The relationship between pregnancy-associated plasma protein-A (PAPP-A) and human intervertebral disc degeneration

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Summary. Pregnancy-associated plasma protein-A (PAPP-A), a metalloproteinase expressed by a number of cell types, has the important role of cleaving insulin-like growth factor (IGF)-binding protein-2, -4 and -5 in the extracellular matrix and thus freeing up IGF and making it available to cells. The objective of the present study was to utilize immunocytochemical analysis to determine the proportion of PAPP-A-positive cells in a large group of disc specimens which covered the spectrum of changes from relatively healthy Thompson grade II discs to extremely degenerate grade V discs. Work was approved by our institutional human subjects review board. Seventy-two intervertebral disc specimens were assessed for immunocytochemical localization of PAPP-A and the proportion of positive cells determined in the outer annulus, inner annulus and nucleus pulposus. The percentage of PAPP-A positive cells in both the outer and inner annulus correlated significantly with increasing stages of disc degeneration in a fashion which was not dependent upon subject age. There was no significant difference in the percentage of PAPP-A positive cells in the inner annulus of herniated vs non-herniated sites, or in the outer annulus of herniated vs non-herniated sites. Data reported here point to the importance of additional work to elucidate the role of PAPP-A in intervertebral disc aging and degeneration.

Key words: PAPP-A, Disc degeneration, Immunocytochemistry

Introduction

Pregnancy-associated Plasma Protein-A (PAPP-A) is a newly recognized metalloproteinase (MMP) expressed by several cell types, including fibroblasts, osteoblasts and smooth muscle cells (Lawrence et al., 1999; Mazerbouga et al., 2001; Ortiz et al., 2003; Sivanandam et al., 2004; Conover et al., 2004, 2006; Kumar et al., 2005; Qin et al., 2006; Jia and Heersche 2007). Recent work has also shown that PAPP-A is present in the human intervertebral disc (Gruber et al., 2008b). PAPP-A has an extremely important role because it cleaves IGF binding proteins (IGFBP) -2, -4 and -5 in the extracellular matrix (ECM), thereby modulating local IGF bioavailability to nearby cells (Kumar et al., 2005; Boldt and Conover 2007). Insulin-like growth factor-I (IGF-I) has long been recognized as an important regulator of chondrocyte phenotype and differentiation. Some cells, such as human osteoarthritic chondrocytes, show a low or hypo-responsive response to IGF-I (Tardig et al., 1996). Compared to cartilage, much less is understood about IGF-I in the human intervertebral disc. Okuda et al., have suggested that the age-related decrease in IGF-I dependent proteoglycan synthesis in disc cells from the rat may be caused by increased IGF-binding proteins (IGF-BPs) during early aging (Okuda et al., 2002). This points to the importance of obtaining a greater understanding of how IGF-I is regulated in the disc.

Conover et al., have studied many aspects of PAPP-A in their recent publications, including its regulation in cultured human osteoblasts (Conover et al., 2004) and smooth muscle cells (Bayes-Genis et al., 2001; Conover et al., 2006), and have reviewed that it is expressed by a variety of cell types (Boldt and Conover 2007). PAPP-A is now viewed as an important determinant of growth and development via its ability to proteolyze IGFBPs.
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and thus regulate IGF bioavailability in the microenvironment around cells. Thus it is timely to further study the specific role of PAPP-A in the pathophysiology of disc degeneration.

There is some evidence that there may be increased IGFBP in the aged disc. In vitro work by Okuda et al., looked at cultured disc cells from aging rats (Okuda et al., 2002). They found an age-related decrease in proteoglycan synthesis and an increase in IGFBPs. Osada et al., have cultured bovine disc cells, and found that fetal cells were much more active than adult cells (Osada et al., 1996). Walsh et al., examined mouse disc cells and also found a muted response to IGF-I in vitro (Walsh et al., 2004). It is important to note, however, that these were data derived from in vitro work which may have influenced cell responses.

The objective of the present study was to utilize immunocytochemical analysis to determine the proportion of PAPP-A-positive cells in a larger group of disc specimens (compared to our previous analyses (Gruber et al., 2008b) which covered the spectrum of changes from relatively healthy Thompson grade II discs (Thompson et al., 1990) to extremely degenerate grade V discs. In addition, we investigated potential differences between cervical compared to lumbar disc sites, and potential relationships with disc herniation.

Material and methods

Clinical study population

Experimental study of human disc specimens was approved prospectively by the authors’ Human Subjects Institutional Review Board at Carolinas Medical Center. The need for informed consent was waived since disc tissue was removed as part of routine surgical practice. Scoring of disc degeneration utilized the Thompson scoring system; this system scores disc degeneration over the spectrum from healthy discs to discs with advanced degeneration (grade V, the most advanced stage of degeneration) (Thompson et al., 1990). Patient specimens were derived from surgical disc procedures performed on individuals with herniated discs and degenerative disc disease. Surgical specimens were transported to the laboratory in sterile tissue culture medium. The study population contained the following Thompson grade distribution: Lumbar specimens: 6 grade II specimens; 21 grade III specimens, 21 grade IV specimens, and 1 grade V specimen; cervical specimens: 2 grade III specimens, 20 grade IV specimens, and 1 grade V specimen.

Immunolocalization of PAPP-A in human disc tissue and cell counts

Specimens were fixed in 10% neutral buffered formalin for no longer than 24 hours, and stored in 70% ethanol until embedding in paraffin without decalcification. Paraffin sections were cut at 4µm, collected on PLUS slides (Allegiance, McGaw Park, IL) and dried at 60°C. Sections were deparaffinized in xylene (Allegiance) and rehydrated through graded alcohols (AAPER, Shelbyville, KY) to distilled water. The remainder of the procedure was performed using the Dako Autostainer Plus (Dako, Carpinteria, CA) automated stainer. Endogenous peroxidase was blocked using 3% H₂O₂ (Humco, Texarcana, TX). Slides were incubated for one hour with anti-human PAPP-A (polyclonal rabbit anti-human; Dako) at a 1:50 dilution. Universal Negative control rabbit (Dako) was used as a negative control. Secondary antibody was Dako LSAB2 biotinylated Link for HRP/AP for 10 minutes followed by peroxidase-conjugated streptavidin (Dako) for 10 minutes and DAB (Dako) for 5 minutes. Slides were removed from the stainer, rinsed in water, counterstained with 0.002% light green, dehydrated, cleared and mounted with resinous mounting media (Cytoseal XYL, Richard-Allan Scientific, Kalamazoo, MI).

The proportion of cells positive for PAPP-A immunolocalization was determined by counting the number of positive and negative cells in outer annulus, inner annulus and nucleus pulposus regions of the disc. No measurements were performed in sites of morphologic evidence of disc herniation. Two authors (HEG and LB) performed cell counts, and data were checked for intraobserver consistency. The scorer was blinded to grade and age at the time of histologic scoring. For cell counts, the average total number of cells scored were as follows: outer annulus, 299±190 (53) (mean ± S.D. (n)); inner annulus, 477±279 (67), and nucleus pulposus, 300±68 (4).

Statistical analyses

Statistical analysis of data utilized standard methods using SAS software (version 11; SAS Institute, Cary, NC), including Fisher’s exact test, ANOVA and Tukey’s test where appropriate, correlation analyses and t-tests. Data were tested for normality using the Shapiro-Wilk test. When testing for linear relationships between Thompson disc grades and the percentage of cells positive for PAPP-A immunolocalization, Spearman’s correlation coefficient, a non-parametric test, was used because Thompson grades were measured on an ordinal scale. Since only four nucleus specimens were available for analyses, these nucleus data were not included in our correlation analyses.

Results

Table 1 summarizes the subject demographic data according to Thompson grade and region of the disc evaluated (outer annulus, inner annulus and nucleus). (Note that some tissue samples contained both regions of outer annulus and inner annulus, and inner annulus and nucleus pulposus.) The study population contained 21 males and 50 females, and disc specimens were obtained from 49 lumbar and 22 cervical disc sites.

Figures 1-3 show representative immunocytochemical localizations in the outer annulus (Fig. 1A),
inner annulus (Fig. 2), and nucleus pulposus (Fig. 3). A negative control is shown for the outer annulus in Figure 1B. When disc cells were present in clusters (Fig. 2B), both positive and negative localization patterns were present. Degenerating discs also can contain annulus cells encapsulated by concentric layers of matrix material. Such cells often displayed PAPP-A localization as shown in Figure 2C).

The percentages of PAPP-A positive cells in the outer annulus and in the inner annulus grouped according to disc degenerative stage are shown in Figure 4A and B. Although both relationships are highly significant (p=0.0063 and <0.0001), there was a stronger correlation present with the inner annulus data. In the outer annulus, the percentage of cells positive for PAPP-A localization was 24% larger in more degenerated grade V discs than in grade II discs (Fig. 5A). In the inner annulus, however, there was a 50% increase in cells positive for PAPP-A localization when comparing grade II discs to grade V discs (Fig. 5B). (Since only four specimens were analyzed from the nucleus pulposus, these data were not included in the analyses relating data to disc grade).

To further investigate the relationship between disc degeneration and the proportion of PAPP-A positive cells in the inner annulus, we utilized ANOVA analysis with Tukey’s test. Since there were few specimens in the Thompson grade V group, these were pooled with the grade IV specimen data for this analysis. Analysis showed that there were significant differences between grade II discs and grade III and pooled grade IV-V discs, and between grade III and pooled grade IV-V discs, p=0.05. As shown in Fig. 4, 5, the proportion of positive cells increased with increasing disc degeneration.

In the present study we also investigated here potential differences due to derivation of data from cervical vs. lumbar disc sites. (The four specimens from the nucleus pulposus were all derived from lumbar disc sites.) As shown in Table 2, there were no significant differences due to site. All data are from lumbar discs.

### Table 1. Patient demographic data and thompson grades*

<table>
<thead>
<tr>
<th>Thompson Grade</th>
<th>Outer Annulus**</th>
<th>Inner Annulus**</th>
<th>Nucleus**</th>
<th>Age (years) *</th>
</tr>
</thead>
<tbody>
<tr>
<td>II</td>
<td>3</td>
<td>5</td>
<td>1</td>
<td>36.1±11.2 (6)</td>
</tr>
<tr>
<td>III</td>
<td>19</td>
<td>19</td>
<td>3</td>
<td>43.4±11.6 (22)</td>
</tr>
<tr>
<td>IV</td>
<td>29</td>
<td>41</td>
<td>0</td>
<td>53.8±10.2 (41)</td>
</tr>
<tr>
<td>V</td>
<td>2</td>
<td>2</td>
<td>0</td>
<td>49.0±11.3 (2)</td>
</tr>
<tr>
<td>Pooled Grades IV and V</td>
<td>31</td>
<td>43</td>
<td>0</td>
<td>53.6±10.2 (43)</td>
</tr>
</tbody>
</table>

* Data expressed as means ± S.D. (n). Age data represent mean age for all subjects of respective disc grade. ** Data are number of specimens for each disc site in the respective disc grade.

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**Fig. 1.** A. Immunolocalization of PAPP-A in cells in the outer annulus. B. Negative control from an adjacent section. x 320

**Fig. 2.** Immunolocalization of PAPP-A in cells in the inner annulus (A). When cells were present in clusters (B) in the inner annulus, both positive and negative cells were present. Cells with negative localization are marked by arrows. C illustrates an inner annulus cell which is surrounded by a concentric encapsulating matrix (arrow) and is positive for PAPP-A localization. Note the prominent degeneration and loss of matrix in this specimen (*). A, x 275; B, x 250; C, x 320
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Table 2. Comparison of data by disc site (lumbar vs cervical)*.

<table>
<thead>
<tr>
<th></th>
<th>Lumbar</th>
<th>Cervical</th>
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<tr>
<td>Age (years)</td>
<td>47.0±13.7 (49)</td>
<td>53.0±6.8 (23)</td>
</tr>
<tr>
<td>Outer annulus: % Positive for PAPP-A localization</td>
<td>75.0±14.9 (39)</td>
<td>82.4±14.8 (14)</td>
</tr>
<tr>
<td>Inner Annulus: % Positive for PAPP-A localization</td>
<td>63.7±20.2 (45)</td>
<td>76.2±10.3 (22) **</td>
</tr>
<tr>
<td>Nucleus pulposus</td>
<td>16.5±9.5 (4)</td>
<td>--</td>
</tr>
</tbody>
</table>

*Data expressed as means ± S.D. (n). ** The mean percentage localization in the cervical inner annulus specimens was significantly greater than that in the corresponding lumbar sites, p=0.0013.

Fig. 3. This image of the nucleus pulposus shows a region of relatively few cells, some of which were positive for PAPP-A immunolocalization. x 720

Fig. 4. Presentation of the proportion of cells positive for PAPP-A immunolocalization in the outer annulus (A) and in the inner annulus (B).

Fig. 5. Mean data for the proportion of cells positive for PAPP-A localization in the outer annulus (A) and inner annulus (B). Data are presented as means ± S.D.
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Table 3. Comparison of data derived from herniated vs non-herniated sites.

<table>
<thead>
<tr>
<th></th>
<th>Herniated Sites</th>
<th>Non-Herniated Sites</th>
</tr>
</thead>
<tbody>
<tr>
<td>Outer annulus: % Positive for PAPP-A localization</td>
<td>80.3±11.36 (28)</td>
<td>73.37±17.98 (25)</td>
</tr>
<tr>
<td>Inner Annulus: % Positive for PAPP-A localization</td>
<td>68.95±17.31 (39)</td>
<td>66.31±20.19 (28)</td>
</tr>
</tbody>
</table>

**Data expressed as means ± S.D. (n)**

Discussion

The proteolytic activity of PAPP-A targets selected IGFBP 2, 4 and 5. We have previously shown that these binding proteins are expressed by the cells of the human intervertebral disc (Gruber et al., 2008b). Findings from the work of Kumar et al. examining conditioned medium of myoblasts showed that PAPP-A did not cleave IGFBP-3 (Kumar et al., 2005). These investigators proposed that PAPP-A plays a critical role by enhancing the bioavailability of IGFs by its degradation of the IGFBPs and the subsequent release of IGFs into the surrounding tissues (Kumar et al., 2005), and it is likely that it plays a similar role in the intervertebral disc extracellular matrix.

IGF-1 is an important cytokine in the disc from a number of perspectives. As noted above, Okuda et al., studied cultured disc cells from aging rats (Okuda et al., 2002) and identified an age-related decrease in proteoglycan synthesis and an increase in IGFBPs. Work from our laboratory has found that IGF-1 has an anti-apoptotic effect and an anti-senescent effect on cultured human annulus cells (Gruber et al., 2000, 2008a). Thus a better understanding of PAPP-A, which has the ability to free up IGF in the disc matrix, is biologically important.

Whether in lumbar discs or cervical disc, the findings reported here show that a greater proportion of disc cells contain PAPP-A in the more advanced stages of disc degeneration (Figs. 4, 5), a relationship which was shown to be independent of subject age by our analysis of covariance assessment.

Studies on matrix synthesis and degradation in the disc have shown that pro-inflammatory cytokines, such as TNF-α, are important cytokines involved in the pathogenesis of disc degeneration (Freemont et al., 2002; Weiler et al., 2005). TNF-α is elevated in the degenerating disc, induces changes in the biochemical features of discs, notably a loss in proteoglycans, stimulates production of nerve growth factor (Abe et al., 2007), and has been found to regulate MMP-3 in nucleus pulposus cells (Seguin et al., 2005). These data are important since there is an accumulating body of evidence showing that PAPP-A is the target of inflammatory cytokines (Resch et al., 2004).

In addition to studies on TNFα, work by Hoyland et al., emphasizes the importance of the role of IL-1 in the degenerating intervertebral disc. LeMaitre et al., have reported increased levels of IL-1 in degenerate human discs (LeMaitre et al., 2005). IL-1 was also found to significantly downregulate types I and II collagen and aggrecan (LeMaitre et al., 2005). There are also data showing that IL-1 beta will stimulate, and the IL-1 receptor agonist inhibit, matrix degeneration in intact human discs (Hoyland et al., 2008). Current three-dimensional in vitro work in our laboratory, however, has shown that both II-1 and TNFα will stimulate in vitro production of PAPP-A by human annulus cells (Gruber et al., 2010); we look forward to additional studies of both in vivo and in vitro effects of inflammatory cytokines on the intervertebral disc.

We should mention that a shortcoming of the present study is the absence of concomitant measurement of IGF-1 and the relevant IGF-binding proteins. This would have been very helpful since data from the aging rat nucleus pulposus by Okuda et al. (2002) have been interpreted to show a loss of proteoglycans caused by downregulation of IGF-1 receptor in addition to an increase in IGF-binding proteins. Clearly future human annulus and nucleus pulposus studies which include such analyses will be important contributions to the literature.

In summary, the work presented here is important because it provides new insight into disc cell expression of PAPP-A during disc degeneration. As the disc degenerates, a significantly greater proportion of cells express PAPP-A. We look forward to additional studies from in vivo and in vitro investigations to further analyze PAPP-A’s role in disc aging and degeneration.

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References


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