Autologous Adipose-Derived Stromal Cells and PRF Extract for Spontaneous Intervertebral Disc Disease in Canine

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Abstract
The administration of autologous adipose-derived stromal cells (ADSC) and platelet-rich fibrin (PRF) extract by intra-venous injections was investigated as a treatment for canine intervertebral disc disease (IVDD). IVDD is a disease that can cause several symptoms in domestic dogs, ranging from pain to partial or complete paralysis. Five dogs suffered from IVDD that caused hind-legs paralysis with deep pain perception, were treated with autologous adipose-derived stromal cells, vascular fraction, and PRF extract. ADSC were expected to ameliorate symptoms of IVDD due to its anti-inflammatory and cell regenerative action. Five sequential intravenous injections applied in one-week interval for five weeks improved condition of treated dogs judged by patients gait score and exercise pattern.

Keywords: Intervertebral disc disease; Adipose stromal stem cells; Platelet-rich fibrin; Stromal vascular fraction


Introduction
Intervertebral disc disease (IVDD) is a condition when the spinal intervertebral disc either bulged or herniated into the spinal cord space. IVDD can present with number of symptoms in domestic animals, ranging from a mild pain to partial or complete paralysis. The symptoms of IVDD can be induced by acute trauma or from other causes of disc herniation. IVDD occurs more commonly in some specific dog breeds, including the Dachshund, Beagle, Basset Hound, Poodle and Shih Tzu, but can occur in any breed and in dogs of any age or gender. There are three Hansen types of IVDD. In the Hansen type I, intervertebral discs become fragile due to calcification of their outer layer. Any forceful impact such as post-jumping can cause one or more discs to burst and the inner material can herniate and compress the spinal cord. Type 1 IVDD commonly affects the neck spine region in smaller dog breeds. With the Hansen type II herniation, discs become fibrous and harder over a long period of time and eventually break down, bulge out, and compress the spinal cord. Type III Hansen is a subtype of Hansen type I that the intervertebral disc degeneration and herniation characterized by extension of disc material “like a carpet over several vertebrae” [1].

A complete neurologic examination may suggest the location of the spine cord compression that can be specified by X-ray and in some cases by MRI (magnetic resonance imaging) or CT (computed tomography) examination. The path physiology of IVDD involves not only the initial mechanical damage, but also the secondary cascades of inflammation, ischemia free-radical formation, and cytotoxicity resulting in neuron death by necrosis or apoptosis. Regenerative medicine therapies based on biomaterials and stem cells application have been tested with promising results [2]. This clinical study reports the results of application of autologous adipose-derived stromal cells and platelet-rich fibrin (PRF) extract to dogs suffering from IVDD (Figure 1).
Figure 1: Dog was diagnosed IVDD by CT examination. The dog exhibited complete paralysis and loss of the deep pain perception. IVDD locations are indicated by red-marked circles.

Case Presentation

In this study, all dogs with IVDD have symptoms that are confirmed by the CT scan and neurological examination as a spinal cord function disturbance (Table 1). All dogs lost the panicculi, urinary bladder continence, proprio reflex and deep pain perception (Table 2). They could not support their own body weight. The IVDD scale used for quantifying spinal cord abnormalities in these individuals indicated presence of the IVDD grade 5. The five dogs that completed the experimental treatment program with ADSC and platelet-rich fibrin (PRF) extract exhibited an excellent gait performance. The paralysis ameliorated in four dogs and they could walk easily and with a good gait pattern after eight weeks (Table 3). The dog 1 did not improve well in eight weeks, but it showed good panicculi, urinary bladder continence and proprio reflex. The dog walked after 16 weeks after ADSC and platelet-rich fibrin (PRF) extract administration.

Table 1: Dogs with IVDD have symptoms that are identified by CT examinations.

<table>
<thead>
<tr>
<th>Examination Finding</th>
<th>Dog 1 (Yang-Mei)</th>
<th>Dog 2 (Pi-Dan)</th>
<th>Dog 3 (Ji-Li)</th>
<th>Dog 4 (Don-Don)</th>
<th>Dog 5 (Shau-Hai)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CT scan Lesion</td>
<td>T3-T4 L1-2-3-4</td>
<td>L1-L2</td>
<td>C4-C5 T12-T13</td>
<td>T5-T6 L1-L2</td>
<td>T13-L1</td>
</tr>
</tbody>
</table>
**Table 2:** Clinical findings of submitted dogs before stem cells therapy.

<table>
<thead>
<tr>
<th>Clinical Finding</th>
<th>Dog 1 (Yang-mei)</th>
<th>Dog 2 (Pi-dan)</th>
<th>Dog 3 (Ji-li)</th>
<th>Dog 4 (Don-don)</th>
<th>Dog 5 (Shau-hai)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Panicculi reflex</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
</tr>
<tr>
<td>Urinary Bladder continence</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
</tr>
<tr>
<td>Proprio. reflex</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
</tr>
<tr>
<td>Weight bearing</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
</tr>
<tr>
<td>Deep pain perception</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
<td>(-)</td>
</tr>
<tr>
<td>IVDD Grade</td>
<td>(V)</td>
<td>(V)</td>
<td>(V)</td>
<td>(V)</td>
<td>(V)</td>
</tr>
<tr>
<td>Age</td>
<td>8y/o</td>
<td>4y/o</td>
<td>5y/o</td>
<td>9y/o</td>
<td>11y/o</td>
</tr>
<tr>
<td>Breed</td>
<td>Dachshund</td>
<td>Poodle</td>
<td>Dachshund</td>
<td>Dachshund</td>
<td>Mixed</td>
</tr>
</tbody>
</table>

**Figure 2:** The procedure for ADSC isolation and purification: Fat tissue was collected under general anaesthesia (A, B). Tissue samples were kept at 4-7°C and were washed three times with sterile saline to remove contaminating blood cells and tissue debris (D). The washed tissues were digested with an equal volume of 0.1 % collagenase at 37 °C for 30min (E). The digested tissues then were inactivated with 10 % fetal bovine serum (FBS) (F). The tissues were washed five times with saline and centrifuged at 300g for 5min. The pallet was collected and resuspended (G). Cells in the pallet was observed under microscope (H) [21].

NSF: no specific finding.
Table 3: Clinical improvement of submitted dogs after stem cells therapy.

<table>
<thead>
<tr>
<th></th>
<th>Dog 1 (Yang-mei)</th>
<th>Dog 2 (Pi-dan)</th>
<th>Dog 3 (Ji-ii)</th>
<th>Dog 4 (Don-don)</th>
<th>Dog 5 (Shau-hai)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Improve time</td>
<td>4wks 8wks</td>
<td>4wks 8wks</td>
<td>4wks 8wks</td>
<td>4wks 8wks</td>
<td>4wks 8wks</td>
</tr>
<tr>
<td>Paniculii reflex</td>
<td>(-) (+)</td>
<td>(+) (+)</td>
<td>(+) (+)</td>
<td>(+) (+)</td>
<td>(+) (+)</td>
</tr>
<tr>
<td>Urinary Bladder continence</td>
<td>(-) (+)</td>
<td>(+) (+)</td>
<td>(+) (+)</td>
<td>(+) (+)</td>
<td>(+) (+)</td>
</tr>
<tr>
<td>Proprio. reflex</td>
<td>(-) (+)</td>
<td>(+) (+)</td>
<td>(-) (+)</td>
<td>(+) (+)</td>
<td>(+) (+)</td>
</tr>
<tr>
<td>Weight bearing</td>
<td>(-) (-)</td>
<td>(+) (+)</td>
<td>(-) (+)</td>
<td>(+) (+)</td>
<td>(+) (+)</td>
</tr>
<tr>
<td>Walk</td>
<td>(-) (-)</td>
<td>(+) (+)</td>
<td>(-) (+)</td>
<td>(+) (+)</td>
<td>(+) (+)</td>
</tr>
</tbody>
</table>

Discussion

ADSC is from the adipose stromal vascular fraction (SVF) (Figure 2), which is comprised of the mononuclear cells derived from adipose tissue. This term is used to describe the mitotically active source of adipocyte precursors. SVF as a source of stem cells was first described by De Ugate et al. [3], who identified MSC-like cells in SVF that could be induced to differentiate into adipogenic, chondrogenic, myogenic, and oestrogenic lineages [3]. Subsequent to the initial description, the same group reported after in vitro expansion, the SVF derived cells had surface marker expression similar to bone marrow derived MSC, comprising CD29, CD44, CD71, CD90, CD105/SH2, and SH3 and lacking CD31, CD34, and CD45 expression.

Autologous platelet-rich fibrin (PRF) extract was isolated from the dog’s blood as described in materials and methods based on the paper [8].

Figure 3: Autologous platelet rich fibrin (PRF) extract was isolated from the dog’s blood as described in materials and methods based on the paper [8].

(A) The PRF clots were retrieved from the tubes and the red blood cell gel was detached and discarded. Four PRF clots were transferred into sterile tubes. The tubes were put on a shaker and agitated gently. After 5 minutes, the tubes were vortexed to form a PRF membrane, and the volume of releasate was measured and returned into the tube. The tubes were further agitated gently.

(B) The extract was mixed with ADSC and injected by intravenously.

In this study, we showed that the administration of ADSC and PRF extract could improve the damage of spinal cord in the intervertebral disc disease (IVDD) in dogs. The severity of IVDD was classified into five grades according to clinical signs and symptoms. Grade 1 includes spinal hyperesthesia, Grade 2 for ambulatory paraparesis, ataxia, proprioceptive deficits, Grade 3 for non-ambulatory paraparesis, Grade 4 for paraplegia with nociception preserved, and Grade 5 for paraplegia with loss of nociception [11].

The stem cells application has demonstrated the promising result in spinal cord injury [12]. Several reports demonstrated effectiveness of bone marrow stromal cells implantation in acute spinal cord injury models, including monkey [13], human [14] and dogs [15,16]. In our study, the five dogs suffering from IVDD Grade V and experimentally treated with autologous cells derived from adipose tissue and PRF extract significantly improved their clinical
condition.

According to Dr. Kim’s research, the full recovery time in decompression surgery for IVDD was about 62.25 days. The decompression surgery plus ADSC local graft was 59.7days for recovery to grade 1 [19]. In this study, we combined ADSC and PRF extract to treat grade 5 IVDD patients. The average recovery time was about 48 days. It thus appears as a better recovery than that from decompression surgery.

The spinal cord undergoes inflammation response by primary hurt [18,19]. The reddish, swelling, heat, hemorrhage and function loss were induced in damaged area. Another physic or mechanical procedures could induce more cell death and hemorrhage. The more serious injuries were induced to form the glia scar in spinal cord [20,21]. ADSC and PRF extract were supposed to decrease the inflammation cytokines such as IL 6 and TNF-α and limited the inflammation cascade pathway [23,24].

This study, though limited in size, successfully tested a novel treatment procedure for dogs suffering from IVDD in canine. The performance of the clinical dog patients in this preliminary study encourages us to go toward a brilliant direction.

**Consent of Patients**

The experimental protocol was approved by the Deng Chuan veterinary hospital, Kaoshiung, Taiwan and all participants provided informed consent before the study.

**Acknowledgement**

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**Conflict of Interest Statement**

The authors have declared no conflict of interest.

**References**
