratory evidence of statins' regulation of epidermal growth factor receptors, some types of small G-proteins, and nuclear factor κ B and toll-like receptor 4 signaling pathways may, in turn, suggest a mechanism behind these reductions in microgliosis and astrogliosis (Takemoto and Liao, 2001; Loane and Faden, 2010; Wu et al., 2010; Uekawa et al., 2014; Wang et al., 2014). Thus, statins may sequester neuroinflammation by modulating several deleterious post-TBI processes. Furthermore, atorvastatin and simvastatin attenuate functional deficits and enhance neuronal survival in preclinical studies (Wang et

Table 1 Milestone studies of drug-based therapies for TBI

Drug	Model or clinical population	Therapeutic effects	Study
Statins			
Simvastatin	Rodent CCI model	Increases neuronal survival and reduces astrocyte activation	Wu et al., 2014
	<i>In vitro</i> astrocytes, oxygen-glucose deprivation (3 h)	Inhibits IL-1 production and alters caveolin-1 expression and restrains epidermal growth factor phosphorylation in lipid rafts, supporting a possible mechanism for the <i>in vivo</i> results	Wu et al., 2014
	Rodent CCI model	Attenuates intercellular adhesion molecule-1 expression, improves grip test score, reduces impact area	Wang et al., 2014
Simvastatin/ Atorvastatin (pretreatment)	Murine weight-drop model	Partially restores cerebral blood flow	Wang et al., 2007
		Reduces neuronal degeneration in hippocampus, improves vestibulomotor function (rotarod times)	Wang et al., 2007
Atorvastatin	Murine weight-drop model	Reduces neuronal degeneration in hippocampus, improves vestibulomotor function (rotarod times), attenuates neurocognitive deficit (Morris water maze times), decreases microglial activation and levels of TNF- α and IL-6 RNA	Wang et al., 2007
Rosuvastatin (pretreatment)	Murine severe subarachnoid hemorrhage model	Improves neurologic score and neuronal survival; decreases edema and immunoglobulin G extravasation (BBB permeability marker); reduces brain superoxide production, NF- κ B activation, and microglial activation; inhibits upregulation of TNF- α , MMP-9, and COX-2	Uekawa et al., 2014
Rosuvastatin	Clinical trial of severe TBI patients aged 16 to 50 years	Lowers amnesia time as assessed by Galveston Orientation Amnesia Test	Tapia-Perez et al., 2008
Minocycline	Murine weight-drop model	Reduces edema and levels of IL-1 β and MMP-9, improves neurological function (string test scores)	Homsy et al., 2009
	Rodent mild blast-induced TBI model	Regulates levels of CRP, MCP-1, claudin 5, neuron-specific enolase, neurofilament-H, tau, S100 β , and corticosterone; microglial growth and activation; anxiety scores (open field and elevated plus maze); spatial memory (Barnes maze times)	Kovesdi et al., 2012
Melatonin	Rodent subarachnoid hemorrhage model	Decreases levels of TNF- α , IL-1 β , IL-6, TLR4 and related agents, NF- κ B, myeloid differentiation factor 88, and inducible nitric oxide synthase; attenuates spatial learning and memory deficits; lowers numbers of apoptosis- and necrosis-positive cells	Wang et al., 2013
	Murine weight-drop model	Decreases levels of TNF- α and IL-1 β and microglial activation, increases peri-impact neuronal survival, dephosphorylates mammalian target of rapamycin pathway	Ding et al., 2014a
	Murine weight-drop model	Attenuates oxidative stress, cortical neuronal degeneration, and edema	Ding et al., 2014b
Melatonin/ Minocycline/ Melatonin + Minocycline	Rodent mild CCI model	No significant differences in Morris water maze, cortical impact area, or microglial activation among the control or treatment groups	Kelso et al., 2011
Progesterone	Rodent moderate CCI model, aged	Improves neurological outcomes (modified neurological severity scores and Morris water maze) and hippocampal long term potentiation, increases number of circulating EPCs, vessel density, and CD31- and CD34-positive cell numbers	Li et al., 2012
	<i>In vitro</i> culture of human peripheral blood mononuclear cells	Increases EPC proliferation when blood mononuclear cells were collected from participants in the menstrual phase but not luteal phase, indicating a hormonal interaction	Matsubara and Matsubara, 2012
	<i>In vitro</i>	Dose-dependently enhances angiogenic potential of EPCs (tube formation, migration, adhesion, VEGF release)	Yu et al., 2017
	Rodent CCI model	Improves neurological function (modified neurological severity scores), vessel density, and occludin and progesterone receptor expression; repairs BBB leakage; reduces brain edema	Yu et al., 2017

BBB: Blood-brain barrier; CCI: controlled cortical impact; COX-2: cyclooxygenase-2; CRP: C-reactive protein; EPCs: endothelial progenitor cells; IL: interleukin; MCP-1: monocyte chemoattractant protein-1; MMP-9: matrix metalloproteinase 9; NF- κ B: nuclear factor- κ B; TBI: traumatic brain injury; TLR4: Toll-like receptor 4; TNF- α : tumor necrosis factor- α ; VEGF: vascular endothelial growth factor.

al., 2007). Echoing this, rosuvastatin pretreatment protects against TBI-induced edema, neurological deficits, neuronal cell death, and BBB disruption in animal models, demonstrating that these anti-inflammatory effects may translate to improved system-wide outcomes (Uekawa et al., 2014). Such findings have helped rosuvastatin advance to a clinical trial, where multiple doses effectuated small reductions of disorientation and amnesia in TBI patients, as measured by the Galveston Orientation Amnesia Test (Tapia-Perez et al., 2008). More elaborate preclinical studies are necessary to assess how clinical administration and outcomes may be best optimized.

Aside from its antibiotic properties, minocycline, a tet-

racycline derivative, may also exert neuroprotective and anti-inflammatory effects. That it has a well-established record as a treatment for several diseases and is approved by the United States Food and Drug Administration advances its translation to TBI. Minocycline also possesses the ability to cross the BBB (Saivin and Houin, 1988), which promotes its use for central nervous system (CNS) disorders. These attributes posit minocycline as an excellent prospect for clinical trials. Expanding this background to TBI, minocycline treatment reduces edema and levels of inflammatory mediators IL-1 β and MMP-9 in a murine weight-drop model (also known as closed head injury model) (Homsy et al., 2009) and normalizes levels of inflammatory markers C-reactive

protein and monocyte chemoattractant protein-1 in a rodent model of mild blast-induced TBI (Kovesdi et al., 2012) when compared to vehicle treatment. Additionally, minocycline may lower nitric oxide, another inflammatory mediator, via its regulation of nitric oxide synthase, demonstrated in an *in vitro* study (Amin et al., 1996). These anti-inflammatory effects are coupled with a reduction in microglial growth and activation (Kovesdi et al., 2012). As microglia support cytokines MMP-9, IL-1 β and IL-6, inhibiting microglial activation may indirectly sequester these cytokines as well (Homsí et al., 2009; Ziebell and Morganti-Kossmann, 2010; Guo et al., 2011). Furthermore, a string test, designed to assess neurological and locomotor functioning by investigating whether TBI mice can climb onto a string, walks, and escapes to a platform, indicates that these anti-inflammatory and anti-edematous effects can translate to functional recovery (Homsí et al., 2009). While the results of some laboratory studies have demonstrated null effects of minocycline on one or more of these dimensions, this may be due to inadequate dosage timing and amounts (Homsí et al., 2009; Kelso et al., 2011). As such, further investigation into minocycline's therapeutic potential is needed.

The neuroprotective and anti-inflammatory qualities of melatonin, a naturally-occurring hormone, similarly posit that this drug serves as another possible pharmacotherapy for TBI. Secreted by the pineal gland, melatonin possesses the ability to diffuse across cell membranes and cross the BBB (Lozano et al., 2015; Lin et al., 2016). Melatonin treatment is well-tolerated by patients and has a history of use for various other diseases (Jahnke et al., 1999; Seabra et al., 2000). Melatonin treatment after induced subarachnoid hemorrhage in rats reduces levels of proinflammatory cytokines TNF- α and IL-1 β (Wang et al., 2013). Likewise, in a murine weight-drop model, the melatonin-treated group exhibits lower levels of these same proinflammatory cytokines when compared to the untreated and vehicle-treated groups (Ding et al., 2014b). These anti-inflammatory effects are coupled with melatonin's demonstrated ability to greatly inhibit microglial activation, as measured by IBA-1 immunofluorescence, a marker of microglia (Ding et al., 2014b). The same model also reveals that melatonin may ameliorate post-TBI neuronal degeneration, oxidative stress, and edema, indicating that melatonin could recapitulate cognitive function in humans (Ding et al., 2014a). However, similar to minocycline, these functional outcomes of melatonin are somewhat inconsistent. When administered alone or in concert with minocycline, melatonin had no effect on cortical tissue integrity or Morris water maze performance in one rodent model (Kelso et al., 2011). This may be due, however, to suboptimal timing and dosages. As it stands, melatonin has been implicated in multiple anti-inflammatory and neuroprotective processes, but these have, at times, failed to translate to functional benefits. To address this disparity, more studies examining the long-term safety and efficacy of melatonin as a TBI therapeutic are needed (Hirschenfang et al., 1968; Lozano et al., 2015).

Overall, pharmacotherapeutic options have demonstrat-

ed promise for attenuating TBI pathology, especially with respect to neuroinflammation. However, these results are at times somewhat tenuous or conflicting. Thus, other options, such as the rapidly emerging field of stem cell transplants, may be considered.

Stem Cell Therapy Ameliorates Inflammation

In addition to anti-inflammatory pharmacotherapy, stem cell therapy represents a promising approach for treating TBI (Table 2). Stem cells have the ability to proliferate and differentiate into a variety of cell lines and entice restorative cytokines to migrate to the injured area (Antonucci et al., 2014; Tajiri et al., 2014a, c). Preclinical data have revealed the efficacy of stem cells to differentiate into neurons after TBI and other CNS disorders (Antonucci et al., 2012; Rodrigues et al., 2012; Liu et al., 2013; Acosta et al., 2014; Tajiri et al., 2014a, b). Thus, direct cell-replacement constitutes one of the ways in which stem cells may attenuate TBI pathology. Secondly, stem cells may secrete various trophic factors that may directly or indirectly support beneficial processes while inhibiting deleterious ones, also known as "bystander effects". Furthermore, evidence of a stem cell-paved "biobridge" by which exogenous stem cells may conduct endogenous stem cells to the impact area to decrease inflammation supports a third method by which stem cells may promote recovery after TBI (Tajiri et al., 2013). Interestingly, this biobridge links the injury site to the subventricular and subgranular zone, which primarily houses neural progenitor cells (Tajiri et al., 2013). This biobridge may thus encourage the migration and differentiation of adult neural progenitor cells. Overall, following brain trauma, stem cell transplantation exhibits a significant ability to ameliorate neural cell death and secondary inflammatory response, leading to improved cognitive and motor outcomes in preclinical settings (Acosta et al., 2014; De la Pena et al., 2014).

Stem cells' secretome plays a major role in their reduction of TBI-induced inflammatory response. Mesenchymal stem/stromal cells (MSCs) possess the ability to travel to the injury site and initiate the cellular effectors of the inflammatory response such as macrophages, neutrophils, T lymphocytes, and microglia (Borlongan, 2011; Borlongan et al., 2011; Zhang et al., 2013). Macrophages represent one component of the inflammatory response and function primarily through phagocytosis (Lin and Du, 2018). In a spinal cord injury model, embryonic stem cell-conditioned media facilitates macrophage phagocytosis of apoptotic cells, thus reducing inflammation and improving recovery (Guo et al., 2016). Moreover, when exposed to inflammatory factors, MSCs may secrete prostaglandin E₂, which, in turn, may induce macrophages to release anti-inflammatory IL-10 and TGF- β (Németh et al., 2009; Chiossone et al., 2016), and which may also influence macrophages toward their anti-inflammatory M2 phenotype or initiate production of new M2 macrophages (Cho et al., 2014; Geng et al., 2014; Vasandan et al., 2016). In addition, promotion of neutrophils, cells that support healing and protect against infection, also holds potential for reducing inflammation. MSCs may secrete

Table 2 Milestone studies of cell-based therapies for TBI

Cell type	Model or clinical population	Therapeutic effects	Study
MSCs			
Bone marrow-derived MSCs	Murine CCI model	Decrease neutrophil extravasation, MMP-9 expression, and BBB leakage, also implicating TSG-6 as a mediator for these effects	Watanabe et al., 2013
	Rodent weight-drop model	Diminish microglia/macrophage and leukocyte density; increase anti-inflammatory cytokines (IL-10, TGF-1 β); decrease pro-inflammatory cytokines (IL-1 β , IL-6, IL-17, TNF- α , IFN- γ) and MCP-1, macrophage inflammatory protein-2, and RANTES; enhance TSG-6 expression; reduce edema and neurological deficit (neurological severity score)	Zhang et al., 2013
	<i>In vitro</i> co-culture of MSCs and bone marrow-derived macrophages	Reduce M1 markers including IL-1, IL-6, monocyte chemoattractant protein-1, and inducible nitric oxide synthase; increase M2 markers including IL-4, IL-10, arginase-1, and CD206	Cho et al., 2014
	<i>In vitro</i> co-culture of MSCs and macrophages incubated with lipopolysaccharide	Influence macrophages toward M2 phenotype, marked by increased IL-10 and CD206	Geng et al., 2014
	<i>In vitro</i> co-culture of monocytes/macrophages and MSCs	Support differentiated monocyte survival, influence macrophages toward M2-like phenotype via prostaglandin E2, increase phagocytic capacity, scavenger receptors, and IL-10 and TGF- β production in M2-like macrophages	Chiossone et al., 2016
	<i>In vitro</i> co-culture of MSCs and bone marrow-derived macrophages	Support macrophage differentiation, amplify respiratory burst, potentiate microbicidal responses in naïve macrophages, increase M2 phenotype activation, shift metabolism, better rescue human endothelial cells, implicating prostaglandin E2-driven mechanism	Vasandan et al., 2016
Bone marrow-derived MSCs (SB623)	Rodent moderate CCI model, chronic	Initiate endogenous cell proliferation and immature neural differentiation, form stem-cell paved biobridge and increase MMP-9 expression/activity, improve motor and neurological functioning (elevated body swing test, rotarod times, Benderson-Neurological scores)	Tajiri et al., 2013
Bone marrow-derived MSCs (SB623)	Phase 2 double-blind, randomized clinical trial of TBI adults	Does not significantly increase risk of adverse effects, improve motor deficits (Fugl-Meyer Motor Scale scores)	Okonkwo et al., 2019
Salivary gland-derived MSCs	<i>In vitro</i> co-culture of MSCs and neutrophils incubated with lipopolysaccharide	Respond to bacterial endotoxin by mobilizing and expressing chemokine receptors, recruit neutrophils by releasing IL-8 and macrophage migration inhibitory factor, increase lifespan and chemokine expression of neutrophils	Brandau et al., 2010
Bone marrow mononuclear cells	Phase 1 clinical trial of severe TBI children	All patients survived, no adverse effects, improve outcomes in majority of patients	Cox et al., 2011
		Attenuate length of treatment for intracranial pressure and severity of injury, suggesting anti-inflammatory effects	Liao et al., 2015
	Phase 1 clinical trial of severe TBI adults	No serious adverse effects, preserve brain structure, improve functional outcomes, and decrease inflammatory cytokines	Cox et al., 2017
Adipose-derived stem cells	Rodent mild CCI model, young	Ameliorate motor (forelimb akinesia, paw grasp) and cognitive (radial arm water maze) functioning, reduce impact and peri-impact area, decrease hippocampal cell loss	Tajiri et al., 2014a
	Rodent mild CCI model, aged	Slightly ameliorate cognitive (radial arm water maze) functioning only, reduce peri-impact area only	Tajiri et al., 2014a
Adipose-derived stem cell-conditioned media	Rodent mild CCI model, young	Ameliorate motor (forelimb akinesia, paw grasp) and cognitive (radial arm water maze) functioning, reduce impact and peri-impact area, decrease hippocampal cell loss	Tajiri et al., 2014a
	Rodent mild CCI model, aged	No significant differences from vehicle group	Tajiri et al., 2014a
Adipose-derived stem cell exosomes	Rodent mild CCI model	Reduce peri-impact area, impact area, and motor deficit, implicating MALAT1 mechanism	Patel et al., 2018
Embryonic stem cell-conditioned media	Murine thoracic spinal contusion injury model/ <i>in vitro</i> myelin-laden macrophages	Rescue myelin-laden macrophage function, enhance exocytosis of internalized lipids, normalize apoptotic cell phagocytosis, decrease TNF- α levels, increase M2 macrophage phenotype, improve locomotor recovery	Guo et al., 2016

BBB: Blood-brain barrier; CCI: controlled cortical impact; IFN- γ : interferon- γ ; IL: interleukin; MCP-1: monocyte chemo-attractant protein-1; MMP-9: matrix metalloproteinase 9; MSCs: mesenchymal stem/stromal cells; NF- κ B: nuclear factor- κ B; TBI: traumatic brain injury; TGF-1 β : transforming growth factor-1 β ; TLR4: Toll-like receptor 4; TNF- α : tumor necrosis factor- α ; TSG-6: TNF- α -stimulated gene/protein 6.

chemokines that attract neutrophils, which in turn curtail inflammation (Brandau et al., 2010; Lin and Du, 2018). Upon recruitment, neutrophils exposed to MSC-derived IL-6 also survive for longer and display enhanced functioning (Lin and Du, 2018). Lastly, MSCs may release various factors to sequester and suppress T cells that would otherwise increase inflammation and secondary cell death (Lin and Du, 2018). Taken together, preclinical models evince that exogenous stem cells may enhance, inhibit, recruit, and generally adjust

the activities of other cell types to reduce inflammation.

Aside from their regulation of other cells to indirectly modulate inflammation, stem cell transplants may also control the secondary cell death cascade by directly releasing anti-inflammatory agents. Key inflammatory signatures, such as TNF- α and IL-1, elicit the secretion of an anti-inflammatory protein known as TNF- α -stimulated gene/protein 6 (TSG-6) by the transplanted MSCs (Watanabe et al., 2013; Zhang et al., 2013). The anti-inflammatory property

of TSG-6 acts by interrupting the inflammatory pathway of toll-like receptors and NF- κ B (Watanabe et al., 2013; Zhang et al., 2013). Furthermore, TSG-6 can mitigate T cell activity by downregulating their production of pro-inflammatory cytokines, such as interferon γ and by upregulating anti-inflammatory cytokines, such as IL-4 (Russo et al., 2011). Thus, TSG-6 exemplifies one of the ways that stem cell therapy may abrogate neuroinflammation via their secretome. Another major way in which stem cells' secretome may reduce inflammation is through their emission of exosomes, a kind of nanoparticle that facilitates the release and transfer of proteins, DNA, mRNA and miRNA (Valadi et al., 2007; Altanerova et al., 2017). Long noncoding RNA metastasis-associated lung adenocarcinoma transcript 1 (MALAT1) is sometimes carried by exosomes originating from human adipose-derived stem cells and represents one of the beneficial agents that may be contained in exogenous stem cells' secretome (Patel et al., 2016, 2018). In brain and spleen tissues, treatment with exosomes containing MALAT1 regulates TBI-induced inflammatory pathways, regenerative pathways, cell cycle and death, and expression of other non-coding RNAs in a rodent controlled cortical impact (CCI) model (Patel et al., 2018). In turn, these MALAT1-controlled mechanisms generate significant post-TBI improvements in impact and peri-impact areas and on locomotor assessments, when compared to groups treated instead with MALAT1-depleted exosomes or with vehicle (Patel et al., 2018). Altogether, stem cell transplants possess multiple mechanisms whereby they may reduce inflammation through secreted factors.

Once the transplanted stem cells reach the host tissue, they are challenged by the harsh environment of the injury site. Granulocyte-colony stimulating factor (G-CSF) can be administered alongside human umbilical cord blood cells (hUCBs) to promote the neuroprotection of the exogenous stem cells (Acosta et al., 2014). Delivering stem cells in combination with factors such as G-CSF yields significant improvements in neurogenesis and the capacity to reduce cell death, when compared to delivering stem cells alone (Acosta et al., 2014). Moreover, G-CSF possesses properties that reduce brain edema, promote the recovery of motor function, and enhance control of glutamate levels (Acosta et al., 2014). G-CSF uses a receptor-mediated transport to mobilize endogenous stem cells from the bone marrow into the peripheral blood. Once in the periphery, the cells can then relocate to the injury site and promote the release of growth factors, chemokines, and cytokines that assist in repairing brain tissue (Acosta et al., 2014). Recent preclinical evidence that stem cells migrate to and survive better in the spleen rather than the brain suggests that direct transplantation may not be as efficacious as a peripheral approach (Acosta et al., 2015b). In fact, there are studies emphasizing systemic delivery of stem cells in TBI and other neurodegenerative models (Acosta et al., 2015b).

Several transplantable cell lines have been tested, with some reaching clinical trials for stroke therapy, such as fetal cells, embryonic stem cells, neural stem/progenitor cells (including NT2N and CTX0E3 cells), umbilical cord blood

cells, amnion cells, adipose cells, and induced pluripotent stem cells (Hara et al., 2008; Li et al., 2008; Stroemer et al., 2009; Kaneko et al., 2011; Liu et al., 2014; De la Pena et al., 2015). Due to its robust safety record in other disease models, bone marrow and its cellular derivatives have been given a particular interest (Borlongan et al., 2011; Steinberg et al., 2016). Various preclinical studies also support that bone marrow-derived stem cells are not only safe, but also effective, in that bone marrow-derived mononuclear cells may preserve the BBB, promote neurogenesis, and facilitate functional outcomes (Cox et al., 2017). Furthermore, different routes of transplantation such as intravenous, intra-arterial, and intranasal as well as direct intracerebral implantation have all demonstrated the functional benefits bone marrow-derived stem cells offer (Borlongan et al., 2004, 2011; Prasad et al., 2014; Acosta et al., 2015b). Such an auspicious record has given researchers the impetus to explore bone marrow mononuclear cells and bone marrow-derived stem cells in clinical studies, where they have demonstrated no difference in adverse events, though further investigations are warranted to evaluate their efficacy (Cox et al., 2011, 2017; Liao et al., 2015; Okonkwo et al., 2019). This notion calls attention to the need for combination therapy in order to improve the outcomes of pharmacotherapeutics and stem cell transplantations in TBI and other neurodegenerative diseases.

Current Efforts in Combined Drug and Stem Cell Therapies for Traumatic Brain Injury

Thus far, the reviewed studies have focused on the status of either stand-alone stem cell transplants or certain pharmacotherapeutics. Interestingly, combinations of the two are emerging as the most promising TBI therapeutics (**Figure 1**). A systematic review of combination therapies for TBI demonstrates the wide variety of potential treatments and groups the results by whether the combination yields enhanced, unaffected, or diminished effects as compared to stand-alone therapies (Kline et al., 2016). Therapies combining stem cell transplants with drugs and other supplements have produced auspicious results, possibly due to the amenability of stem cells to combined use and the ability of some drugs and supplements to facilitate and optimize the ameliorative actions of stem cells (**Table 3**).

Many hormones may produce benefits when administered alone or in concert with stem cells. The sex hormone progesterone has many applications for various CNS disorders due to its penchant for anti-inflammatory and neuroprotective effects. Moreover, that it may encourage the proliferation of endothelial progenitor cells (EPCs) and improve post-TBI neuroregeneration advances progesterone as an attractive option for combination with stem cells (Li et al., 2012; Matsubara and Matsubara, 2012). In an *in vitro* model, administering progesterone to rat EPCs dose-dependently increases secretion of angiogenesis precursors, including VEGF (Yu et al., 2017). Furthermore, as demonstrated by a CCI model of TBI, male rats who receive intraperitoneal progesterone

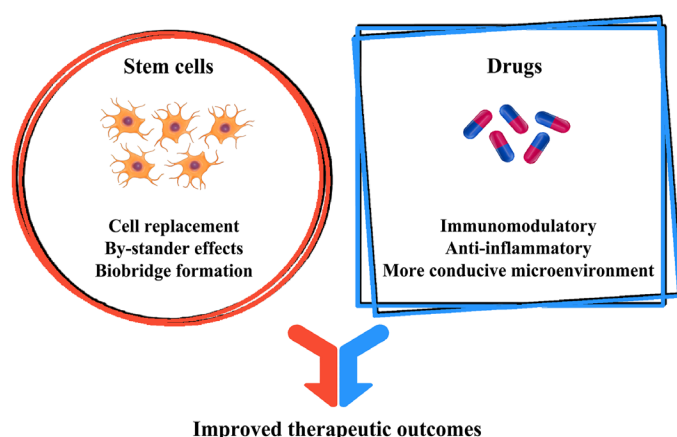


Figure 1 Drugs and stem cells work together to yield enhanced therapeutic outcomes.

During the chronic phase of traumatic brain injury, persisting neuroinflammation contributes significantly to secondary neural cell death. While stand-alone cell therapy can be effective to an extent, combining stem cell transplantation with drug therapies may synergistically attenuate inflammatory neurodegeneration and promote regenerative mechanisms within the brain, thereby enhancing functional recovery following traumatic brain injury.

Table 3 Milestone studies of combination therapies for TBI

Drug, scaffold, precondition, or overexpression	Cell type	Model	Therapeutic effects	Study
Granulocyte-colony stimulating factor	Umbilical cord blood cells	Rodent moderate CCI model, chronic	Significantly reduce neuroinflammation and hippocampal cell loss, improve endogenous neurogenesis and motor function (elevated body swing test and rotarod times); better than monotherapy	Acosta et al., 2014
Progesterone	Embryonic neural stem cells	Rodent CCI model	Improve functional outcomes (open field test, Barnes maze, Morris water maze, rotarod times), especially when additionally combined with enriched environment; better than monotherapy	Nudi et al., 2015
Erythropoietin	Cord blood-derived CD34 MSCs	Rodent CCI model	Engage glial cells, support cell proliferation, increase microvessel density, decrease damaged area, improve inclined plane test scores; all better than monotherapy	Tunc Ata et al., 2016
Propranolol	Bone marrow-derived MSCs	Rodent moderate-severe CCI	Retain benefits of monotherapies (propranolol: decrease activated microglia counts; MSCs: decrease BBB permeability, improve neurogenesis, inhibit activated microglia/macrophage accumulation) and further improve spatial learning and memory (Morris water maze times)	Kota et al., 2016
Hypoxic precondition	Neural progenitor cells	Rodent CCI model, juvenile	Enhance social behaviors (interaction, novelty, transmission of food preference tests) and expression of oxytocin and oxytocin receptors; better than unconditioned cells	Wei et al., 2016
Chitosan, heparin, fibronectin, bFGF scaffold	Radial glial cells	Rodent CCI model	Supports feasibility of scaffolds for promoting cell transplant survival and proliferation	Skop et al., 2016
bFGF-loaded sodium hyaluronate collagen scaffold	Neural stem cells	Rodent hippocampal injury	Modulate microenvironment to improve survival and neuronal differentiation of transplanted cells and increase synaptic formation between endogenous and exogenous cells, improve cognitive outcomes (Morris water maze); better than monotherapy	Duan et al., 2016
BDNF overexpression	Neural stem cells	Rodent CCI model	Increase expression of microtubule-associated protein 2, neurofilament 200, calmodulin, actin, and β -catenin; better than naïve neural stem cells	Chen et al., 2017
PEP-1 and SOD1	Neural stem cells	Rodent CCI model	Dose-dependently promote proliferation and differentiation of transplanted cells, upregulate aquaporin-4 mRNA and protein expression levels, improve functional outcomes (Bederson scores); better than monotherapy	Jia et al., 2018
Docosahexaenoic acid	Neural stem cells (neonatal)	Murine mild CCI model	Reduce motor function deficits (pole climbing test), promote endogenous neurogenesis, increase cortical glial reactivity and microglia, attenuate dopaminergic neuron depletion; better than monotherapy	Ghazale et al., 2018
Chitosan scaffold	Bone marrow-derived MSCs	Rodent Feeney's free fall combat injury model	Promote the survival, proliferation, and differentiation of transplanted cells; improve functional outcomes (modified neurological severity scores, Morris water maze) and brain repair; better than monotherapy	Tan et al., 2018
MDL28170	Bone marrow-derived MSCs	Rodent CCI model	Inhibit inflammation, promote transplanted cell survival/reduce apoptosis of transplanted cells, reduce lesion volume, improve functional outcomes (modified neurological severity scores); better than monotherapy	Hu et al., 2019
IL-10 overexpression	Bone marrow-derived MSCs	Rodent severe CCI model	Improve fine motor functional outcomes (Morris water maze, ladder rung walking task), alter macrophage expression, reduce glial fibrillary acidic protein cell numbers and TNF- α expression	Peruzzaro et al., 2019
		Rodent severe CCI model	Reduce cortical and hippocampal cell loss and neuroinflammation, increase autophagy, mitophagy, and cell survival; better than naïve MSCs	Maiti et al., 2019
FGF-21 overexpression	Bone marrow-derived MSCs	Murine moderate CCI model	Enhance the homing abilities (speed, precision, stability) of transplanted cells to the injury site; better than naïve MSCs	Shahror et al., 2019

BBB: Blood-brain barrier; BDNF: brain derived neurotrophic factor; bFGF: basic fibroblast growth factor; CCI: controlled cortical impact; FGF-21: fibroblast growth factor 21; IL: interleukin; MSCs: mesenchymal stem/stromal cells; SOD1: superoxide-dismutase-1; TBI: traumatic brain injury.

exhibit enhanced vessel density, occludin expression, and BBB integrity, as well as reduced edema and neurological deficit scores, as compared to vehicle-treated rats (Yu et al., 2017). Additionally, that a progesterone receptor-antagonist reverses all of these benefits indicates that progesterone is a key facilitator of endogenous EPC brain repair (Yu et al., 2017). To further support that it is the interaction between progesterone and stem cells that produces these benefits, another study using the CCI model was performed with a double control. Combined therapy of embryonic neural stem cell (eNSC) transplants and progesterone begets improved performances on Morris water maze, open field, and rotarod functional assessments when compared to the control treatment, progesterone-only treatment, or eNSC-only treatment, as well as enhanced survival, migration, and differentiation by the stem cells when compared to the eNSC-only treatment (Nudi et al., 2015). Taken together, these studies indicate that progesterone may greatly stimulate and enable the neuroprotective and anti-inflammatory effects of stem cells, although more double-controlled studies using alternate TBI models are needed (Zibara et al., 2019).

Another naturally-occurring hormone, erythropoietin (EPO), helps to produce red blood cells under normal conditions but may also be applied to TBI treatment because of its neurotrophic, angiogenic, anti-inflammatory, and anti-apoptotic effects (Gonzalez et al., 2007; Bath et al., 2013). In a rodent model, combining EPO treatment with cord blood-derived CD34 MSCs yields enhanced post-TBI cell proliferation, neuronal damage attenuation, glial cell activation, microvessel density, impact area healing, and functional recovery beyond that which is produced by EPO-only or stem cell-only treatment (Tunc Ata et al., 2016). Thus, this demonstrates that EPO and stem cell combination therapy is another promising area for further research.

Various other pharmacotherapeutics may be combined with stem cells to attenuate TBI pathology. Some of these drugs, including G-CSF; peptide carrier PEP-1; antioxidant enzyme Cu,Zn-superoxide dismutase (SOD1); and polyunsaturated fatty acid docosahexaenoic acid (DHA), primarily work by synergizing with stem cells to directly stimulate and support beneficial endogenous processes. As previously discussed, co-administration of G-CSF with hUCBs may confer neuroprotective and anti-inflammatory effects greater than those conferred by stand-alone G-CSF or hUCBs in a CCI model of moderate TBI, possibly due to G-CSF's recruitment of stem cells to the injury site and its suppression of MHC-II microglia activation (Acosta et al., 2014; Liska and de la Pena, 2017). Furthermore, *in vitro* measures of PEP-1 and SOD1's effects on neural stem cells (NSCs) reveal that these drugs may dose-dependently increase NSC proliferation in a rodent model (Jia et al., 2018). This is supported by further tests demonstrating that co-administering PEP-1, SOD1 and NSCs *in vivo* produces higher levels of brain aquaporin-4, which is beneficial for regulating edema in chronic TBI, and improves neurological function, as quantified by the modified Benderson score, when compared to vehicle treatments and stand-alone treatments (Jia et al., 2018). In addition,

DHA may exhibit neuroprotective properties when combined with NSC transplants. Co-administering these two in a CCI murine model increases neurogenesis, glial reactivity, and dopaminergic neurons; decreases calpain/caspase activation; and improves motor function scores, as measured by the pole climbing test, when compared to vehicle treatments and NSC-only treatments (Ghazale et al., 2018).

Other drugs, including propranolol, a β -adrenergic receptor antagonist, and MDL28170, a calpain inhibitor, primarily work by suppressing deleterious factors that may interfere with endogenous or stem cell transplant-mediated reparative processes. Combining propranolol with later administration of human bone marrow-derived MSCs (BM-MSCs) yields complementary benefits greater than either treatment by itself in laboratory settings (Kota et al., 2016; Dekmak et al., 2018). Along with other improvements commensurate with the stand-alone therapies, combined treatment produces greater recovery of cognitive and memory faculties, as compared to vehicle treatments and stand-alone treatments (Kota et al., 2016). Furthermore, MDL28170 facilitates BM-MSC survival and reduces BM-MSC apoptosis after transplantation in a rodent model (Hu et al., 2019). Thus, after experimental TBI, administering MDL28170 followed by BM-MSCs renders lower levels of inflammation, reduces lesion volume, and improves neurological function scores as compared to the vehicle treatment and the stand-alone treatments (Hu et al., 2019). Taken together, many drugs may confer complementary and additive effects when combined with stem cell transplants by enhancing positive processes or by suppressing negative processes.

While the synergy of stem cells and anti-inflammatory drug therapies may represent the most direct approach to counteracting secondary neurodegeneration in chronic TBI, the primary hindrance to stem cell transplants continues to be ensuring their adequate migration, survival, proliferation, and differentiation within the hazardous environment of the TBI brain (Skop et al., 2016; Zibara et al., 2018). Therefore, a number of studies have employed the use of biocompatible scaffolding to more effectively deliver and secure stem cells at injury sites. For example, radial glial cells fixed within an extensively interlinked framework of chitosan, fibronectin, heparin, and basic fibroblast growth factor may be implanted and successfully incorporated into the brains of TBI rats (Skop et al., 2016). Similarly, BM-MSCs complexed with chitosan porous scaffolds can be directly delivered into the lesions of TBI rats without adverse effects (Tan et al., 2018). In fact, this combination treatment increases the survival, proliferation, and differentiation of the BM-MSCs, thereby improving functional recovery (Tan et al., 2018). Moreover, joint transplantation of NSCs and a basic fibroblast growth factor-loaded sodium hyaluronate collagen scaffold enhances the survival of NSCs as well as their differentiation into functional neurons within injured regions of the TBI rat brain (Duan et al., 2016). Along with these effects, the promotion of synaptogenesis in the presence of this scaffold further contributes to the significant amelioration of cognitive deficits (Duan et al., 2016).

In addition to combinatorial treatments with cell-based therapy and pharmaceuticals or scaffolding biomaterials, other recent preclinical studies reveal that modifying the behavior and secretome of stem cell transplants may significantly enhance their anti-inflammatory effects, as well as their migration, viability, and potency in injured regions of the brain. In the context of reprogramming the behavioral aspect of stem cells for TBI treatment, preconditioning NSCs in a hypoxic environment *in vivo* before transplantation activates cell survival mechanisms which prime their resistance to the hostile conditions of the damaged brain (Wei et al., 2016; Zibara et al., 2018). Compared to NSCs cultured in healthy conditions prior to infusion *in vivo*, hypoxia-preconditioned NSCs more effectively attenuate inflammation, elevate release of trophic factors (i.e., glial cell line-derived neurotrophic factor, VEGF, brain-derived neurotrophic factor (BDNF), and EPO), and promote neurological recovery in a juvenile rodent model (Wei et al., 2016). In specific regard to augmenting the secretory functions of exogenous stem cells, inducing the overexpression of various trophic factors has proven to be promising therapeutic approach. Notably, genetically engineered BM-MSCs overexpressing the anti-inflammatory cytokine IL-10 increase the prevalence of autophagy, mitophagy, and cell-survival markers in a rat model, indicating the protection of neural cells from inflammation and oxidative stress (Maiti et al., 2019). These modified BM-MSCs also exert immunomodulatory effects, likely by altering inflammatory cell marker expression by macrophages, which may produce an environment that is more conducive to brain repair and functional recovery after experimental TBI (Peruzzaro et al., 2019). Indeed, MSCs induced to overexpress FGF-21 with this technique display more accurate migratory homing to TBI lesions in a murine model (Shahror et al., 2019). Furthermore, NSC transplants excessively producing BDNF exhibit improvements in survival, growth, proliferation, and differentiation into neurons at lesion sites in a rodent CCI model, potentially due to BDNF-induced upregulation of cytoskeletal protein expression (Chen et al., 2017; Zibara et al., 2018).

Overall, stem cell-based combination therapies stand as attractive strategies for TBI treatment. As previously mentioned, one review categorizes a wide variety of TBI-targeted combination therapies and classifies them by whether the combination produces enhanced, null, or reduced effects (Kline et al., 2016). Among the categories, studies of stem cell-based combination therapies consistently generate the highest proportion of enhanced effects and the lowest proportion of diminished effects when compared to studies of combination therapies not involving stem cells (Kline et al., 2016). Nevertheless, recent research of stem cell-based combination therapy for TBI is somewhat limited. This lack of studies is difficult to explain in light of stem cell-based combination therapies' relative success. The findings of stem cell-based combination therapies are promising, and thus warrant greater examination to advance these therapies to clinical trials.

Conclusion

In this review, a plethora of studies outline different treatment regimens for TBI with pharmacotherapy, cell therapy, or a combination of the two. Throughout the chronic phase of TBI, cyclically perpetuated neuroinflammatory cascades—characterized by upregulated levels of pro-inflammatory molecules, downregulated levels of anti-inflammatory molecules, and dysfunction of the BBB—cause devastating secondary cell death in the brain. While other mechanisms contribute to this neurodegeneration, the delayed onset and long-term progression of inflammatory processes are indicative of a wide therapeutic window for intervention. Thus, reducing aberrant neuroinflammation represents a promising approach to ameliorate secondary cell loss and promote neurological and functional recovery in TBI victims. Several pharmacological therapies demonstrate anti-inflammatory effects; however, inadequate dosages and poor clinical models have hindered their observed efficacy as TBI treatments. Instead, stem cells have attracted the attention of clinicians due to their capacity to replace lost brain cells, secrete neurotrophic factors, and recruit restorative cytokines and endogenous stem cells to the injured area. Furthermore, a combination of both pharmacotherapeutics and stem cells has yielded the most favorable outcomes. Certain drugs amplify the neuroregenerative properties of stem cells and provide a more suitable environment for the cells of the brain. Altogether, further investigation is needed to optimize treatment parameters and ensure safety in larger animal models, as well as to design more effective controlled small animal studies, before stem cell combination therapy can be translated to the clinic.

Author contributions: Drafting, editing, and formatting the manuscript: BB, MH, CK, and CVB; figure creation: CK and JYL; literature retrieval and collection: BC and NS. All authors approved the final manuscript.

Conflicts of interest: CVB was funded and received royalties and stock options from Astellas, Asterias, Sanbio, Athertsys, KMPHC, and International Stem Cell Corporation; and also received consultant compensation for Chiesi Farmaceutici. He also holds patents and patent applications related to stem cell biology and therapy. The other authors have no other relevant affiliations or financial involvement with any organization or entity with a financial interest in or financial conflict with the subject matter or materials discussed in the manuscript apart from those disclosed.

Financial support: CVB was funded by National Institutes of Health (NIH) R01NS071956, NIH R01NS090962, NIH R21NS089851, NIH R21NS094087, and Veterans Affairs Merit Review I01 BX001407.

Copyright license agreement: The Copyright License Agreement has been signed by all authors before publication.

Plagiarism check: Checked twice by iThenticate.

Peer review: Externally peer reviewed.

Open access statement: This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-Non-Commercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

Open peer reviewer: Elizabeth D. Kirby, The Ohio State University, USA.

Additional file: Open peer review report 1.

References

- Acosta SA, Tajiri N, de la Pena I, Bastawrous M, Sanberg PR, Kaneko Y, Borlongan CV (2015a) Alpha-synuclein as a pathological link between chronic traumatic brain injury and Parkinson's disease. *J Cell Physiol* 230:1024–1032.

- Acosta SA, Tajiri N, Hoover J, Kaneko Y, Borlongan CV (2015b) Intravenous bone marrow stem cell grafts preferentially migrate to spleen and abrogate chronic inflammation in stroke. *Stroke* 46:2616-2627.
- Acosta SA, Tajiri N, Shinozuka K, Ishikawa H, Sanberg PR, Sanchez-Ramos J, Song S, Kaneko Y, Borlongan CV (2014) Combination therapy of human umbilical cord blood cells and granulocyte colony stimulating factor reduces histopathological and motor impairments in an experimental model of chronic traumatic brain injury. *PLoS One* 9:e90953.
- Altanerova U, Jakubecova J, Repiska V, Altaner C (2017) Exosomes of human mesenchymal stem/stromal/medicinal signaling cells. *Neoplasma* 64:809-815.
- Amin AR, Attur MG, Thakker GD, Patel PD, Vyas PR, Patel RN, Patel IR, Abramson SB (1996) A novel mechanism of action of tetracyclines: effects on nitric oxide synthases. *Proc Natl Acad Sci U S A* 93:14014-14019.
- Antonucci I, Pantalone A, Tete S, Salini V, Borlongan CV, Hess D, Stuppia L (2012) Amniotic fluid stem cells: a promising therapeutic resource for cell-based regenerative therapy. *Curr Pharm Des* 18:1846-1863.
- Antonucci I, Di Pietro R, Alfonsi M, Centurione MA, Centurione L, Sancilio S, Pelagatti F, D'Amico MA, Di Baldassarre A, Piattelli A, Tete S, Palka G, Borlongan CV, Stuppia L (2014) Human second trimester amniotic fluid cells are able to create embryoid body-like structures in vitro and to show typical expression profiles of embryonic and primordial germ cells. *Cell Transplant* 23:1501-1515.
- Bath PM, Sprigg N, England T (2013) Colony stimulating factors (including erythropoietin, granulocyte colony stimulating factor and analogues) for stroke. *Cochrane Database Syst Rev* 6:CD005207.
- Borlongan CV (2011) Bone marrow stem cell mobilization in stroke: a 'bonehead' may be good after all! *Leukemia* 25:1674-1686.
- Borlongan CV, Lind JG, Dillon-Carter O, Yu G, Hadman M, Cheng C, Carroll J, Hess DC (2004) Intracerebral xenografts of mouse bone marrow cells in adult rats facilitate restoration of cerebral blood flow and blood-brain barrier. *Brain Res* 1009:26-33.
- Borlongan CV, Glover LE, Tajiri N, Kaneko Y, Freeman TB (2011) The great migration of bone marrow-derived stem cells toward the ischemic brain: therapeutic implications for stroke and other neurological disorders. *Prog Neurobiol* 95:213-228.
- Brandau S, Jakob M, Hemeda H, Bruderek K, Janeschik S, Bootz F, Lang S (2010) Tissue-resident mesenchymal stem cells attract peripheral blood neutrophils and enhance their inflammatory activity in response to microbial challenge. *J Leukoc Biol* 88:1005-1015.
- Chen T, Yu Y, Tang LJ, Kong L, Zhang C, Chu H, Yin L, Ma H (2017) Neural stem cells over-expressing brain-derived neurotrophic factor promote neuronal survival and cytoskeletal protein expression in traumatic brain injury sites. *Neural Regen Res* 12:433-439.
- Chiossone L, Conte R, Spaggiari GM, Serra M, Romei C, Bellora F, Becchetti F, Andaloro A, Moretta L, Bottino C (2016) Mesenchymal stromal cells induce peculiar alternatively activated macrophages capable of dampening both innate and adaptive immune responses. *Stem Cells* 34:1909-1921.
- Cho DI, Kim MR, Jeong HY, Jeong HC, Jeong MH, Yoon SH, Kim YS, Ahn Y (2014) Mesenchymal stem cells reciprocally regulate the M1/M2 balance in mouse bone marrow-derived macrophages. *Exp Mol Med* 46:e70.
- Colmano G, Gross WB (1971) Effect of metyrapone and DDD on infectious diseases. *Poult Sci* 50:850-854.
- Cox CS, Baumgartner JE, Harting MT, Worth LL, Shah SK, Ewing-Cobbs L, Hasan KM, Day MC, Lee D, Jimenez F, Gee A (2011) Autologous bone marrow mononuclear cell therapy for severe traumatic brain injury in children. *Neurosurgery* 68:588-600.
- Cox CS Jr, Hetz RA, Liao GP, Aertker BM, Ewing-Cobbs L, Juranek J, Savitz SI, Jackson ML, Romanowska-Pawliczek AM, Triolo F, Dash PK, Pedroza C, Lee DA, Worth L, Aisiku IP, Choi HA, Holcomb JB, Kitagawa RS (2017) Treatment of severe adult traumatic brain injury using bone marrow mononuclear cells. *Stem Cells* 35:1065-1079.
- De la Pena I, Sanberg PR, Acosta S, Lin SZ, Borlongan CV (2014) Umbilical cord blood cell and granulocyte-colony stimulating factor: combination therapy for traumatic brain injury. *Regen Med* 9:409-412.
- De la Pena I, Sanberg PR, Acosta S, Lin SZ, Borlongan CV (2015) G-CSF as an adjunctive therapy with umbilical cord blood cell transplantation for traumatic brain injury. *Cell Transplant* 24:447-457.
- Dekmak A, Mantash S, Shaito A, Toutonji A, Ramadan N, Ghazale H, Kassem N, Darwish H, Zibara K (2018) Stem cells and combination therapy for the treatment of traumatic brain injury. *Behav Brain Res* 340:49-62.
- Ding K, Wang H, Xu J, Li T, Zhang L, Ding Y, Zhu L, He J, Zhou M (2014a) Melatonin stimulates antioxidant enzymes and reduces oxidative stress in experimental traumatic brain injury: the Nrf2-ARE signaling pathway as a potential mechanism. *Free Radic Biol Med* 73:1-11.
- Ding K, Wang H, Xu J, Lu X, Zhang L, Zhu L (2014b) Melatonin reduced microglial activation and alleviated neuroinflammation induced neuron degeneration in experimental traumatic brain injury: possible involvement of mTOR pathway. *Neurochem Int* 76:23-31.
- Duan H, Li X, Wang C, Hao P, Song W, Li M, Zhao W, Gao Y, Yang Z (2016) Functional hyaluronate collagen scaffolds induce NSCs differentiation into functional neurons in repairing the traumatic brain injury. *Acta Biomater* 45:182-195.
- Geng Y, Zhang L, Fu B, Zhang J, Hong Q, Hu J, Li D, Luo C, Cui S, Zhu F, Chen X (2014) Mesenchymal stem cells ameliorate rhabdomyolysis-induced acute kidney injury via the activation of M2 macrophages. *Stem Cell Res Ther* 5:80.
- Ghazale H, Ramadan N, Mantash S, Zibara K, El-Sitt S, Darwish H, Chamaa F, Boustany RM, Mondello S, Abou-Kheir W, Soueid J, Kobeissy F (2018) Docosahexaenoic acid (DHA) enhances the therapeutic potential of neonatal neural stem cell transplantation post-traumatic brain injury. *Behav Brain Res* 340:1-13.
- Gonzalez FF, McQuillen P, Mu D, Chang Y, Wendland M, Vexler Z, Ferriero DM (2007) Erythropoietin enhances long-term neuroprotection and neurogenesis in neonatal stroke. *Dev Neurosci* 29:321-330.
- Guo L, Rolfe AJ, Wang X, Tai W, Cheng Z, Cao K, Chen X, Xu Y, Sun D, Li J, He X, Young W, Fan J, Ren Y (2016) Rescuing macrophage normal function in spinal cord injury with embryonic stem cell conditioned media. *Mol Brain* 9:48.
- Guo YS, Singh P, Draviam E, Greeley GH Jr, Thompson JC (1989) Peptide YY inhibits the insulinotropic action of gastric inhibitory polypeptide. *Gastroenterology* 96:690-694.
- Guo ZD, Wu HT, Sun XC, Zhang XD, Zhang JH (2011) Protection of minocycline on early brain injury after subarachnoid hemorrhage in rats. *Acta Neurochir Suppl* 110:71-74.
- Hara K, Yasuhara T, Maki M, Matsukawa N, Masuda T, Yu SJ, Ali M, Yu G, Xu L, Kim SU, Hess DC, Borlongan CV (2008) Neural progenitor NT2N cell lines from teratocarcinoma for transplantation therapy in stroke. *Prog Neurobiol* 85:318-334.
- Hirschenfang S, Silber M, Benton JG (1968) Personality patterns in peripheral neuropathy. *Dis Nerv Syst* 29:46-50.
- Homsy S, Federico F, Croci N, Palmier B, Plotkine M, Marchand-Leroux C, Jafarian-Tehrani M (2009) Minocycline effects on cerebral edema: relations with inflammatory and oxidative stress markers following traumatic brain injury in mice. *Brain Res* 1291:122-132.
- Hu J, Chen L, Huang X, Wu K, Ding S, Wang W, Wang B, Smith C, Ren C, Ni H, ZhuGe Q, Yang J (2019) Calpain inhibitor MDL28170 improves the transplantation-mediated therapeutic effect of bone marrow-derived mesenchymal stem cells following traumatic brain injury. *Stem Cell Res Ther* 10:96.
- Jahnke G, Marr M, Myers C, Wilson R, Travlos G, Price C (1999) Maternal and developmental toxicity evaluation of melatonin administered orally to pregnant Sprague-Dawley rats. *Toxicol Sci* 50:271-279.
- Jia J, Chen F, Wu Y (2018) Recombinant PEP-1-SOD1 improves functional recovery after neural stem cell transplantation in rats with traumatic brain injury. *Exp Ther Med* 15:2929-2935.
- Kaneko Y, Hayashi T, Yu S, Tajiri N, Bae EC, Solomita MA, Chheda SH, Weinbren NL, Parolini O, Borlongan CV (2011) Human amniotic epithelial cells express melatonin receptor MT1, but not melatonin receptor MT2: a new perspective to neuroprotection. *J Pineal Res* 50:272-280.

- Kelso ML, Scheff NN, Scheff SW, Pauly JR (2011) Melatonin and minocycline for combinatorial therapy to improve functional and histopathological deficits following traumatic brain injury. *Neurosci Lett* 488:60-64.
- Kline AE, Leary JB, Radabaugh HL, Cheng JP, Bondi CO (2016) Combination therapies for neurobehavioral and cognitive recovery after experimental traumatic brain injury: Is more better? *Prog Neurobiol* 142:45-67.
- Kota DJ, Prabhakara KS, van Brummen AJ, Bedi S, Xue H, DiCarlo B, Cox CS Jr, Olson SD (2016) Propranolol and mesenchymal stromal cells combine to treat traumatic brain injury. *Stem Cells Transl Med* 5:33-44.
- Kovesdi E, Kamnakh A, Wingo D, Ahmed F, Grunberg NE, Long JB, Kasper CE, Agoston DV (2012) Acute minocycline treatment mitigates the symptoms of mild blast-induced traumatic brain injury. *Front Neurol* 3:111.
- Kumar A, Loane DJ (2012) Neuroinflammation after traumatic brain injury: opportunities for therapeutic intervention. *Brain Behav Immun* 26:1191-1201.
- Li Z, McKercher SR, Cui J, Nie Z, Soussou W, Roberts AJ, Sallmen T, Lipton JH, Talantova M, Okamoto S, Lipton SA (2008) Myocyte enhancer factor 2C as a neurogenic and antiapoptotic transcription factor in murine embryonic stem cells. *J Neurosci* 28:6557-6568.
- Li Z, Wang B, Kan Z, Zhang B, Yang Z, Chen J, Wang D, Wei H, Zhang JN, Jiang R (2012) Progesterone increases circulating endothelial progenitor cells and induces neural regeneration after traumatic brain injury in aged rats. *J Neurotrauma* 29:343-353.
- Liao GP, Harting MT, Hetz RA, Walker PA, Shah SK, Corkins CJ, Hughes TG, Jimenez F, Kosmach SC, Day MC, Tsao K, Lee DA, Worth LL, Baumgartner JE, Cox CS Jr (2015) Autologous bone marrow mononuclear cells reduce therapeutic intensity for severe traumatic brain injury in children. *Pediatr Crit Care Med* 16:245-255.
- Lin C, Chao H, Li Z, Xu X, Liu Y, Hou L, Liu N, Ji J (2016) Melatonin attenuates traumatic brain injury-induced inflammation: a possible role for mitophagy. *J Pineal Res* 61:177-186.
- Lin L, Du L (2018) The role of secreted factors in stem cells-mediated immune regulation. *Cell Immunol* 326:24-32.
- Liska MG, de la Pena I (2017) Granulocyte-colony stimulating factor and umbilical cord blood cell transplantation: Synergistic therapies for the treatment of traumatic brain injury. *Brain Circ* 3:143-151.
- Liu R, Zhang Z, Lu Z, Borlongan C, Pan J, Chen J, Qian L, Liu Z, Zhu L, Zhang J, Xu Y (2013) Human umbilical cord stem cells ameliorate experimental autoimmune encephalomyelitis by regulating immunoinflammation and remyelination. *Stem Cells Dev* 22:1053-1062.
- Liu SP, Fu RH, Wu DC, Hsu CY, Chang CH, Lee W, Lee YD, Liu CH, Chien YJ, Lin SZ, Shyu WC (2014) Mouse-induced pluripotent stem cells generated under hypoxic conditions in the absence of viral infection and oncogenic factors and used for ischemic stroke therapy. *Stem Cells Dev* 23:421-433.
- Loane DJ, Faden AI (2010) Neuroprotection for traumatic brain injury: translational challenges and emerging therapeutic strategies. *Trends Pharmacol Sci* 31:596-604.
- Lozano D, Gonzales-Portillo GS, Acosta S, de la Pena I, Tajiri N, Kaneko Y, Borlongan CV (2015) Neuroinflammatory responses to traumatic brain injury: etiology, clinical consequences, and therapeutic opportunities. *Neuropsychiatr Dis Treat* 11:97-106.
- Maas AI, Stocchetti N, Bullock R (2008) Moderate and severe traumatic brain injury in adults. *Lancet Neurol* 7:728-741.
- Maiti P, Peruzzaro S, Kolli N, Andrews M, Al-Gharaibeh A, Rossignol J, Dunbar GL (2019) Transplantation of mesenchymal stem cells overexpressing interleukin-10 induces autophagy response and promotes neuroprotection in a rat model of TBI. *J Cell Mol Med* 23:5211-5224.
- Mashkouri S, Crowley MG, Liska MG, Corey S, Borlongan CV (2016) Utilizing pharmacotherapy and mesenchymal stem cell therapy to reduce inflammation following traumatic brain injury. *Neural Regen Res* 11:1379.
- Matsumoto Y, Kitazono S, Ohtsuru Y, Himeno R (1986) Orthodontic treatment for 3 impacted upper anterior teeth. *Fukuoka Shika Daigaku Gakkai Zasshi* 13:60-66.
- Matsubara Y, Matsubara K (2012) Estrogen and progesterone play pivotal roles in endothelial progenitor cell proliferation. *Reprod Biol Endocrinol* 10:2.
- Neal EG, Acosta SA, Kaneko Y, Ji X, Borlongan CV (2019) Regulatory T-cells within bone marrow-derived stem cells actively confer immunomodulatory and neuroprotective effects against stroke. *J Cereb Blood Flow Metab* 39:1750-1758.
- Németh K, Leelahavanichkul A, Yuen PS, Mayer B, Parmelee A, Doi K, Robey PG, Leelahavanichkul K, Koller BH, Brown JM, Hu X, Jelinek I, Star RA, Mezey E (2009) Bone marrow stromal cells attenuate sepsis via prostaglandin E(2)-dependent reprogramming of host macrophages to increase their interleukin-10 production. *Nat Med* 15:42-49.
- Neuwelt EA, Bauer B, Fahlke C, Fricker G, Iadecola C, Janigro D, Leybaert L, Molnar Z, O'Donnell ME, Povlishock JT, Saunders NR, Sharp F, Stanimirovic D, Watts RJ, Drewes LR (2011) Engaging neuroscience to advance translational research in brain barrier biology. *Nat Rev Neurosci* 12:169-182.
- Nudi ET, Jacqmain J, Dubbs K, Geeck K, Salois G, Searles MA, Smith JS (2015) Combining enriched environment, progesterone, and embryonic neural stem cell therapy improves recovery after brain injury. *J Neurotrauma* 32:1117-1129.
- Offit PA, Shaw RD, Greenberg HB (1986) Passive protection against rotavirus-induced diarrhea by monoclonal antibodies to surface proteins vp3 and vp7. *J Virol* 58:700-703.
- Okie S (2005) Traumatic brain injury in the war zone. *N Engl J Med* 352:2043-2047.
- Okonkwo DO, McAllister P, Kawabori M, Chen J, Frishberg B, Achrol A, Gross R, Steinberg G, Imai H, Lu D, Yasuhara T, Bates D (2019) Late Breaking Abstract - Safety and Clinical Outcomes in Traumatic Brain Injury Patients: Interim Analysis of the STEMTRA Trial. Presented at; 2019; American Association of Neurological Surgeons Annual Scientific Meeting. San Diego.
- Pabon MM, Acosta S, Guedes VA, Tajiri N, Kaneko Y, Borlongan CV (2016) Brain region-specific histopathological effects of varying trajectories of controlled cortical impact injury model of traumatic brain injury. *CNS Neurosci Ther* 22:200-211.
- Patel NA, Moss LD, Lee JY, Tajiri N, Acosta S, Hudson C, Parag S, Cooper DR, Borlongan CV, Bickford PC (2018) Long noncoding RNA MALAT1 in exosomes drives regenerative function and modulates inflammation-linked networks following traumatic brain injury. *J Neuroinflammation* 15:204.
- Patel RS, Carter G, El Bassit G, Patel AA, Cooper DR, Murr M, Patel NA (2016) Adipose-derived stem cells from lean and obese humans show depot specific differences in their stem cell markers, exosome contents and senescence: role of protein kinase C delta (PKCδ) in adipose stem cell niche. *Stem Cell Invest* 3:2.
- Peruzzaro ST, Andrews MMM, Al-Gharaibeh A, Pupiec O, Resk M, Story D, Maiti P, Rossignol J, Dunbar GL (2019) Transplantation of mesenchymal stem cells genetically engineered to overexpress interleukin-10 promotes alternative inflammatory response in rat model of traumatic brain injury. *J Neuroinflammation* 16:2.
- Prasad K, Sharma A, Garg A, Mohanty S, Bhatnagar S, Johri S, Singh KK, Nair V, Sarkar RS, Gorthi SP, Hassan KM, Prabhakar S, Marwaha N, Khandelwal N, Misra UK, Kalita J, Nityanand S, InveST Study Group (2014) Intravenous autologous bone marrow mononuclear stem cell therapy for ischemic stroke: a multicentric, randomized trial. *Stroke* 45:3618-3624.
- Ransohoff RM (2016) A polarizing question: do M1 and M2 microglia exist? *Nat Neurosci* 19:987-991.
- Rodrigues MC, Voltarelli J, Sanberg PR, Allickson JG, Kuzmin-Nichols N, Garbuzova-Davis S, Borlongan CV (2012) Recent progress in cell therapy for basal ganglia disorders with emphasis on menstrual blood transplantation in stroke. *Neurosci Biobehav Rev* 36:177-190.

- Russo I, Barlati S, Bosetti F (2011) Effects of neuroinflammation on the regenerative capacity of brain stem cells. *J Neurochem* 116:947-956.
- Seabra ML, Bignotto M, Pinto LR Jr, Tufik S (2000) Randomized, double-blind clinical trial, controlled with placebo, of the toxicology of chronic melatonin treatment. *J Pineal Res* 29:193-200.
- Shahmoradian RA, Ali AAA, Wu CC, Chiang YH, Chen KY (2019) Enhanced homing of mesenchymal stem cells overexpressing fibroblast growth factor 21 to injury site in a mouse model of traumatic brain injury. *Int J Mol Sci* doi:10.3390/ijms20112624.
- Shlosberg D, Benifla M, Kaufer D, Friedman A (2010) Blood-brain barrier breakdown as a therapeutic target in traumatic brain injury. *Nat Rev Neurol* 6:393-403.
- Skop NB, Calderon F, Cho CH, Gandhi CD, Levison SW (2016) Optimizing a multifunctional microsphere scaffold to improve neural precursor cell transplantation for traumatic brain injury repair. *J Tissue Eng Regen Med* 10:E419-432.
- Steinberg GK, Kondziolka D, Wechsler LR, Lunsford LD, Coburn ML, Billigen JB, Kim AS, Johnson JN, Bates D, King B, Case C, McGrogan M, Yankee EW, Schwartz NE (2016) Clinical outcomes of transplanted modified bone marrow-derived mesenchymal stem cells in stroke: a phase 1/2a Study. *Stroke* 47:1817-1824.
- Stroemer P, Patel S, Hope A, Oliveira C, Pollock K, Sinden J (2009) The neural stem cell line CTX0E03 promotes behavioral recovery and endogenous neurogenesis after experimental stroke in a dose-dependent fashion. *Neurorehabil Neural Repair* 23:895-909.
- Tajiri N, Acosta SA, Shahaduzzaman M, Ishikawa H, Shinozuka K, Pabon M, Hernandez-Ontiveros D, Kim DW, Metcalf C, Staples M, Dailey T, Vasconcellos J, Franyuti G, Gould L, Patel N, Cooper D, Kaneko Y, Borlongan CV, Bickford PC (2014a) Intravenous transplants of human adipose-derived stem cell protect the brain from traumatic brain injury-induced neurodegeneration and motor and cognitive impairments: cell graft biodistribution and soluble factors in young and aged rats. *J Neurosci* 34:313-326.
- Tajiri N, Duncan K, Antoine A, Pabon M, Acosta SA, de la Pena I, Hernandez-Ontiveros DG, Shinozuka K, Ishikawa H, Kaneko Y, Yankee E, McGrogan M, Case C, Borlongan CV (2014b) Stem cell-paved bio-bridge facilitates neural repair in traumatic brain injury. *Front Syst Neurosci* 8:116.
- Tajiri N, Duncan K, Borlongan MC, Pabon M, Acosta S, de la Pena I, Hernandez-Ontiveros D, Lozano D, Aguirre D, Reyes S, Sanberg PR, Eve DJ, Borlongan CV, Kaneko Y (2014c) Adult stem cell transplantation: is gender a factor in stemness? *Int J Mol Sci* 15:15225-15243.
- Tajiri N, Kaneko Y, Shinozuka K, Ishikawa H, Yankee E, McGrogan M, Case C, Borlongan CV (2013) Stem cell recruitment of newly formed host cells via a successful seduction? Filling the gap between neurogenic niche and injured brain site. *PLoS One* 8:e74857.
- Takemoto M, Liao JK (2001) Pleiotropic effects of 3-hydroxy-3-methylglutaryl coenzyme a reductase inhibitors. *Arterioscler Thromb Vasc Biol* 21:1712-1719.
- Tan K, Wang X, Zhang J, Zhuang Z, Dong T (2018) [Effect of chitosan porous scaffolds combined with bone marrow mesenchymal stem cells in repair of neurological deficit after traumatic brain injury in rats]. *Zhongguo Xiu Fu Chong Jian Wai Ke Za Zhi* 32:745-752.
- Tapia-Perez J, Sanchez-Aguilar M, Torres-Corzo JG, Gordillo-Moscote A, Martinez-Perez P, Madeville P, de la Cruz-Mendoza E, Chali-Williams J (2008) Effect of rosuvastatin on amnesia and disorientation after traumatic brain injury (NCT003229758). *J Neurotrauma* 25:1011-1017.
- Tunc Ata M, Turgut G, Akbulut M, Kocyigit A, Karabulut A, Senol H, Turgut S (2016) Effect of erythropoietin and stem cells on traumatic brain injury. *World Neurosurg* 89:355-361.
- Uekawa K, Hasegawa Y, Ma M, Nakagawa T, Katayama T, Sueta D, Toyama K, Kataoka K, Koibuchi N, Kawano T, Kuratsu J, Kim-Mitsuyama S (2014) Rosuvastatin ameliorates early brain injury after subarachnoid hemorrhage via suppression of superoxide formation and nuclear factor-kappa B activation in rats. *J Stroke Cerebrovasc Dis* 23:1429-1439.
- Valadi H, Ekström K, Bossios A, Sjöstrand M, Lee JJ, Lötvall JO (2007) Exosome-mediated transfer of mRNAs and microRNAs is a novel mechanism of genetic exchange between cells. *Nat Cell Biol* 9:654-659.
- Vasandan AB, Jahnvi S, Shashank C, Prasad P, Kumar A, Prasanna SJ (2016) Human mesenchymal stem cells program macrophage plasticity by altering their metabolic status via a PGE(2)-dependent mechanism. *Sci Rep* 6:38308.
- Wang H, Lynch JR, Song P, Yang HJ, Yates RB, Mace B, Warner DS, Guyton JR, Laskowitz DT (2007) Simvastatin and atorvastatin improve behavioral outcome, reduce hippocampal degeneration, and improve cerebral blood flow after experimental traumatic brain injury. *Exp Neurol* 206:59-69.
- Wang KW, Chen HJ, Lu K, Liliang PC, Liang CL, Tsai YD, Cho CL (2014) Simvastatin attenuates the cerebral vascular endothelial inflammatory response in a rat traumatic brain injury. *Ann Clin Lab Sci* 44:145-150.
- Wang Z, Wu L, You W, Ji C, Chen G (2013) Melatonin alleviates secondary brain damage and neurobehavioral dysfunction after experimental subarachnoid hemorrhage: possible involvement of TLR4-mediated inflammatory pathway. *J Pineal Res* 55:399-408.
- Watanabe J, Shetty AK, Hattiangady B, Kim DK, Foraker JE, Nishida H, Prockop DJ (2013) Administration of TSG-6 improves memory after traumatic brain injury in mice. *Neurobiol Dis* 59:86-99.
- Wei ZZ, Lee JH, Zhang Y, Zhang YB, Deveau TC, Gu X, Winter MM, Wei L, Yu SP (2016) Intracranial transplantation of hypoxia-preconditioned iPSC-derived neural progenitor cells alleviates neuropsychiatric defects after traumatic brain injury in juvenile rats. *Cell Transplant* 25:797-809.
- Werner C, Engelhard K (2007) Pathophysiology of traumatic brain injury. *Br J Anaesth* 99:4-9.
- Wu H, Mahmood A, Lu D, Jiang H, Xiong Y, Zhou D, Chopp M (2010) Attenuation of astrogliosis and modulation of endothelial growth factor receptor in lipid rafts by simvastatin after traumatic brain injury. *J Neurosurg* 113:591-597.
- Yu P, Li S, Zhang Z, Wen X, Quan W, Tian Q, Gao C, Su W, Zhang J, Jiang R (2017) Progesterone-mediated angiogenic activity of endothelial progenitor cell and angiogenesis in traumatic brain injury rats were antagonized by progesterone receptor antagonist. *Cell Prolif* doi: 10.1111/cpr.12362.
- Zhang R, Liu Y, Yan K, Chen L, Chen XR, Li P, Chen FF, Jiang XD (2013) Anti-inflammatory and immunomodulatory mechanisms of mesenchymal stem cell transplantation in experimental traumatic brain injury. *J Neuroinflammation* 10:106.
- Zhao J, Moore AN, Clifton GL, Dash PK (2005) Sulforaphane enhances aquaporin-4 expression and decreases cerebral edema following traumatic brain injury. *J Neurosci Res* 82:499-506.
- Zibara K, Ballout N, Mondello S, Karnib N, Ramadan N, Omais S, Nabouh A, Caliz D, Clavijo A, Hu Z, Ghanem N, Gajavelli S, Kobeissy F (2019) Combination of drug and stem cells neurotherapy: Potential interventions in neurotrauma and traumatic brain injury. *Neuropharmacology* 145:177-198.
- Ziebell JM, Morganti-Kossmann MC (2010) Involvement of pro- and anti-inflammatory cytokines and chemokines in the pathophysiology of traumatic brain injury. *Neurotherapeutics* 7:22-30.

P-Reviewer: Kirby ED; C-Editors: Zhao M, Li JY; T-Editor: Jia Y