ABSTRACT

A cytokine release syndrome (CRS), associated with elevated circulating levels of several cytokines including interleukin (IL)-6 and interferon (IFN)-γ, might be seen in some infectious insults, for instance in severe acute respiratory syndrome (SARS) induced by Coronavirus (Cov)-2, as well as following administration of natural and bispecific antibodies and, more recently, following adoptive T-cell therapies for cancer. Normally, inflammatory conditions activate the innate and adaptive immune systems, which results in the release of cytokines, responsible for the phagocytosis of apoptotic vesicles and resolution of inflammation. Pro-inflammatory cytokines such as IL-1β, tumor necrosis factor alpha (TNFα) and, especially in chronic inflammatory diseases, autoimmune diseases, cancer and cytokine storms, IL-6 play crucial roles in inflammation. In some instances, however, this release gets out of hand, and features of overzealous immune responses (macrophage activation syndromes) might occur, leading to cytokine release syndromes (CRS) with inflammatory signs such as fever, fatigue, nausea, and sometimes secondary organ dysfunction or multi-organ failure. Apart from specific vaccines and maybe the anti-viral remdesivir and/or dexamethasone for treatment of CRS, there are no convincing disease-modifying interventions. So far, though, non-antiviral and immune-targeted interventions, also affecting non-target cells, were found associated with many side affects. A more targeted or focused approach is thus needed. Pending the site of the CRS-inducing insult, CRSs may occur systemic or compartmental. Recently, preclinical research yielded a beneficial anti-inflammatory effect of fresh naive bone marrow-derived stem cells (bm-SCs) in the treatment of various compartmental CRSs in the immune-privileged central nervous system (CNS). Therefore, it is argued that bm-SCs might also play a disease-modifying role in the systemic CRS. Bm-SCs have the advantage of targeting only the cells of interest as they are very selective in their actions. In addition, they actively move to the site of inflammation.
respiratory syndrome (MERS), for instance, viruses bind to alveolar epithelial cells and then activate the innate and adaptive immune systems, which results in the release of many cytokines, including (IFN)γ and IL-6. Normally, this process is adequately controlled by the anti-inflammatory mechanisms of the body. However, in some instances, this mechanism gets out of hand, and features of overzealous immune responses (CRS) might occur. This phenomenon might explain why some people have a severe reaction to for instance coronaviruses while others only experience mild symptoms. The reason why younger people are less involved, though, is still an enigma. CRS may be best formulated as an infectious or sterile-induced production of circulating cytokines beyond a normal response, to a specific condition, leading to inflammatory signs with fever, severe fatigue, nausea and in some cases even secondary organ dysfunction or multi-organ failure.

The precise reason for this excessive inflammatory response is not entirely understood but may be caused by new or specific highly pathogenic conditions, able to initiate a macrophage activation syndrome (MAS). Cytokines play an important role in normal immune responses but when they overshoot their target, due to a MAS, they can be extremely harmful. In those conditions, the expression of a healthy immune system results in the pathological release of multiple inflammatory mediators (cytokines, oxygen free radicals, and coagulation factors). These include both pro-inflammatory mediators such as TNF (an inflammatory cytokine produced by macrophages/monocytes during acute inflammation, responsible for a diverse range of signaling events within cells, leading to necrosis or apoptosis), IL-1 (a group of 11 cytokines that plays a central role in the regulation of immune and inflammatory responses to infections or sterile insults, stimulating phagocytosis and programmed cell death) and IL-6 (a pleiotropic cytokine, secreted by T cells and macrophages o activate immune responses during infections or after trauma), as well as anti-inflammatory mediators such as IL-1 receptor antagonists and the cytokine synthesis inhibitory factor IL-10 (a cytokine with potent anti-inflammatory properties that plays a central role in limiting host immune response to pathogens, thereby preventing damage to the host and maintaining normal tissue homeostasis). So, it is not the illiciting pathogenic insult itself but rather the overreactive immune response to this condition, which induces a cytokine release syndrome.

A CRS can be systemic or compartmental, such as in immune-privileged areas such as the eyes, joints and the central nervous system. Following traumatic brain injury (TBI) and spinal cord injury (SCI), there are no disease-modifying interventions, but preclinical research suggests a beneficial (anti-inflammatory) effect of autologous naive stem cells to treat this overactive inflammatory condition.

Stem cells are not only acting on local concentrations of circulating pro- and anti-inflammatory proteins, they are rather selective in their actions, thus sparing non-target tissue cells. The risk of stem cells for eventual adverse events in humans can be comprehensively reduced when applying fresh non-substantial manipulated autologous stem cells.

**Compartmental Overreactive Immune Responses (TBI, SCI)**

Pro-inflammatory cytokines play an essential role in maintenance of normal and repair functions, for instance in the central nervous system compartment, after TBI and SCI. Massive and uncontrolled release of these cytokines such as IL-1β and TNF-α, and especially IL-6, though, may result in a great deal of collateral damage around the site of the injury. After an initial acute injury-induced necrosis, a profound and uncontrolled, local secondary inflammatory reaction will develop, characterized by the abundant release of several cytokines, mainly IL-6, with pro- and anti-inflammatory functions. In terms of molecular pathogenesis, traumatic injuries of brain and spinal cord share a negative contribution of non-neuronal
cells (immune cells, glial cells) expressing and activating the inflammatory process as a significant commonality. Microglia, the resident immune cells in the CNS, comprise an entire spectrum of phenotypes that span the range from deleterious to regulatory to remodeling effects. Gene expression analyses led to the identification of homeostatic state 1 microglia and environmental-dependent, disease-associated, reactive state 2 and state 3 microglia. State 2 microglia are the classical acute necrosis-activated cytotoxic microglia (interferon response microglia: IRMs), affecting neuronal survival and secreting pro-inflammatory TNF, IL-1β and IL-12, and nitric oxide. State 3 microglia are the alternatively activated, cytoprotective microglia (activated response microglia: ARM) that express genes involved in innate immune response, thus supporting an anti-inflammatory response, and preventing classical microglial activation. ARMs secrete the anti-inflammatory cytokines IL-4 (stimulating activated B-cell and T-cell proliferation, and the differentiation of B cells into plasma cells), the immunoregulatory IL-13 (regulates the function of human B cells and monocytes), and TGF-β. Once the inciting inflammatory event has been adequately resolved, a lower production of pro-inflammatory cytokines will convert IRMs into ARMs which facilitate phagocytosis of cell debris, promote tissue repair, and support cell survival. However, when the pathogenic stimulus for some reason cannot be adequately cleared, a persistent IRM response will follow. As a consequence, a progressive unintended injury might follow, as is the case in many neurodegenerative disorders. This condition is visualized in the perpetual cell death-inducing process in figure 1.

In the central nervous system, pending the function of the blood-brain-barrier, CRS may lead to highly elevated levels of IL-6 (1,400–3,550,000 pg/ml), normally marginally detectable (10–230 pg ml) as compared to corresponding serum levels (0–11,000 pg/ml). Significant higher levels were also seen for other cytokines, including IL-1α, IL-1β, IL-8, IL-10, and IL-12. Some of these cytokines can have both beneficial as well as harmful effects on the brain. The timing of the rise in cytokines may be crucial.

Acute or chronic pathological cytokine release is established in some neurologic conditions, spinal/cerebral trauma, amyotrophic lateral sclerosis (ALS), Alzheimer’s and Parkinson’s disease as well as multiple sclerosis, and explained as an underlying cause of the non-infectious pathology. Here, the body is not capable to adequately handle these inflammatory processes, which then become chronic, and progressively affect neurological functioning. Ultimately, in animal models for amyotrophic lateral sclerosis (ALS) and spinal cord injury (SCI) this process could be aborted, and pro-inflammatory cytokine levels lowered by an intervention with Bm-SCs. In spinal cord injury, acute injury-induced apoptosis is followed by a necrosis-induced secondary inflammatory response, deteriorating the direct trauma-induced neurological functioning. Activated microglia and macrophages play an important role in this secondary inflammation. Cytotoxic IRMs (see figure 1), here were found responsible for the enhanced production of pro-inflammatory cytokines. The intrathecal application of a standardized human Bm-SC product (fresh not-manipulated autologous bone marrow containing high concentrations of viable hematopoietic and mesenchymal stem cells obtained by centrifugation and positive depletion of erythrocytes, monocytes and lymphocytes, suspended in Ringers lactate, and reduced in volume for intrathecal application: Neuro-Cells, manufactured by Neuroplast BV, The Netherlands) did reduce the levels of pro-inflammatory cytokines as well as the neuropathologic signs in the central nervous system. Most importantly, the resolution of the inflammatory process did improve functional outcome (which was not the case in steroid treated control groups). Indeed, the benefit of Bm-SCs over anti-inflammatory drugs, including steroids, was shown in both spinal cord injury and ALS-like animals. The exact mechanisms are still unknown, but the fact that stem cells are not only decision-making cells, moving to and acting upon local concentrations of circulating pro- and anti-inflammatory proteins, is a likely explanation for the beneficial roles of Bm-SCs. The risk for eventual adverse events of stem cells, when limited to the exclusive use of non-substantial manipulated autologous stem cells, is negligible.

Another compartment are the synovial spaces of joints in which an overwhelming immune reaction can exacerbate an ongoing arthritis. The onset is still not clear, but in the affected joints, cytokines such as TNF and IL-1 here play an important role in apoptosis and loss of cartilage cells. The levels of TNF and IL-1 are increased in patients with rheumatoid arthritis in serum as well as in the synovial fluid.

**Systemic Overreactive Immune Responses (SARS-CoV-2)**

Lately we have been faced by the devastating effects of COVID-19 infections in which fatalities were linked to a dysregulated host immune response to a severe acute respiratory syndrome (SARS-CoV-2). In the most serious affected patients, SARS-induced MAS displayed significantly increased plasma levels of several cytokines, chemokines, and inflammatory markers (such as C-reactive protein, procalcitonin, and ferritin), levels correlating with serious systemic signs of inflammation, and sometimes organ dysfunction or multi-organ failure with fatal outcome. Severe COVID-19 and SARS infections are notable for severe T-cell depletion. In COVID-19, there is unique suppression of interferon signaling by infected respiratory tract cells with intact cytokine signaling. A decreased naïve T-cell
response likely contributes to an excessive inflammatory response and increases the odds of a cytokine storm. Serum cytokine levels that were found elevated in patients with CoV-19–associated hyperreactive immune responses included IL-1β, IL-6, TNF, IFN-γ and IFN-γ-induced IP-10, macrophage inflammatory protein (MIP) 1α and 1β, and vascular endothelial growth factor (VEGF). Especially IL-6 is hypothesized as one of the main mediators. The pooled mean serum interleukin-6 concentration in these patients was about 100 times higher (3110.5 pg/mL) when compared to Covid patients not suffering from this overreactive immune reaction (36.7 pg/mL). In addition to the activated immune cells and elevated systemic cytokine levels, other clinical and laboratory abnormalities were found to predict worsening outcomes in CoV-19, such as an elevated C-reactive protein, hypoalbuminemia, and renal dysfunction. Using a logistic regression model to analyze the factors influencing the severity of the disease, Wang and colleagues identified 3 of 19 predictive variables as significant (white blood cells, IL-10, and IL-6 levels) using the new Chinese National Health Commission Coronavirus Pneumonia Classification, and 2 of 19 predictive variables as significant (lower CD3 percentages and higher IL-6 levels) using the Sequential Organ Failure Assessment (SOFA) score. The multivariable regression analysis results showed only IL-6 levels as indicators related to the severity of Covid patients in both classifications.

Treatment of Overreactive Immune Responses

Systemic cytokine release syndromes

Pending the severity of cytokine release syndromes, supportive treatment may be necessary to protect organs and prevent further damage. Symptomatic treatment includes intravenous fluids, oxygen or ventilator support, medicines to support heart function, blood product transfusions, medicine or dialysis for the kidneys. Cantini et al. studied the reports of antiviral therapies in Covid-19 patients, and concluded that only Remdesivir, not Lopinavir/ritonavir, Favipiravir, Atazanavir, Interferon α2b, Ribavirin or Umifenovir reduced both some time to recovery and mortality. IL-6 expression, activated by IL-1β and TNFα, plays an important role in CRS. Although anti-IL-6 antibodies should lower IL-6 expression and thus dampen the downstream IL-6 signaling pathways, as in the treatment of rheumatoid arthritis, Castleman’s disease and CAR-T-induced CRS, IL-6R blockers Tocilizumab, Sarilumab, and Siltuximab have been ineffective. Cantini et al. also evaluated the results of other non-antiviral and immune targeted agents. They concluded that both anakinra and dexamethasone reduced mortality, that results from chloroquine and hydroxychloroquine use were inconclusive; and that Baricitinib resulted in a significant reduction of intensive care unit admission and deaths. The promising results of anakinra, though, await confirmation, and corticosteroids, which reduce inflammation without targeting a specific cytokine, as in the treatment of TBI and SCIs were found to increase mortality risk and to delay recovery with a longer hospital stay.

Compartmental cytokine release syndromes

In the central nervous system, microglia, the resident immune cells in the CNS, comprise an entire spectrum of environment-adapted phenotypes that span the range from deleterious to regulatory to remodeling effects. Although microglial activation provides a first defense against acute injury or infection, in these cases chronic or excessive activation is considered highly detrimental. Indeed, in TBI and SCI, direct trauma-induced necrosis is followed by a secondary inflammatory reaction, which incites an excessive IRM response that will eliminate even the surviving cells in the immediate vicinity (the penumbra). In brain and spinal cord injuries, unfortunately, treatment with anti-inflammatory drugs have produced mixed results at best and can produce serious adverse events, unless the result of insufficient dosage or bioavailability, delayed time points of delivery, and/or the taking-over of the eventually inhibited pathway by other single or downstream pathways. As the blood-brain-barrier is usually very tight, the systemic access of these drugs to the brain might also be hampered.

A more recent approach of treating such disorders with intrathecal transplants of fresh bone marrow-derived stromal cells (bm-SCs) seemed more promising. Intrathecal application of naïve bm-SCs in SCI-rats within 24 h after the lesioning inhibited the conversion of state-2 (IRM) into state-3 (ARM) microglia, decreased neuronal apoptosis as well as the activators of IL-6 expression (TNF and IL-1β) in serum and spinal tissue when compared to the vehicle-treated animals. Intrathecal application of naïve bm-SCs in SCI-rats within 24 h after the lesioning inhibited the conversion of state-2 (IRM) into state-3 (ARM) microglia, decreased neuronal apoptosis as well as the activators of IL-6 expression (TNF and IL-1β) in serum and spinal tissue when compared to the vehicle-treated animals. These bm-SCs were found to dampen the downstream IL-6 signaling pathways, with improved clinical outcomes in experimental SCI-rats with evident features of a cytokine-driven hyperimmune reaction.

In case of another compartmental CRS, rheumatoid arthritis, one of the interventional targets is to lower TNF concentration by specific TNF-α inhibitors such as infliximab or certolizumab. Despite this specific approach, there is still a non-responder rate of more than...
 Which make this kind of therapy less effective. The experience with stem cells in rheumatoid arthritis is limited but, in general, their (immune-modulating) effects have led to a considerable improvement in the management of RA patients.

Conclusion

Cytokine release/secretion syndromes can be the endpoint of many different triggers ranging from immunotherapies to infectious diseases. It is not surprising that there is, yet no general therapy established, since the causes are so numerous and different in kind. Preclinical research suggests a beneficial (anti-inflammatory) effect of naïve bm-derived stem cells in the treatment of this overactive inflammatory condition in the central nervous system compartment. However, it is even in well controlled experimental models not clear, whether these anti-inflammatory effects are mediated by the resident cells if the stem cells have quickly passed their effects onto those cells. The inherent benefit of bm-derived stem cells over a drug modulating a single inflammatory effector pathway has been shown in head-to-head comparisons. An alternative explanation could be the activation of pathways higher up in the inflammatory cascades. Bm-SCs are decision-making cells that coordinate their operations with their immediate environment. Their cell-to-cell communication here may play a major role and future research must elucidate the content of stem cell-secreted EVs in modulating inflammatory micro-environment(s) and the systemic effect thereof. If the effects of bm-SCs are directly related to the way of administration and the differentiation state of the stem cells, naïve bm-derived stem cells may be superior. Therefore, subsequent studies on the systemic intravenous application of bm-derived stem cells in systemic CRS, and in compartmental CRS diseases via a direct injection (intercompartmental like intrathecal or injections in the joint) need to be done to address this question.

In acute SCI animal models, the intrathecal application of bm-SCs was found to reduce the initial increase of IL-1β, TNF and IL-6 levels, and thus to prevent for the secondary inflammation-induced neurological deficits. This effect might be exerted by the stem cells immediately after application or, indirectly, after passing their effects to resident cells.

Acknowledgment

The research is done by Neuroplast bv and Neuroplast bv receives an innovation loan of the Ministry of economic affairs of the Netherlands. No additional funding is involved.

Conflict of Interest Statement

Johannes de Munter is CEO of Neuroplast bv and has a staff position at the department of Neuroscience and Mental Health at the University Maastricht, The Netherlands. Kimberly de Hoo is employed by Neuroplast bv as clinical associate.

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