The Microscopic Pictures of Degenerative Intervertebral Disc of Different Diseases with Additional Safranin-O Stain

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Abstract

Background: The lumbar spine and cervical spine are vulnerable with degeneration, which is related to idiosyncratic problem, aging or improper stress of the spine. Biochemical and structural changes of disc, narrowing of intervertebral disc space, acute or chronic disc protrusion related to disc degeneration with/without neural symptoms are well documented. We are interested in the morphologic changes of the excised herniated or degenerative discs from the spinal surgeries for herniated intervertebral disc (HIVD), spondylolisthesis or spinal stenosis.

Methods: The specimens are collected under the permission of institutional review board (IRB), and are divided into three main groups, including 9 specimens (9 cases) of HIVD, 12 specimens (10 cases) of spondylolisthesis, and 10 specimens (10 cases) of spinal stenosis. Five disc specimens of 3 cadavers (3 from lumbar L4-5 disc and 2 from thoracic T9-10 disc) with no mentioned disc disease are obtained for control group. Histochemical stains of hematoxylin and eosin (H and E) stain and safranin-O-fast green stain are performed to evaluate the content of glycosaminoglycans and chondrocytic proliferation. The photos of disc lesions are taken with consistent LED light of Nikon n Coolscope II to analyze the discoloration within the H and E and safranin-O stained sections. Kruskal-Wallis test (one-way ANOVA) was performed for significance check.

Results: HIVD cases have younger age (p=0.036). The excised disc specimens of HIVD group show mild loss of glycosaminoglycans over nucleus pulposus, and even less eosinophilic change in the nucleus pulposus. The discs of spondylolisthesis and spinal stenosis groups may display uneven loss of glycosaminoglycans with occasionally massive loss, and more eosinophilic change than that of HIVD group (p=0.014). Clonal proliferation of chondrocytes over the areas other than endplate and outer rim of annulus fibrosus is a consistent phenomenon in all three groups of degenerative discs, and displaying lesser degree of chondrocytic proliferation among the HIVD cases without significant difference.

Conclusion: We notice that focal eosinophilic change or even necrosis of extracellular matrix is probable an important pathologic picture of degenerative disc and clonal proliferation of chondrocytes is present in all 3 groups of degenerative intervertebral discs. The chondrocytic proliferation in the disc fibrocartilage might be similar to the chondrocytic proliferation within the articular cartilage of grade 3-4/6 knee osteoarthritis.

Keywords: Chondrocytic proliferation; Colorimetry analysis; Degenerative intervertebral disc; Safranin-O-fast green stain

Introduction

Intervertebral disc is belonged to the kind of fibrocartilage, and composed of nucleus pulposus, i.e., a gelatin-like center, and annulus fibrosus, a thick capsule of air-bag-like structure. The end plate, a dense fibrocartilage interface, functions as a buffer between vertebral body and intervertebral disc, and may influence the nutrition support from verbebral capillaries [1]. The lumbar spine and cervical spine are two regions vulnerable with degenerative change due to idiosyncratic tendency, aging or improper stress. HIVD is most frequent cause of sciatica [2]. Spinal stenosis and spondylolisthesis are other common causes of low back pain or even lumbosacral radiculopathy with functional disturbance of lower limbs. Biochemical and structural changes of disc, narrowing of intervertebral disc space, acute or chronic disc protrusion with/without neural symptoms related to disc degeneration are well documented, and also called spondylosis. Osteoarthritits of knee joint with loss of articular cartilage and narrowing joint space on X-ray film is the most well-known degenerative joint disease, can be graded by Safranin-O stain. Therefore, we are interested in safranin-O stain for comparing the glycosaminoglycans (GAG) content in addition to the conventional microscopic findings on the excised disc specimens.
### Materials and Methods

The specimens are collected under the permission of institutional review board (IRB: B09901002-1), and are divided into three main groups, including 9 specimens (9 cases) of intervertebral disc herniation (HIVD), 12 specimens (10 cases) of spondylolisthesis, and 10 specimens (10 cases) of spinal stenosis (Table 1). If more than one type of spinal diseases is diagnosed, the case will be distributed into the group of severer one. Three lumbar disc (L4-5) specimens and 2 thoracic disc (T9-10) specimens from 3 autopsy cases with no mentioned disc disease are enrolled for control group. One is a victim of Wilms’ tumor with mild cachexia died at the age of 34 years old. Another one is a victim of breast cancer with minimal cachexia expired at the age of 42. The rest case expired at the age of 81 due to chronic disease without past history of intervertebral disc disorder (Figure 1).

| Sex | Age | HIVD |
|-----|-----|------|
| M   | 57  | L4/5 HIVD, Rt |
| M   | 31  | L4/5 HIVD, Rt, recurrent |
| M   | 48  | L4/5 HIVD, Lt |
| F   | 59  | L3-5 Spondylolisthesis with disc herniation |
| M   | 79  | L4/5 recurrent disc herniation |
| F   | 57  | L3/4 disc herniation |
| M   | 45  | L4/5 HIVD |
| M   | 49  | L5/S1 Spondylolisthesis with disc herniation |
| M   | 51  | C6 Vertebral body fracture with C6/7 HIVD |
| M   | 64  | L4/5 disc herniation, L4/5 Spondylolisthesis |
| F   | 68  | L4/5 Spondylolisthesis |
| F   | 56  | L4/5 Spondylolisthesis with canal stenosis |
| F   | 63  | L4-5 Spondylolisthesis |
| M   | 55  | L4/5 Spondylolisthesis with spinal stenosis |
| F   | 63  | X2: L2/3 and L3/4 degenerative kyphoscoliosis |
| F   | 56  | L4-5 Spondylolisthesis, L3-5 spinal stenosis |
| F   | 61  | L4-5 Spondylolisthesis with L4/5 spinal stenosis |
| F   | 70  | X2: L2-4 and L5/S1 degenerative scoliosis |
| M   | 64  | S/P L4/5 PDF with L3/4 adjacent degeneration |
| F   | 78  | L2/3 scoliosis with canal stenosis |
| M   | 82  | L4/5 spinal stenosis |
| F   | 69  | L4/5 spinal stenosis |
| M   | 57  | S/P L3-5/S1 posterior fusion with L2/3 stenosis |
| F   | 72  | L3-S1 Spondylolisthesis with stenosis |
| M   | 51  | L4/5 segmental instability with stenosis |
| F   | 62  | L4 compression fracture with L4/5 spinal stenosis |
| F   | 62  | L4/5 spinal stenosis |
| F   | 87  | L3-5 spinal stenosis |
| M   | 66  | L3-5/S1 spinal stenosis |

Table 1: The genders, ages and preoperative diagnoses (made by medical images and clinical findings) of 3 groups of specimens.

Histochemical stains, including hematoxylin and eosin stain and safranin-O-fast green stain, are performed with usual standard procedures, except for Safranin-O stains are performed using 0.25 g/100 ml of SIGMA fast green ECF as working solution, which can demonstrate the extracellular matrix changes between GAGs (orange) and collagens (green).

The photos are taken by Nikon COOLSCOPE II digital microscope with unadjustable and consistent LED light source. We use Adobe Photoshop CS4 to calculate the ratios of blue/red peaks for safranin-O stained at most green discoloration sites (Figure 2A). We also calculate the color intensity ratios of red/blue peaks for selected H and E stained most eosinophilic discoloration sites to get the quantitative data and simulate colorimetry (Figure 2B).

Averages of top 3 clonal proliferative fibrochondrocyte numbers are counted to evaluate the severity of chondrocytic proliferation.
Kruskal-Wallis test was performed for significance check. The p-value was set at 0.05 for all statistical analysis.

| Group          | N  | Mean Rank |
|----------------|----|-----------|
| Index 1        |    |           |
| HIVD           | 9  | 10.00     |
| Spondylolisthesis | 10 | 15.10     |
| spinal stenosis | 7  | 15.71     |
| Total          | 26 |           |
| Index 2        |    |           |
| HIVD           | 9  | 7.50      |
| Spondylolisthesis | 10 | 16.85     |
| spinal stenosis | 7  | 16.43     |
| Total          | 26 |           |

**Test Statistics**

|                | Chi-Square | Df | Asymp.sig |
|----------------|------------|----|-----------|
|                | 2.913      | 2  | 0.233     |

Table 2: Statistical analysis of B/R ratios for safranin-O stain (index 1), and R/B ratios for H and E stain (index 2) (calculated at the mid-term of study, not include all the cases of spondylolisthesis and spinal stenosis).

**Results**

The average age (mean=52.9) of HIVD patients is significantly younger than those of spinal stenosis (68.6) and spondylolisthesis (62.0) (p=0.036). Normal disc with gray colored nucleus pulposus, and reddish annulus fibrosus due to more type I and type II collagen fibers [3] are discernible on H and E stain (Figure 3A). The safranin-O stain demonstrates outer rim of annulus fibrosus with green colour and indicating very low GAG substance. Middle and inner rim of fibrosus reveals highest amount of GAG content (orange), and intermediate intensity of GAG substance over nucleus pulposus (Figure 3B). The eosinophilic changes (Figure 3C) of focal nucleus pulposus on H and E stain of spinal stenosis (mean rank=16.43) or spondylolisthesis cases (mean rank=16.85) are significantly more than those of HIVD cases (mean rank=7.5) (Table 2) (p=0.014). However, the green discoloration (loss of GAG)(Figure 3D) detected on safranin-O-fast green stain of spinal stenosis (15.71) or spondylolithesis cases (15.10) is non-significantly more than those of HIVD cases (mean rank=10.00) (Table 2) (p=0.233).

Focal clonal proliferation of chondrocytes over the areas other than endplate and outer annulus fibrosus is a consistent phenomenon in all three types of degenerative disc, while lesser degree of the proliferation among the HIVD cases (p=0.169). Indeed, all three groups have abnormal chondrocytic proliferation (Figure 4) comparing with the control group (p=0.009, Table 3).

The excised disc of HIVD may show mild loss of glycosaminoglycans over nucleus pulposus, but relatively intact nucleus pulposus. The excised disc tissue of spinal stenosis has adequate glycosaminoglycans, actively proliferative chondrocytes, but seems to have more possibility of fibrinoid change with loss of glycosaminoglycans or even necrosis over focal nucleus pulposus tissue.

**Cell Number**

|                | N  | Mean  | Std. Deviation | Std. Error |
|----------------|----|-------|----------------|------------|
| Cadaver        | 5  | 4.6000| 2.60778        | 1.16619    |
| HIVD           | 9  | 13.7778| 7.66123        | 2.55374    |
| Spondylolisthesis | 12 | 18.4167| 9.61493        | 2.77559    |
| Spinalstenosis | 10 | 19.1000| 7.41212        | 2.34392    |
| Total          | 36 | 15.5278| 9.04043        | 1.50674    |

**Anova**

|                | Sum of Squares | Df | Mean Square | F    | Sig. |
|----------------|----------------|----|-------------|------|------|
| Between Groups | 852.400        | 3  | 284.133     | 4.528| 0.009|
| Within Groups  | 2008.128       | 32 | 62.754      |      |      |
| Total          | 2860.528       | 35 |              |      |      |

**Table 3:** Statistical analysis of the degrees of clonal chondrocytic proliferation reveals no significant differences among the study groups, but abnormal increase observed comparing from the small control group (p=0.009).
The excised disc tissue of spinal stenosis has adequate glycosaminoglycans, actively proliferative chondrocytes, but have more possibility of eosinophilic degeneration change with focal loss of glycosaminoglycans or even necrosis over nucleus pulposus tissue (p=0.014). The spondylolisthesis disc may display uneven distribution of glycosaminoglycans, including focal loss of it, over nucleus pulposus tissue.

Figure 4: Dot plotting chart displays the distribution of average numbers of top 3 chondrocytic colonies of each specimens.

Discussion

Low back pain frequently involves some pathophysiologic situations of intervertebral disc or osteoarticular compartments of spine. However, HIVD, spondylolisthesis and spinal stenosis are 3 main causes with different etiologies. We choose 5 specimens from autopsy cases as control group to compare the tissue characters and morphology with histochemical stains, which are easier than ultra-structural studies or molecular studies to be kept in acceptable experimental quality. Degeneration of intervertebral disc has been partially proved by our semi-quantitative study. The nature of eosinophilic change on the degenerative disc is correlated from preservation of GAG to loss of GAG or even necrotic change. So the eosinophilic changes should be subdivided into more than one mechanism that is not clarified in this study. The uneven loss of GAGs expressed by safranin-O stain within the degenerative discs unlike the mild even loss of orange coloration on the nucleus pulposus of the 81-year-old cadaver. Disc cell clusters in pathological human intervertebral discs is well documented [4]. Clonal chondrocytic proliferation may develop in brown or greenish areas of nucleus pulposus with safranin-O stain (Figure 5A and 5B) or the inner rim of annulus fibrosus, as nidus of disc degeneration mentioned by Shamji et al. [5]. Ovoid to polymorphus eosinophilic globules can seldomly be seen within the clonal condrocytic territory or in between the territories.

Occasional foci of proliferative “isolated” chondrocytes with an indistinct boundary, may be a kind of chondroid metaplasia, are observed. Taking the example of a focus found in a 79-year-old patient with recurrent herniation of intervertebral disc (Figure 5C), and the other example of involved end plate happened to a 40-year-old patient with L5/S1 spondylolisthesis and disc herniation (Figure 5D). Migration of the atypical chondroid metaplastic tissue from nucleus pulposus into surrounding annulus fibrosus or end plate are suspected, and supported by observable tissue gap round the atypical metaplastic tissue foci after tissue processing. It might be hypothetically considered a tissue defect made by the bullet-like chondroid tumorlet over inner rim of annulus fibrosus may weaken the annulus fibroosus and eventually disc bulging or herniation.

Neovascularization by ingrowth of capillares are noted in few cases over focal fibrous perichondrium or nearby outer rim of annulus fibrosus, but vary rarely found over other areas of the intervertebral disc.

The pathologic microsection technique for disc specimens are relatively difficult due to variable textures within the intervertebral disc, ranging from soft to dry tough and even bony hard. Decalcification can either affect the result of histochemical staining unless adequate neutralization of the decalcified specimens in running water. Dividing the key specimens of operated degenerative disc into separate jars for further recognition of the characteristic pathologic changes and then molecular pathologic changes are the next goal of this study group.
Conclusion

We notice that a few pathologic pictures do exist in all 3 groups of degenerative disc, but HIVD specimens have less severity of morphologic changes than those of spinal stenosis and spondylolisthesis with the eosinophilic change of nucleus pulposus having the significant difference. Spondolthesis and spinal stenosis cannot exclude the possibility of disc-induced diseases. However, regenerative proliferation (clustering) of chondrocytes is present in all of degenerative intervertebral disc, no matter what the clinical course is chronic or acute. The chondrocytic proliferation in the fibrocartilage of disc might be similar to the chondrocytic clustering within the synovial joint cartilage of grade 3-4/6 osteoarthritis which is a grading system proposed by Pritzker et al. [6].

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