Transplantation of Neural Progenitor Cells within Hyaluronic Acid Hydrogel in Traumatic Brain Injury in Experiment

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The aim of the study was to evaluate the possibilities of transplantation of autologous neural progenitor cells from C57BL/6 mouse nasal olfactory lamina propria in the hydrogel based on low-, medium-, and high-molecular weight hyaluronic acid during a reconstructive operation after an open traumatic brain injury (TBI) in experiment.

Materials and Methods. A reconstructive therapy was performed on the model of an open traumatic brain injury using the autologous neural progenitor cells from C57BL/6 mouse nasal olfactory lamina propria in a hydrogel based on high, medium, and low molecular hyaluronic acid 7 days after injury. Short-term and long-term impairments of neurological functions and memory due to TBI were assessed using a series of behavioral and cognitive tests (modified neurological severity score (mNSS)), open field test, novel object recognition test and passive avoidance behavior). High-field magnetic resonance tomography was used for visualization of a lesion cavity.

Results. Implantation of autologous neural progenitor cells in the hydrogel based on high-molecular weight hyaluronic acid in contrast to low- and medium-molecular weight had a protective effect reducing neurological deficit and restoring motor functions, short-term and long-term memory in the posttraumatic period. The effect of hydrogel molecular weight on the morphological characteristics of the injury cavity was less effective during the autologous neural progenitor cells transplantation after TBI.

Conclusion. Transplantation of autologous neural progenitor cells from C57BL/6 mouse nasal olfactory lamina propria in the hydrogel based on high-molecular weight hyaluronic acid into the injury focus after open brain trauma promotes functional recovery of reflex and cognitive behavior of animals in the posttraumatic period.

Key words: autologous neural progenitor cells; hyaluronic acid hydrogel; traumatic brain injury; TBI; functional recovery.

Treatment of the consequences of traumatic brain injury (TBI) and a number of other neurodegenerative processes is one of the most complex and socially significant problems of medicine, as in the majority of cases it leads to disability and decline in cognitive capacity [1, 2]. Absence of significant success of drug therapy [3], limited ability of neurogenesis in adults organism [4] cause the study of fundamental regeneration mechanisms of the injured brain areas and developing principally new methods of TBI treatment. The development of adequate matrix-carrier for transplanted cells, which would create a specific microenvironment with prolonged recovery process of neural networks and tissue structure regeneration is one of the most promising methods of the treatment of TBI [5–7].

At present, the works in the field of neurotransplantation are mainly focused on the use of hydrogel systems of various composition with progenitor cells. Firstly, the pore structure of the hydrogel allows cells and nutrients to penetrate into the matrix and metabolic products to dispose in the volume of the body [8]. Second, the hydrogel systems are optimal in terms of...
mechanical compatibility with the tissues of the brain and spinal cord tissue. The compatibility of the mechanical characteristics is determining during the differentiation of progenitor cells [9] and influences overall success of implantation [10]. Third, hydrogels are promising drug compounds in the injury site.

Extracellular matrix of the brain consists of the molecules synthesized and secreted by neurons and glial cells, which form stable aggregates in intercellular space in various combinations [11]. The most pronounced and studied clusters of extracellular matrix molecules in the CNS are perineuronal nets [12]. They are rich in hyaluronic acid (HA), chondroitin sulfate proteoglycans and link proteins. HA being a framework for the extracellular matrix take part in regulation of cellular differentiation, migration, proliferation, and angiogenesis. Perisynaptic matrix comprising HA can create barriers to the diffusion of synaptic molecules in the membrane and thus promoting compartmentalization synaptic mechanism of signal transmission.

In addition, HA can form hydrogels, which not only contain a large amount of water that promotes the viability of cells, but also have biophysical, biochemical properties, similar to those of the body tissues. Moreover, hyaluronic hydrogels are biologically compatible. However, their use for neurotransplantation is not yet sufficiently developed.

The aim of the investigation was to assess the possibilities of transplantation of autologous neural progenitor cells of C57BL/6 mouse nasal olfactory lamina propria in the hydrogel based on low-, medium-, and high-molecular weight hyaluronic acid hydrogel in the reconstructive surgery of open traumatic brain injury in experiment.

Materials and Methods. Adult male C57BL/6 mice (10–12 weeks of age, 20–22 g) were purchased from the Department of Experimental Animals, Pushchino, Russia, and maintained under standard housing conditions (temperature: 18–24°C; relative humidity: 45%; light and dark cycle: 12:12 h). Food and water were provided ad libitum. All animals were treated according to the protocols evaluated and approved by the Guide for the Care and Use of Laboratory Animals (National Research Council, 2011), Russian Federation standard 33044-2014 “Principles of Good Laboratory Practice” and the Ethical Committee of the Nizhny Novgorod State Medical Academy. The number of mice used and their suffering were minimized.

Hyaluronic acid-based hydrogel. The hydrogel which was used for neurotransplantation was a sterile 1% water solution of sodium hyaluronate (Scientific Production Enterprise (SPE) Tula Industry, Ltd, Russia) with different molecular weight of the active substance: low-molecular weight form (55.0 kDa); medium-molecular weight form (500.0 kDa); high-molecular weight form (1,250.0 kDa). Sodium chloride (8.5 mg) and phosphate buffered saline (PBS) (320 μg/ml) were used as auxiliary substances.

Culture preparation. Neural progenitor cells were obtained from the olfactory mucosa of C57BL/6 mice according to the previously developed protocol [13].

Weight-drop traumatic brain injury model. In this study, we used the Feeney weight-drop method with some modifications [14] by dropping the weight with a blunt surface which provides acceleration with a minimum of local action at the point of application of the traumatic force. This model produces a complete clinical picture of focal injury including brain contusion, and allows exploration of traumatic and local changes that are accompanied by secondary death of neurons in distant parts of the brain, which are sensitive to trauma, such as hippocampus, dentate gyrus and thalamus, and also evaluation of the movement disorders and cognitive impairment.

Spontaneously breathing mice were anesthetized with 1.5% isoflurane in oxygen and placed in a stereotaxic frame adapted for mice (Narishige, Japan) on a 37.8°C heating pad. For inducing focal brain injury, the animals were placed on a non-flexible steel platform in order to minimize energy dissipation, to reduce the risk of jaw fractures and achieve the horizontal location of the calvarium to the tip of the weight. All animals underwent craniotomy under aseptic conditions using a surgical microscope. The scalp was shaved and cleansed with chlorhexidine and povidone-iodine wash. The scalp and periosteum were opened, and left craniotomy (4 mm in diameter, 2 mm posterior to stereotaxic bregma, and 2 mm lateral to the midline) was performed with a fine drill, leaving the dura intact. A weight-drop device was placed over the dura. A sterile polypropylene cylindrical tube was manufactured with a 4-mm opening at one end. Then, a 4 g weight stainless steel cylinder was dropped from 80 cm height through a guiding tube onto the burr hole to create a contusion brain injury model. TBI results in an ipsilateral injury with cortical contusion, hemorrhage, dura rupture that is typical for severe head trauma. After injury, the scalp incision was closed tightly with polypropylene suture and treated with an antiseptic solution.

Experimental groups. Adult male C57BL/6 mice (n=60) were randomly assigned to six groups. The first group (n=10) included intact animals. The second group (n=10) comprised sham-operated animals. Mice of control third group (n=10) were injected PBS into the lesion cavity. The fourth group (n=10) was with implantation of progenitor cells in the high-molecular weight hyaluronic acid hydrogel (HA “D”); the fifth group — implantation of progenitor cells in the medium-molecular weight hyaluronic acid hydrogel (HA “E”) (n=10); the sixth group — implantation of progenitor cells in the low-molecular weight hyaluronic acid hydrogel (HA “K”) (n=10).

Injury day was considered as day 0. Mice in the “HA hydrogel with cells” groups were subjected to TBI, and 7 days later, a hyaluronic acid-based hydrogel with engrafted progenitor cells (30 μl) was directly
Experimental investigations on magnetic resonance imaging (MRI) analysis and testing of neurological and cognitive functions of mice.

The modified neurological severity score. This scale includes the assessment of 10 clinical parameters characterizing motor functions, ability to balance, and some behavioral reactions of animals (alertness, searching behavior). One point is given for failure of the task and no point for succeeding. A maximal neurologic severity score of 10 points indicates severe neurological dysfunction, with failure at all tasks [15–18].

A day before TBI modeling, neurological status was evaluated in all groups of animals to assess the initial level and then on 1–10, 15, 20, 30 days post injury.

Open field test. To assess cortical function, mice were subjected to behavioral testing using of the open field task on day 4 prior to injury (baseline) and on days 10 and 30 post injury. Locomotor and exploratory behavior was analyzed. At the beginning of the test, individual animals were placed in the center of the open field box (LE800S; PanLab/Harvard Apparatus Spain; Stoelting Co., USA) and allowed to freely explore it for 5 min. Traveled distance, mean speed and the number of rearing as well as duration of anxiety-like behavior such as grooming and freezing acts were automatically recorded with a video camera (SONY SSC-G118; Japan). The data acquisition and analysis were controlled with the SMART v.3.0.03 software (Panlab/Harvard Apparatus Spain; Stoelting Co., USA).

Passive avoidance test. The passive avoidance task was used to assess simple non-spatial learning ability on day 10 post injury [19]. The passive avoidance device comprised of two adjoining compartments, one illuminated and one darkened, divided by a guillotine door. The dimensions of each compartment were 46×27×25 cm. The floor of the compartments consisted of steel rods capable of delivering a slight electric foot shock (50 Hz, 1.5 mA for 10 ms) to the animal (Shuttle Box LE895; PanLab/Harvard Apparatus Spain; Stoelting Co., USA). The test began by placing the mouse in the lighted compartment. The feeding of electric foot shock, the position of the guillotine door between the compartments and the latency for crossing to the dark compartment were recorded with the Shutavoid v.1.8.03 software (PanLab/Harvard Apparatus Spain; Stoelting Co., USA). The test was over when the animal entered the dark compartment or did not do it during 3 min. Cognitive functions were studied during retention test after 24 h (on day 10) and on day 30 of posttraumatic period.

Novel object recognition test. To evaluate impairments of nonspatial hippocampus-mediated short-term memory, a novel object recognition test was performed. The procedure consisted of three sessions — habituation, training, and retention [20, 21]. On days 23 and 24 after TBI, the animals were placed in the center of the square arena (45×45×40 cm) (LE802S; PanLab/Harvard Apparatus Spain; Stoelting Co., USA) for 10 min for habituation and stress reduction — the animals explored the empty arena. On day 24 of the posttraumatic period, in the first trial, two identical objects were placed in the right and left corners of the box. Mice were placed into the box for 10 min, where they explored the two objects, and the exploratory activity was recorded. During the retention session 24 h after the training session, the animals were placed into the same box, but one of the familiar objects used during the training was replaced with a novel object. The objects were different color but similar in size. The animals were then allowed to explore freely for 10 min, and the time spent in exploring each object was recorded.

In the course of the experiment, the cumulative time spent by the mouse exploring each of the familiar and novel objects was recorded. Exploration of an object was defined as follows: directing the nose to the object at a distance of 2 cm and touching it with the nose. Animal behavior during training, and retention was considered as explorative if total exploring time to the objects lasted not less than 10 s. Registration of single explorative acts of animals was carried out using video camera (SONY SSC-G118; Japan). The data acquisition and analysis were controlled with the SMART v.3.0.03 software (Panlab/Harvard Apparatus Spain; Stoelting Co., USA). Memory function was operationally defined by the discrimination ratio (DIR) for the novel object (discrimination ratio, DIR) by the formula: DIR=[(novel object exploration time – familiar object exploration time)/total exploration time]×100 [17].

Magnetic resonance imaging. Imaging of the mice brain was carried out on the high field MRI machine Agilent Technologies DD2-400 9.4 T (USA) (400 MHz) with a volume coil M2M (H.). Monitoring of the physiological parameters of animals (temperature, respiration and ECG) during imaging was performed on a SA Instruments company (USA) equipment using PC-SAM program. During imaging the animals were under isoflurane anesthesia. Animals were heated by 37° C warm air.

MGEMS (multi gradient echo multi slice) pulse sequence with the following parameters: repetition time is 1,300 ms, echo time is 8 ms, echo number — 8, the amount of savings — 4, number of slices — 15, the thickness of slice was 1 mm, the field of vision — 20×20 mm², matrix size — 128×128 pixels was used for T2*-weighted imaging. The total duration of the sequence was 8 min 32 s.

Statistical analysis. Results were expressed as mean ± standard error of the mean (SEM). All data...
were evaluated for normal distribution by Shapiro-Wilk normality test (W-criterion). All data were analyzed using Statistica 8.0 software (StatSoft, USA). Nonparametric analyses were performed using a Wilcoxon signed-rank test for repeated measures of outcomes of the TBI. Statistical comparisons between behavioral data of treatments on each evaluated time-point were performed using a Mann–Whitney U test. A p value of <0.05 was considered significant.

**Results and Discussion**

**Assessment of neurological status of the animals in the posttraumatic period.** The functional assessment for mice behavior showed the sustained fault of neurological functions on day 1 after TBI (Figure 1). The modified neurological severity score in the groups with brain injury was significantly higher compared with initial level as well as intact and sham-operated groups of animals (p<0.05).

Asymmetrical behavior was characteristic of all animals due to limb paresis on the contralateral side to the injury cavity, impairments of motor functions, inability to perform reactions connected with maintaining the balance and movement coordination, which is the evidence of the damage of the motor cortex, midbrain, and pons Varolii. Impairments of reflexes associated with the midbrain and cortex damage were also observed.

The neurological deficit was reduced by day 10 of the posttraumatic period in all groups of animals with transplantation of autologous progenitor cells in HA hydrogel into the injury cavity on day 7 after TBI. However, no statistically significant differences compared to the control group were detected (p>0.05).

The tendency to the decrease of the neurological deficit in the group with implantation of the HA “D” hydrogel with engrafted neural progenitor cells into the injury cavity was observed on day 20 of the posttraumatic period (p>0.05).

The neurological status of mice in this group significantly improved compared to the control group (PBS) (p<0.05) in contrast from the groups with implantation into the lesion site on the day 7 of the posttraumatic period of neural cells in the HA “E” and HA “K” on the day 30 after injury. It should be noted that throughout the observed posttraumatic period the neurological functions in animals with TBI were not completely restored. Paresis on the contralateral side to injury and partial impairments of reflexes (searching behavior) were observed. The level of neurological deficit of mice was significantly increased compared to the intact and sham-operated groups of animals.

**Open field test.** A preliminary testing of the animals (before TBI) did not reveal any differences between the groups in the main behavior characteristics. TBI resulted in the impairment of the animal behavior structure. The data obtained in the open field test showed statistically significant the motor and explorative activity decreasing in comparison with the initial level also as the values in the intact and sham-operated groups of the animals (p<0.05). There was no restoration of exploratory activity in any of the mice groups with TBI.

The activity speed recovery occurred in the group of animals with implantation of neural progenitor cells based on HA “D” hydrogel into the injury cavity on day 10 of the posttraumatic period. This factor was significantly higher compared to the control group (p<0.05), but did not differ either from the values of the intact and sham-operated group or from other experimental groups with TBI (p>0.05) (Figure 2).

On day 30 of the posttraumatic period, the speed of motor activity in this group was statistically higher compared to the control (PBS) mice group (p<0.05). It signified the restoration of locomotor activity and functions associated with maintaining the balance.

The activity speed was not restored in the groups of mice with implantation of HA “E” and HA “K” hydrogels engrafted with neural progenitor cells and was not different from the value of the control (PBS) group.
**Experimental Investigations**

Figure 2. The speed of locomotor activity of C57BL/6 mice in the open field test in the posttraumatic period

* $p<0.05$ significant differences between pre- and post-injured values, Wilcoxon test; * $p<0.05$ values in injured brain mice differ significantly from the values of the intact mice, Mann–Whitney U test; ** $p<0.05$ significant difference compared to sham-operated group, Mann–Whitney U test; * $p<0.05$ significant difference compared with the control (PBS) group; Mann–Whitney U test; * $p<0.05$ significant difference compared with the group of high-molecular weight HA + cells, Mann–Whitney U test; n=10 per group

![Graph showing speed of locomotor activity](image)

**Passive avoidance behavior.**

The results of behavioral tests revealed the impairment of learning processes of animals in the control (PBS) group as a result of TBI. The entry latency to the dark compartment in the control (PBS) group of mice significantly increased compared to intact and sham-operated group ($p<0.05$) during the acquisition test. It was caused by the impairment of the motor functions of the CNS of animals owing to the TBI.

On day 10, the entry latency into the dark compartment of the shuttle box in the group with neural progenitor cells in HA “D” hydrogel into the injury cavity significantly increased in the retention test compared with acquisition session as well as in intact and sham-operated groups ($p<0.05$). Moreover, the latency in this group was statistically higher compared to the control (PBS) group ($p<0.05$) (Figure 3).

![Graph showing latency](image)

In the groups with the implantation hydrogel of medium- and low-molecular weight HA engrafted with the progenitor cells into injury cavity 1 week after TBI, the time spent in the illuminated part of chamber at the retention test compared with acquisition session did not differ (p>0.05).

Moreover, the activity speed was significantly lower in compared with the group with implantation of the neural cells in the HA “D” hydrogel (p<0.05).
(p>0.05). At the same time, the latency in the group with the implantation of cells in the hydrogel of low-molecular weight HA was significantly reduced compared to the group of high-molecular weight HA hydrogel engrafted with the cells (p<0.05).

Long-term memory functions were assessed by ability of mice to retain passive avoidance behavior on day 30 of the posttraumatic period. The entry latency at the retention test significantly differed from acquisition session (p<0.05) in the group of animals with implantation of progenitor cells based on HA “D” hydrogel into the injury cavity. Moreover it was significantly exceeded the time of the control (PBS) group (p<0.05).

In the groups with transplantation of neural cells in the medium- and low-molecular weight HA hydrogel as well as in the control group the time spent in the lite compartment was significantly less than the time in the intact group (p<0.05) and in the group with implantation of progenitor cells in the hydrogel of the high-molecular weight HA (p<0.05) on day 30 of the posttraumatic period.

Thus, implantation of HA “D” hydrogel engrafted with progenitor cells into the injury cavity 7 days after TBI modeling caused an optimizing effect on the ability of the animals to acquire conditioned reflexes on day 10 and to update traces of long-term memory in the remote period.

**Novel object recognition test.** The data obtained showed that TBI results in the significant impairment of hippocampus-dependent working memory. Discrimination ratio in the control (PBS) group was significantly decreased compared to the intact and the sham-operated groups (p<0.05) (Figure 4).

The time spent to explore the novel object was statistically higher than the time spent for the familiar object in the group of mice with progenitor cells in HA “D” hydrogel implantation on day 7 following the injury. Discrimination ratio in this animal group did not differ from that in the intact and sham-operated groups of mice over the whole posttraumatic period. At the same time, the DIR value in the group with progenitor cells in HA “D” hydrogel implantation was increased compared to control (PBS) group (p<0.05) that indicates the recovery of CA1–CA3 field hippocampus-dependent short-term recognition memory functions.

In the groups of mice with the neural cells in the hydrogel of medium- and low-molecular weight HA implantation after 7 days after TBI, the discrimination ratio, on the contrary, was not differ from the control (PBS) group (p>0.05). It was significantly decreased in compare to the value in the group of high-molecular weight HA hydrogel engrafted with progenitor cells implantation as well as in the intact and sham-operated groups (p<0.05).

**Magnetic resonance imaging data.** MRI data obtained 30 days after injury showed that in all groups with implantation of neural progenitor cells with HA hydrogel into the injury cavity the volume of the lesion area did not significantly differ from the control (PBS) group (p<0.05) (Figure 5).

The injury area was detected as a defect on the surface of the hemisphere on the magnetic resonance images on day 30 of the posttraumatic period in the group with implantation of the progenitor cells in the HA hydrogel “D”, while the lesion area had a rounded form in the sensorimotor cortex in the group with progenitor cells transplanted with the HA hydrogel “E”. Cystic lesion of a round form with irregular margins filled with fluid in the area of the sensorimotor cortex was visualized on the magnetic resonance images of the group of mice with implantation of HA “K” hydrogel engrafted with progenitor cells.

Thus, TBI results in neurologic, motor and cognitive behavior of animals impairments. Implantation of neural progenitor cells in the hydrogel of the high-molecular weight HA into the injury cavity 7 days after TBI had the protective effect by restoring the synaptic plasticity (cortex, hippocampus) underlying the motor functions, learning and memory processes, in contrast to the transplantation of the medium- and low-molecular weight HA hydrogel with engrafted neural progenitor cells.
A facilitative effect of transplantation of neural progenitor cells in HA hydrogel on the ability of the animals to acquire passive avoidance reflex in acute posttraumatic period and to update traces of short and long-term memory in the remote period has been established. The effect of implantation of the autologous neural progenitor cells in HA hydrogel on the restoration of morphological parameters of the injured brain tissue was less evident. However, an attempt to regenerate the brain tissue structure was observed in the group with implantation of progenitor cells in the hydrogel of the high-molecular weight HA.

**Conclusion.** Transplantation of autologous neural progenitor cells from C57BL/6 mouse nasal olfactory lamina propria in the hydrogel based on high-molecular weight hyaluronic acid into the injury cavity after open...
brain trauma promotes functional recovery of reflex and cognitive behavior of animals in the postraumatic period.

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