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Citation for published version (APA):

Document license:
TAVERNE

DOI:
10.1089/ten.tea.2015.0121

Document status and date:
Published: 01/01/2016

Document Version:
Publisher’s PDF, also known as Version of Record (includes final page, issue and volume numbers)

Please check the document version of this publication:
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The Stimulatory Effect of Notochordal Cell-Conditioned Medium in a Nucleus Pulposus Explant Culture

Stefan A.H. de Vries, MSc,1 Marina van Doeselaar,1 Björn P. Meij, DVM, PhD,2 Marianna A. Tryfonidou, DVM, PhD,2 and Keita Ito, MD, ScD1,3

Objectives: Notochordal cell-conditioned medium (NCCM) has previously shown to have a stimulatory effect on nucleus pulposus cells (NPCs) and bone marrow stromal cells (BMSCs) in alginate and pellet cultures. These culture methods provide a different environment than the nucleus pulposus (NP) tissue, in which the NCCM ultimately should exert its effect. The objective of this study is to test whether NCCM stimulates NPCs within their native environment, and whether combined stimulation with NCCM and addition of BMSCs has a synergistic effect on extracellular matrix production.

Methods: Bovine NP tissue was cultured in an artificial annulus in base medium (BM), porcine NCCM, or BM supplemented with 1 µg/mL Link N. Furthermore, BM and NCCM samples were injected with 10^6 BMSCs per NP sample. Samples were cultured for 4 weeks, and analyzed for biochemical contents (water, glycosaminoglycan [GAG], hydroxyproline, and DNA), gene expression (COL1A1, COL2A1, ACAN, and SOX9), and histology by Safranin O/Fast Green staining.

Results: Culture in NCCM resulted in increased proteoglycan content compared to day 0 and BM, similar to Link N. However, only minor differences in gene expression compared to day 0 were observed. Addition of BMSCs did not result in increased GAG content, and surprisingly, DNA content in BMSC-injected groups was not higher than in the other groups after 4 weeks of culture.

Discussion: This study shows that, indeed, NCCM is capable of stimulating NPC matrix production within the NP environment. The lack of increased DNA content in the BMSC-injected groups indicates that BMSCs have died over time. Identification of the bioactive factors in NCCM is crucial for further development of an NCCM-based treatment for intervertebral disc regeneration.

Introduction

The intervertebral disc (IVD) separates vertebral bodies and provides flexibility to the spine. Its hydrated core, the nucleus pulposus (NP), is rich in negatively charged proteoglycans, which attract water. However, a fibrous ring, the annulus fibrosus (AF) constrains the NP and keeps it from swelling. Therefore, the NP is highly pressurized and plays a crucial role in the distribution of loads. With degeneration, nucleus pulposus cells (NPCs), responsible for production of the NP matrix components, such as proteoglycans and collagens, decrease in number and change their phenotype, resulting in a more catabolic behavior. Also, the gel-like NP changes toward a more fibrous tissue, where collagen type 2 is replaced by collagen type 1. The proteoglycan content and, therefore, the water content decreases, resulting in decreased IVD height. This changes the load distribution properties of the NP, causing more load to be exerted on the AF, which can further contribute to the degenerative process. IVD degeneration is associated with low back pain, and current treatment methods mostly aim at pain alleviation, but do not address the underlying mechanisms of IVD degeneration.

To replenish the decreasing number of NP cells and restore the IVD to a healthy state, bone marrow stromal cells (BMSCs) have been employed in regenerative treatment strategies and are currently being tested in clinical trials phase I/II (ClinicalTrials.gov: NCT01860417). BMSCs can either differentiate toward a chondrogenic phenotype and produce matrix themselves, or provide a trophic effect to the...
NPCs. Injection of BMSCs in the IVD has been shown to inhibit degenerative changes in a canine model. The majority of the BMSCs were present in the NP tissue, 12 weeks after injection and degeneration was arrested. However, addition of the BMSCs did not restore disc height. Therefore, to be able to restore the IVD to a healthy state, other or additional methods need to be investigated.

The use of notochordal cells (NCs) seems promising in IVD regeneration. NCs are large vacuolated cells present in the NP tissue of young individuals. It has been observed that their disappearance coincides with the onset of IVD degeneration. Furthermore, species retaining their NC population, such as pigs or certain dog breeds, keep a healthy IVD throughout the largest part of their lives. They may, therefore, play a role in maintaining a healthy IVD, and could be exploited to reverse the degenerative process.

Notochordal cell-conditioned medium (NCCM) obtained from NCs encapsulated in alginate beads, or directly from NC-rich NP tissue has been shown to have a stimulatory effect on NPCs as well as BMSCs in alginate beads and pellet cultures. Porcine NCCM stimulated matrix production by bovine NPCs and BMSCs and human NPCs in alginate bead cultures. Porcine NCCM also increased matrix production and chondrogenic gene expression in pellet cultures of human BMSCs. Furthermore, canine NCCM stimulated proliferation, proteoglycan production, and expression of genes associated with the chondrogenic phenotype by bovine NPCs. Furthermore, in a homologous canine model, NCCM increased proteoglycan production by NPCs and BMSCs, and anabolic gene expression by NPCs.

Altogether these results are promising; NCCM has only been tested on isolated cells in alginate bead and pellet cultures. Such culture methods provide a markedly different environment from the NP tissue, which could alter the cell’s response to stimuli. For example, several studies demonstrated stimulation of isolated NPCs by osteogenic protein-1 (OP-1) However, in vivo the results were mixed. In the degeneration-induced rabbit model, which has a NP populated mostly by NCs, a stimulatory effect of OP-1 was observed. However, in a more relevant chondrodystrophized canine model of spontaneous disc degeneration, with NPC-rich NP, no stimulation after OP-1 injection was found. Furthermore, OP-1 has been used in human clinical trials, but these results were never released, making it likely that no large therapeutic effect was observed. Finally, when OP-1 was eventually tested in moderately degenerated human NP tissue culture, only insufficient stimulatory effects were found. These findings indicate that the translation from cell to tissue scale is not necessarily a straightforward one, and testing an approach in a near in vivo situation will give a better idea of the in vivo feasibility. Similar to OP-1, NCCM, or its bioactive factors, should exert a stimulatory effect in the NP tissue. However, it is unknown whether NPCs within the NP tissue respond to NCCM stimulation, similar to isolated NPCs. Therefore, the aims of this study are 2-fold. First, to test the stimulatory effect of NCCM in an NP explant culture. Second, to test whether the addition of BMSCs, and hence combined stimulation of NP tissue with NCCM and BMSCs, can lead to a synergistic stimulatory effect on extracellular matrix synthesis.

Materials and Methods

Porcine NCCM

NC-rich NP tissue was harvested from the lumbar IVDs of 10 pigs (<3-month-old), obtained from a local abattoir. The IVDs were dissected and opened under aseptic conditions and the NP tissue was taken out and placed in high-glucose Dulbecco’s modified Eagle’s medium (hgDMEM, Gibco; Invitrogen, Carlsbad, CA) +1% penicillin/streptomycin (PS; Lonza, Basel, Switzerland) with 30 mL/g of tissue, as described previously. The porcine NPs were cultured for 4 days at 37°C, 5% O2, and 5% CO2. At the end of culture, the NP tissue was removed from the medium by filtration with a 70-µm pore-size cell strainer, and medium of the different donors was pooled, two by two, to have enough medium for each repeat. The medium was stored at −80°C until further use.

Bovine BMSCs

Bovine BMSCs were isolated by adherence to plastic from iliac bone marrow aspirates from five 4- to 12-month-old cows. They were expanded in hgDMEM supplemented with 1% PS and 10% fetal calf serum (FCS Gold; PAA laboratories, Pasching, Austria) until subconfluence, before being stored at P0 in liquid nitrogen. A new vial of P0 BMSCs from a single donor was thawed and expanded up to P3 for each experiment.

NP explant culture

NP tissue explants were harvested from the caudal discs of five ~2-year-old cows obtained from a local abattoir. The IVDs were dissected, opened, and the NP explant was taken out aseptically using an 8 mm diameter biopsy punch and scalpel, rendering cylindrical explants with a diameter of 8 mm and a height of ~5 mm. Care was taken not to include AF tissue, or to scratch the endplate. The explants were placed in 15 kDa molecular weight cut-off (MWCO) dialysis tubing (Spectra-Por, Rancho Dominguez, CA), and closed tightly to constrain the NP explant and prevent it from swelling. The cell pellet was aspirated, leaving behind 15–20 µL medium by slowly pipetting up and down. The cell suspension was aspirated with a 27G needle and a 100-µL Hamilton syringe (Da Vinci, Rotterdam, The Netherlands) and injected into the designated NP explants. The explants were then wrapped in 100 kDa MWCO dialysis tubing and placed in a Dynemeca fiber jacket (Varodem, Saint-Léger, Belgium), which was sewn closed tightly to constrain the NP explant and prevent it from swelling. Non-BMSC-injected explants were directly wrapped in the 100 kDa membrane and placed in the fiber jacket after PEG shrinkage. This established method of culturing NP tissue has previously shown to maintain tissue integrity and cellular behavior.

For each repeat, explants of one tail were cultured for 4 weeks in base medium (BM: hgDMEM +1% PS, 3% FCS
Histology

One-quarter of each sample was fixed overnight in 3.7% formalin at room temperature, and embedded in paraffin with a tissue processor (STP-120; Microm, Walldorf, Germany). Sections with a thickness of 8 μm were cut using a microtome (Leica, Wetzlar, Germany) and stained with Safranin O/Fast Green, to visualize proteoglycans and collagen and with Hematoxylin for cell nuclei. Pictures were taken with a brightfield microscope (Observer Z1; Carl Zeiss, Jena, Germany).

Gene expression

At day 0 and 28, two-quarters of the sample were snap frozen in liquid nitrogen and stored at −80°C until RNA isolation. The frozen tissue pieces were disrupted with a Mikro-Dismembrator (Sartorius, Goettingen, Germany) for 30 s at 2000 rpm, which was repeated if necessary. When the sample was pulverized, RNA was isolated using TRIzol, and purified using the Qiagen Mini Kit, with an on-column DNA digestion step. RNA quantity and purity was measured using a spectrophotometer (NC-1000; Isogen, de Meern, The Netherlands) and the absence of genomic DNA contamination in isolated RNA was verified with a minus-RT control reaction (iCycler; Bio-Rad, Veenendaal, The Netherlands). cDNA was synthesized using the VILO Kit (Invitrogen). Genes of interest were ACAN, COL1A1, COL2A1, and SOX9. Both reference genes (18S ribosomal RNA [18S]) and glyceraldehyde 3-phosphate dehydrogenase (GAPDH), remained stable throughout the experimental conditions. Gene expression results are shown relative to GAPDH. Gene expression was investigated using real-time PCR (CFX384; Bio-Rad), and expression was calculated according to the 2−ΔΔCt method. Genes of interest and their corresponding primer pairs are summarized in Table 1.

Statistics

Statistics were performed with Statistical Package for Social Sciences (SPSS, version 21; IBM, Armonk, NY). Normality was tested using the Shapiro–Wilks test. A Kruskal–Wallis test was used, followed by a Mann–Whitney test with post hoc Bonferroni corrections. Statistical significance was accepted with p-values <0.05.

Results

NCCM in a NP explant culture

No differences were observed in water content between culture groups (Fig. 1a). The combined GAG content in the

<table>
<thead>
<tr>
<th>Gene</th>
<th>Accession number</th>
<th>Oligonucleotide sequence (5′–3′)</th>
<th>Product size (bp)</th>
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<tr>
<td>GAPDH</td>
<td>NM_001034034</td>
<td>FW: GGGGTGAAACCACGAGA AT</td>
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<td></td>
<td></td>
<td>RV: CCGCCACCTTCAACGGTG</td>
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<td>COL1A1</td>
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<td></td>
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<tr>
<td>COL2A1</td>
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<tr>
<td>ACAN</td>
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<tr>
<td></td>
<td></td>
<td>RV: CACGAACGGGCGCTT</td>
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GAPDH, glyceraldehyde 3-phosphate dehydrogenase; COL1A1, collagen type I alpha 1; COL2A1, collagen type II alpha 1; ACAN, aggrecan; SOX9, SRY (sex determining region Y)-box 9.

Biochemical content

At day 0 and 28, NP explants were weighed and cut in four equal parts. One-quarter was weighed before and after lyophilization. The water content was calculated as the percent difference between the wet weight and the dry weight of the sample. The dried samples were digested overnight at 60°C in papain digestion buffer (100 mM phosphate buffer, 5 mM L-cystein, 5 mM ethylene diamine tetraacetic acid, and 140 μg/mL papain, all from Sigma). The sample digests were assayed for glycosaminoglycan (GAG) content using a dimethylmethylen blue (DMMB) assay with shark cartilage chondroitin sulfate (Sigma) as a reference. DNA content was measured using the Qubit Quantification Platform (Invitrogen). Hydroxyproline, as a measure for collagen content, was assayed for glycosaminoglycan (GAG) content using a di-methylmethylene blue (DMMB) assay with shark cartilage chondroitin sulfate (Sigma) as a reference. DNA content was measured using the Qubit Quantification Platform (Invitrogen). Hydroxyproline, as a measure for collagen content, was assayed using the Chloramine-T assay with a trans-4-methylmethylene blue (DMMB) assay. For the NCCM-treated groups, the GAG in the sample’s wet weight.

Statistics

Statistics were performed with Statistical Package for Social Sciences (SPSS, version 21; IBM, Armonk, NY). Normality was tested using the Shapiro–Wilks test. A Kruskal–Wallis test was used, followed by a Mann–Whitney test with post hoc Bonferroni corrections. Statistical significance was accepted with p-values <0.05.

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tissue sample and medium was significantly increased with NCCM compared to day 0 and BM (Fig. 1b). These results were verified with a histological Safranin O/Fast Green staining, showing a more intense red color in the NCCM group (Fig. 1e). There were no significant differences in the DNA (Fig. 1c) and hydroxyproline (Fig. 1d) content between culture groups.

No differences between culture groups in gene expression of COL1A1, COL2A1, ACAN, and SOX9 were found (Fig. 2). COL1A1 increased in BM (4.9-fold) and Link N (3.7-fold) compared to day 0. COL2A1 decreased in BM (10-fold) compared to day 0.

**BMSCs with NCCM in a NP explant culture**

No significant differences in water content were observed between groups (Fig. 3a). GAG content in the BMSCs and NCCM+BMSCs group was not significantly different from GAG content at day 0 and with NCCM alone (Fig. 3b). These results were verified by histological Safranin O/Fast

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**FIG. 1.** (a) Water (b) GAG in the tissue sample and culture medium (c) DNA and (d) hydroxyproline content as a percentage of the tissues’ wet weight. Statistics for GAG content were performed for the combined medium and tissue GAG content. Bars indicate \( p < 0.05 \). (e) Safranin O/Fast Green staining for each of the culture groups. Link N, BM supplemented with 1 \( \mu \)g/mL Link N. \( n = 5 \), results are represented as mean ± SD. BM, base medium; NCCM, notochordal cell-conditioned medium; GAG, glycosaminoglycan; SD, standard deviation. Color images available online at www.liebertpub.com/tea

**FIG. 2.** Gene expression relative to GAPDH and normalized to expression at day 0. *\( p < 0.05 \) compared to expression at day 0. \( n = 4 \) for BM, \( n = 5 \) for NCCM and Link N. Results are represented as mean ± SD.
Green staining (Fig. 3e). Also, no significant differences in DNA (Fig. 3c) and hydroxyproline (Fig. 3d) content were found between culture groups.

No differences between culture groups were observed for COL1A1, COL2A1, ACAN, and SOX9 expression (Fig. 4). COL2A1 expression slightly decreased in the BMSC (0.28×day 0) and NCCM+BMSC groups (0.34×day 0).

Discussion

This study verifies that NCCM provides a stimulatory effect on NPCs in their natural environment, that is, the NP tissue. This was demonstrated by increased proteoglycan content in the NCCM group compared to day 0 and BM, which was verified by histological Safranin O/Fast Green staining. Addition of BMSCs alone and addition of BMSCs...
combined with culture in NCCM did not lead to an upregulation of matrix production.

The stimulatory effect of NCCM has so far only been tested on NPCs in alginate bead and pellet cultures. In these studies, NCCM had stimulatory effects on matrix production, cell proliferation, and expression of genes associated with a healthy NP phenotype. However, alginate bead and pellet cultures provide a very different environment from the NP tissue. In alginate bead and pellet cultures there is initially no or hardly any matrix present, and different cell–matrix interactions could alter the cell’s response to stimulation. For example, expression of COL1A1 and COL2A1 of NPCs increased when cultured in alginate beads that has been lost during the isolation procedure.

In vivo disruption of the matrix network and add to further fragmentation. These studies, NCCM had stimulatory effects on matrix production, which was verified by Safranin O/Fast Green staining. The anabolic effect of NCCM is not limited to NPCs within the NP tissue, which was observed by the anabolic effect of NCCM in isolated cell cultures.

The increased proteoglycan content in the NCCM group in the current study suggests that, indeed, NCCM stimulates NPCs within the NP tissue, which was verified by Safranin O/Fast Green staining. The anabolic effect of NCCM is not observed on the gene level, as no increase of ACAN or COL2A1 was observed compared to day 0 or BM. The timing of harvest can be important to see changes in gene expression, and possibly, NCCM stimulation was desensitized after 4 weeks of culture and peaked at some point before analysis of gene expression, which could explain why the effect is not visible at the gene level anymore. Nonetheless, the finding that the end product, that is, GAG content significantly increased is more important than finding increased gene expression, as not all signals are translated into end products. Regardless of the treatment, GAGs were also released in the culture medium of all culture groups.

Given that GAGs are also present in NCCM, the GAG content in the culture medium of the NCCM group was corrected for the GAG content in the stock NCCM. This resulted in a positive value, which indicates that GAGs were produced by the cells rather than being incorporated from the medium itself. The latter is further corroborated by the absence of GAG accumulation at the border of the tissue, which was not observed in Safranin O/Fast Green staining.

It was previously found that aggrecan fragments were already present in mildly degenerated human NP tissue, which could have also been the case in the NP samples in this study, given that the NP explants were extracted from degenerated bovine IVDs. When small enough (i.e., smaller than the 100 kDa pore size of the membrane) these fragments could diffuse out of the sample over time. Furthermore, harvesting the NP sample using a biopsy punch could disrupt the matrix network and add to further fragmentation. In an in vivo application of NCCM for IVD regeneration, that is, when NCCM’s bioactive factors are injected in the NP, small aggrecan fragments would likely remain in the disc, thereby still adding to the IVD’s osmotic pressure.

Injection of BMSCs alone, as well as injection of BMSCs combined with culture in NCCM did not increase proteoglycan production in this study. Only 15–20 μL of cell suspension was injected in the NP explant, without any sign of leakage. Also, quickly after injection, the sample was wrapped in the 100 kDa membrane, which should not allow the BMSCs to migrate out. Therefore, an increased DNA content was expected in the BMSC-injected groups, but this was not the case, suggesting that the BMSCs have not survived the 4-week culture period. Recent studies support the idea that in in vitro chondrocyte–BMSC cocultures, BMSCs have a trophic effect on chondrocytes. In coculture without stimulants toward differentiation, the majority of the undifferentiated BMSCs disappeared over time, whereas only a small number of BMSCs differentiated.

This could also explain a loss of BMSCs in the current study, and the lack of increase in SOX9 expression, associated with differentiation toward the chondrogenic phenotype. Furthermore, the question still remains how this translates to the current study, as undifferentiated BMSCs also did not have a trophic effect that resulted in increased matrix production. Previous studies investigating the trophic effects of BMSCs on NPCs and chondrocytes, found only a modest increase in matrix production in isolated cell cultures. Since the NP tissue is already rich in proteoglycans, it is possible that in the current study, addition of BMSCs did have a trophic effect, although not strong enough to result in a noticeable increase in proteoglycan content. Furthermore, trophic effects of BMSCs on NPCs and chondrocytes were established by culture of isolated cells, and it is unknown whether this effect may hold in a tissue culture. The latter is in line with the in vivo findings, where BMSC transplantation alone arrested degeneration, but did not improve the disc height.

Surprisingly, also the combined stimulation with injection of BMSCs and culture in NCCM did not promote GAG synthesis, although NCCM alone did. Possibly, if the BMSCs slowly disappeared over time, the NCCM’s bioactive factors were used by a higher total number of cells in the NCCM+BMSC group, not providing a strong enough stimulation per cell to upregulate their matrix production. Alternatively, caspases activated in dying cells are known to be able to cleave proteins. This way, the apoptotic BMSCs in this study could have counteracted the stimulatory effect of NCCM.

Although the current results further underscore the promising role for NC-secreted factors in IVD regeneration, the question still remains how this translates to the in vivo situation. Intradiscal injection of Link N in an annular puncture rabbit model of IVD degeneration resulted in partial restoration of disc height and expression of ACAN and COL2A1. Also in this study, a positive, but not significant effect of Link N was observed on proteoglycan production. The effect of NCCM in nonoptimized form, however, was as strong as that of Link N. Furthermore, recent studies found that NCCM has a role in inhibiting neurite growth and blood vessel formation, which can together lead to innervation of the IVD and, therefore, cause back pain. These findings further add to the potential of NCCM in IVD regeneration.
NCCM IN A NUCLEUS PULPOSUS EXPLANT CULTURE

The NP explant culture method provides an environment that is more comparable to the in vivo situation than pellet or alginate bead cultures, but still differences exist that could give rise to a different cellular response to NCCM, such as connective tissue growth factor. Identification of the active factors in NCCM will be required to overcome these problems. Recent studies have identified factors in NCCM, that could potentially be involved in its stimulatory activity, such as connective tissue growth factor.

In conclusion, NCCM is capable of promoting matrix production in NP tissue, similar to alginate bead and pellet cultures. This raises further interest in the identification of the active factors secreted by NCs, and their application in IVD repair. In the current study, no stimulatory effect of addition of BMSCs to the NP tissue was observed. Although it is possible that the BMSCs need to differentiate to survive, the reason for their disappearance is uncertain. Therefore, we are unable to conclude on their stimulatory effect when injected in the NP tissue.

Acknowledgments

This work was supported by AOSpine International through an AOSpine Research Network grant (SRN2011_11). Marianna Tryfonidou was supported by the Dutch Arthritis Foundation (LLP22). Furthermore, the authors gratefully acknowledge Jackson Mwale for supplying them with Link N.

Disclosure Statement

No competing financial interests exist.

References


Address correspondence to:
Keita Ito, MD, ScD
Orthopaedic Biomechanics, Department of Biomedical Engineering
Institute for Complex Molecular Systems
Eindhoven University of Technology
P.O. Box 513
5600 MB Eindhoven
The Netherlands
E-mail: k.ito@tue.nl

Received: March 11, 2015
Accepted: September 22, 2015

Online Publication Date: November 3, 2015