Comparative analysis of movements after experimental spinal cord injury treated with amniotic membrane: Pilot study

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Abstract

Purpose: This study evaluated the action of human amniotic membrane (HAM) in rats after induced spinal cord injury (SCI).

Methods: Twenty-five Wistar rats were divided into five groups (n=5): health (healthy animals underwent to surgical procedure simulation), control (C-injury simulation), injury (I - spinal cord injury) and HAM (SCI + immediate application of HAM). We analyzed the effect of application of HAM fragment on SCI using gait speed analysis (kinematics) and the sciatic functional index (SFI) to the animal gait. Both tests were performed in triplicate at 5 and 12 days after the surgery.

Results: Movement analyses demonstrated that the groups which received the HAM application showed an increase in average speed (after 12 days) and in the sciatic functional index (at both 5 and 12 days) when compared to other groups.

Conclusion: The gait speed kinematics and the functional gait (SFI) analyses indicated that the application of HAM in induced SCI in rats led to the recovery of motion and functional gait, which suggests a positive action of this biomaterial in SCI sequels.

Introduction

Spinal cord injury (SCI) is a reduction or loss of motor function and/or sensory trauma of neuronal elements within the spinal canal where the neurological level of involvement may be complete or incomplete [1]. Complete SC injuries are one of the most common causes of loss of movement and sensitivity at and below the level of the lesion. SCI can lead to spinal shock followed by loss all sensory-motor functions, which can be caused by tumors, infections, vascular disorders, degenerative, and malformations [7,8]. SCI sequels may include expenses with high medical treatment for physical and pathological complications, as urinary tract infection, spasticity, chills and fever, decubitus ulcers and autonomic dysreflexia, as well as social complications as job loss and the permanent need for emotional support systems and medical assistance [6,9].

This scenario may lead to the research of new therapeutic approaches to repair SCI using stem cell-based therapies, such as transplantation of stem cells / neural progenitor. Previous studies have reported functional recovery after stem cell transplantation for spinal cord injury in rodents and nonhuman primates [10]. In this context, application of cells or fragments obtained from human Amniotic Membrane (HAM) has gained importance due to their ability to reduce inflammation and scarring and to serve as a substrate for cell proliferation and differentiation [11-14].

Therefore, considering HAM properties could promote tissue repair as a source of stem cell, we have hypothesized that their use after...
SCI could represent a potential new treatment for SCI sequel’s in motor function. This study aimed to analyze the functional gait of rats with induced SCI comparing the action of HAM applied immediately after the injury.

Methods

After approval by the Research Ethics Committee of University of Vale of Paraíba (CAAE: 16381213.2.0000.5503; CEAU nºA15), 20 adult male rats (Rattus norvegicus, Albinus, Wistar, 60 days, 230 ± 20 g) were divided into four groups:

1. H (health – healthy animals who suffered no surgical nor injury procedures, n=5);
2. C (control-simulation procedures surgical, n=5);
3. I (Injury - spinal cord injured animals, n=5);
4. HAM (SCI + immediate application of HAM, n=5) and

The animals stayed in individual cages with a 12:12 light-dark cycle, at controlled temperature (20 ± 2 ºC), feed with diet and water ad libitum. The HAM was obtained from human placentas collected at the Hospital and Maternity of Santa Casa in São José dos Campos, São Paulo, Brazil. All donors signed a Free, Prior, and Informed Consent, as well as met the inclusion criteria and all laboratory procedures for processing HAM occurred in sterile conditions inside a laminar flow cabinet in accordance with the processing protocol described by Sant’Anna, et al. [13]. First, we manually peeled the HAM from the chorion and extensively washed it with saline solution containing 100 U/ml penicillin, 100 mg/ml streptomycin, and 2.5 ug/ml of amphotericin B. Then, we cut the HAM into pieces of 2 cm x 2 cm and we stored them separately at room temperature in 50 ml tubes containing DMEM (Dulbecco’s Modified Eagle’s medium) without addition of serum and red phenol, in a sterile condition until the application into the animal [13,15].

Surgical procedures

Spinal cord injury was induced by direct trauma with a mini guillotine in groups I and HAM following the protocol described by Paula, et al. [16]. We submitted Group C rats only to surgical procedures and simulation of HAM application, without any kind of injury. For the surgical procedures, the animals received intramuscularly 10% ketamine (anesthetic) associated with xylazine (muscle relaxant) at doses 0.01 ml/kg and 0.005 ml/kg, respectively. We performed a longitudinal incision (2.5 cm) in the dorsal region, between T9 and T10, to expose and visualize the spinal cord. Hereafter, an impactor system (Figure 1) was used to induce a SCI by dropping the guillotine weight from 25 mm free fall, causing a trauma injury by direct contusion. Fifteen seconds were waited and then it was removed [16].

After removal the guillotine weight, we applied a 2.5 cm fragment of HAM in the group of HAM animals. It is emphasized that the MAH fragment was applied with the mesenchymal side in contact with the SCI area, completely covering this region. The edges of HAM fragment were adhered to the surrounding bone tissue with one drop of methacrylate. Tissues are repositioned and sutured with 3-0 polyamide monofilament thread (nylon) suture (Figure 2).

All animals received antibiotic (enrofloxacin, 5 mg/kg subcutaneously once per day for 4 days) and analgesic therapy (tramadol hydrochloride, 10 mg/kg, intraperitoneally, every 12 hours for 5 days) and housed at heated cage (25ºC) for 45 min to control the postoperative hypothermia.

Figure 1. Induction of SCI between T9-T10, by direct trauma, with the use of the mini guillotine.

Figure 2. Surgical procedure. (A) T9-T10 area with spinal cord injury; (B) HAM edges attached to the bone tissue around the region of interest; (C) suturing the tissues.

Evaluation of motor function

Two analyses of movements were performed in triplicate on groups H, C, I and HAM at 5 and 12 days after the surgery. For gait speed analysis (kinematics), we used a box made of acrylic (46 cm longX 8 cm wideX30 cm high) open at the surface. The box was placed in the angle with two cinematic cameras (Optitrack, Flex model: V100 R2) supported on tripods, and positioned laterally to the collection area. For three-dimensional image reconstruction, a reflective marker, adhered to the femur of the mouse’s left hind leg, remained visible simultaneously for both cameras. The positioning of the box and the camera was marked on the floor to standardize data collection in relation to cameras and their calibration.

The software Motive (OptiTrack®) captured the movement of each animal and exported the data in CVS model for analysis in Origin Pro 8.5® software. Each animal performed the test three times (triplicate) and the average gait speed (cm/s) was obtained by calculating the linear coefficient held in the analysis software and expressed by mean and standard deviation.

Also, for the sciatic functional index (SFI) run (collected in triplicate), we used the same box (Figure 3). We coupled a dark box at the end of the trial box where the animals headed for refuge. The trial box was suspended, resting on two aluminum stands. A mobile device...
was positioned below, with the camera directed to the center of the acrylic box, enabling the capture of images from the lower plane.

The video images were exported to the computer to capture the “Frame Grabs” (frames) of the phases of the gait and subsequent measurement [17] using Microsoft® Windows Movie Maker®, Version 5.1 to capture three frames of each animal (triplicate). The frames were stored and exported to the program Image J (Version 1.48, National Institute of Health) to calculate the parameters length of the footprint of the experimental paw (PL), distance between the first and fifth toes of the experimental paw (TS), and distance between the second and the fourth toe of the experimental paw (IT). The SFI values below zero show functional impairment and near -100 configured losses of total function [18].

One-way ANOVA statistical test could be applied after all tests were run in triplicate generating mean and standard deviation values (T-test; p ≤ 0.05).

Results

Table 1 and Figure 4 show the motor assessments of the H, C, I and HAM groups, at 5 and 12 days after surgeries represented as mean ± standard deviation, at significance level p ≤ 0.05.

Table 1 shows that 5 days after induced SCI, the injured group (I) had a significant reduced mean velocity when compared to health (p = 0.0349), to control (0.0480) group. HAM group presented slower movement when compared to health (p = 0.0449) and control (p = 0.0094) at the same experimental time. In the intragroup analyses, both injury group (p < 0.001) and HAM group (p = 0.386) had mean gait speed increased, while Control group showed no significant change of mean gait speed. It is important to observe that regarding 12th post-operative day, there was no significant difference in the intergroup.

Figure 4 presents the data relating Sciatic Functional Index (SFI) to functional gait. The experimental groups I (n=5) and HAM-0 (n=5) showed significant motor impairment of the hind paw when compared to H (n=5) and C (n=5) groups both 5 and 12 days after surgery (p < 0.05). Post-operative groups did not differ among themselves at 5 days. However, the movement impairment of HAM group was significantly lower when compared to groups I (p = 0.0093) and at the twelfth post-operative day. In the intragroup analysis at the same experimental times, only the HAM-0 group showed significant recovery of motor function at the twelfth post-operative day (p = 0.0335).

Discussion

This is an innovative study evaluating the effect of human amniotic membrane (HAM) fragments applied immediately (HAM group) after the induction of spinal cord injuries in rats. The application of a HAM fragment aimed to preserve the tissue microenvironment of this biomaterial, since it contains several growth factors and undifferentiated mesenchymal cells, on the assumption that this would facilitate the repair of nervous tissue lesion [11-14]. Further, the application of HAM in fragments requires less manipulation than the application of stem cells, keeping the growth factors present in it microenvironment could stimulate the differentiation of mesenchymal cells as response to stimuli received from the injured tissue [13]. The number and viability of cells in the human amniotic membrane fragments, when stored for 24 h, suffer no significant change when compared to the fresh membrane [12].
The application of HAM immediately after the induction of SCI aimed to evaluate possible differences in the action of the biomaterial applied immediately after the injury. From our results, we verified that the application of HAM was effective to improve gait speed and functional gait after induced SCI in rats.

SCI can cause loss of neurological functions that affect all levels at and below the injury site, which means that the movements and sensations can be totally or partially reduced below the lesion area [19]. The inflammatory process following trauma can start a secondary lesion characterized by edema, hemorrhage, progressive tissue loss, neuronal loss, and glial cell dysfunctions [16]. The recovery process in humans is limited and not very functional, because the tissue environment that hinders the regenerative process of the axon [19,20].

SCI is also associated with motor and sensory symptoms that lead to changes in posture and gait. The recovery of the gait is one of the main goals of rehabilitation treatment from the point of view of patients and therapists [21].

Researchers have proposed several therapeutic strategies to repair secondary injury via spinal cord regeneration and neuronal growth [22]. In this context, SCI treatment protocols have evolved a plurality of options, such as anti-inflammatory non-steroidal drugs; physiotherapy; Low-Level Laser Therapy (LLLT); ultrasound; corticosteroid injections; shock wave therapy; surgery; biomaterials, growth factor and stem cell application [14,16,20,23].

Up to this moment the reports in the literature pertaining to neurological tissue repair involving the amniotic membrane are still scarce. Although, there are consistent reports of its effectiveness when applied, in the clinical ophthalmology field [24]; as a fragment, about ischemic heart of rats [12]; on the experimental biliary fibrosis of rats, reducing the degeneration fibrotic [13]; about inducted lesion in mouse Achilles tendon [14], among others.

A pilot study from Oliveira, et al. which investigated by histopathological qualitative analysis the application of the HAM fragment in rats SCI, could observe that the HAM visibly reduced the formation of cavitation with cystic aspect, determining less destruction of the spinal cord tissue, with consequent preservation of healthy tissue, when compared to the animal who did not receive HAM after SCI. They also observed the reduced presence of chronic inflammatory infiltrate in the animals of the HAM group, which was a positive response factor, considering that the presence of these cells after the 28-day period would be classified as a chronic response, and could lead to tissue degeneration and necrosis, as well as the formation of large cavities in the nerve tissue, with consequent loss of coordinated functions.

The kinematic analysis demonstrated that the HAM did not differ from health and control groups. However, the HAM group showed a significant increase in average speed on the 12th day after the induced SCI, which was confirmed by improved function observed by the SFI (Table 1). The gait evolution of groups with the HAM suggests a promising positive action of this biomaterial in the repair of nervous tissue.

Other studies showed that HAM has other properties that can aid in tissue repairs, such as anti-adhesive effect, wound protection, adhesion and migration of basal epithelial cells, reduction of fibrosis, prevention of apoptosis, and restoration of the epithelial phenotype. HAM also contains various growth factors, which stimulate re-epithelialization [24,25].

According to Nicodemo, et al. [14] the use of HAM in wound healing in Achilles tendon of rats proved to be very efficient, because there was no increased induction in the inflammatory response, and HAM presented excellent biocompatibility with the injured tendon tissue. Histological analysis of histological sections showed that HAM fragment was incorporated into the area of tendon injury after 7 days, and the number of mature fibroblasts and collagen fibers significantly increased compared to the injured group at 14th day.

Bottai, et al. [26] analyzed the effects of amniotic fluid cells in the third trimester of pregnancy applied in rats with spinal cord injury. These cells grew in different culture media and when transplanted to animals, some of the tested samples were able to induce significant improvement in motor recovery Basso Mouse Scale (BMS). The improved motor functional recovery shows a correlation to the greater preservation of myelin in the ventral horn of white matter and increased vascularization in the peri-wound area.

According to Brown, et al. [27] HAM is a biologically active tissue and has been used to repair myelomeningocele (MMC) in human fetuses. In their study, sheep fetuses underwent surgical creation of MMC at the gestational age of 75 days followed by intrathecal repair at the gestational age of 100 days. The lambs recovered after applying a fragment of autologous HAM, followed by skin closure. The autopsy and histopathology of the spinal cords showed an increase in motor neurons and decreased scarring of wounds in the skin.

Our results are consistent with the properties described for HAM in several studies. HAM acts in the tissue repair process probably as a “basement membrane transplant”, as a substrate for cell proliferation that occurs after spinal cord injury. Articles previously mentioned showed the action antibacterial, anti-inflammatory and inhibitory of proteinase of HAM, as well as the presence of various growth factors such as EGF, TGF-, HGF, which stimulate and promote the functional recovery [13,14,24,25,28].

The high availability and ease of obtaining HAM makes this biomaterial an excellent option for the treatment of SCI. However, clinical application depends on the validation of new controlled experimental studies.

Conclusion

The results of this study suggested that HAM could repair surgically induced spinal cord injuries in rats, and lead to better performance in the recovery of average speed gait and sciatic functional index.

References


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