Original Article
Early acetabular cartilage degeneration in a rabbit model of developmental dysplasia of the hip

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Received May 17, 2015; Accepted July 6, 2015; Epub August 15, 2015; Published August 30, 2015

Abstract: Background: Mild developmental dysplasia of hip (DDH) causes high morbidity of osteoarthritis (OA) on adult. It is thought that change of collagen and proteoglycans in cartilage may be the direct reasons for osteoarthritis. Objective: To detect the changes of the expressions of type II collagen of acetabular cartilage in early DDH and to investigate the relevance between type II collagen and the degeneration mechanism of the acetabular cartilage. Methods: The rabbit model of DDH was successfully established by applying the method of knee extending and fixing with cylinder cast in which left lower extremity as experimental group and right one as control group, checking with X-ray after 5 weeks. The stains of H&E and toluidine blue were applied on the samples of acetabular cartilage to observe the morphological changes of chondrocytes and extracellular matrix (ECM). The immunohistochemical staining and Western-blot were employed to respectively qualify and quantitate the expression of type II collagen. Results: Pathohistology observing indicated the signs of retrogressive changes of acetabular cartilage in experimental group. Also, the positive stained cells in type II collagen in experimental group was higher based on immunohistochemical staining. The quantitative amounts of type II collagen by Western-blot in experimental group was higher significant difference existed between two groups (t = 2.18, P < 0.05). Conclusions: The expression of type II collagen is correlated to a degeneration of acetabular cartilage and increase obviously in early DDH.

Keywords: Hip, bone disease, developmental, collagen, osteoarthritis

Introduction
Developmental dysplasia of the hip (DDH) refers to hip dysplasia in various forms on pediatric patients at different ages. This is the one of the most common diseases in pediatric orthopedics [1-3]. The reported morbidity of DDH throughout the world varies from 0.14% to 3.5%, averagely 3% [4-6], while the highest morbidity goes to Poland, 6.8% as reported in 2006 [7]. A lot of experiments showed that a perfect DDH model can be obtained with hip joint inflexion and with knee joint fixed and straightened [8-10]. The reason lies in that poses like sitting or squatting of rabbits are similar to the gesture of a fetus in uterus. The extended knee joint makes the hip joint bended and tight hamstring for long time, thus results in hip joint capsule and round ligament get stretched and gradually reaches subluxation or total dislocation.

Mild DDH causes high morbidity of osteoarthritis (OA) on adult. OA is the most common degenerative disease, characterized by cartilage degradation, loss of joint space, subchondral sclerosis, and formation of osteophytes [11, 12]. Some long-term follow-up studies have shown that DDH patients are prone to OA at an earlier age than non-DDH controls, and the symptoms of OA in the DDH patients are more severe and require a higher rate of hip replacement surgery [13, 14]. Pathology basis of OA is primarily degeneration and degradation of articular cartilage, including degradation of enzymes of extracellular matrix (ECM) and/or inhibition of new substrate synthesis, etc. At present, it is thought that change of quantity and quality of collagen and proteoglycans in cartilage are the direct reasons for OA losing its normal biomechanics features. During infancy and early childhood, there is increased turnover of type II collagen in the articular cartilage. Etiological study also
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holds the view that type II collagen gene variation may be related to development of DDH [15]. So far, researches on expression of macromolecule protein such as collagen in acetabular cartilage of DDH model only have few reports. In order to investigate the articular cartilage degenerative changes in early-stage DDH, this experiment detected the expression and changes of type II collagen in early acetabular cartilage of DDH model with immunohisto-

Figure 1. A: Anteroposterior X-ray before fixation; B: Anteroposterior X-ray 5 weeks after fixation: acetabulum dysplasia, enlarged inclination of acetabulum, Shenton’s line was continuous.

Figure 2. A: Anteroposterior X-ray before fixation; B: Anteroposterior X-ray 5 weeks after fixation: acetabulum dysplasia and subluxation, femoral head moved outward, Shenton’s line was not continuous.
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Subjects and methods

Materials

This experiment introduced New Zealand albino rabbits of either gender with big ears, born after 4 to 5 weeks and weigh 0.5-0.8 kg (provided by department of laboratory animals from Fudan University). This study was carried out in strict accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the National Institutes of Health. The animal use protocol has been reviewed and approved by the Institutional Animal Care and Use Committee (IACUC) of Fudan University. Rabbits were intramuscular injected with 5% ketamine hydrochloride 5-10 mg/kg; left lower extremities of rabbits were as experimental sides, make knee joints straightened and fixed with long leg cylinder cast; right lower extremities were as control sides which kept the way it is. Anteroposterior X-rays were taken for pelvis 5 weeks later and got 8 successful rabbit models of DDH. Specific indexes of X-ray for DDH diagnosis [8] were as follow: 1) enlarged inclination of acetabulum, Shenton’s line was continuous (Figure 1A, 1B); 2) femoral head moved outward, Shenton’s line was not continuous (Figure 2A, 2B). After euthanasia, obtained hip joints of both sides under asepsis conditions; take part of acetabular cartilage on each side as sample and kept them in liquid nitrogen for two usages: one was to detect protein expression with Western-blot immunoblotting; the rest was treated with routine histopathology staining and immunohistochemical staining.

Morphological observation

Samples were fixed with 4% paraformaldehyde and decalcified in EDTA decalcifying fluid for 3 to 4 weeks, then dehydrated, cleared and embedded with paraffin. Paraffin was cut into pieces and stained with H&E and toluidine blue. Observe morphous, size and arrangement of acetabular cartilage cells and changes of ECM under Leica-DMRA2 optical microscope.

Immunohistochemical staining

Here we introduced with immunohistochemical EnVision Two-Step Method. First antibodies was type II collagen antibody of 1:200 anti-mouse polyclone (product of Labvision company), second antibodies were ChemMate TM EnVision kit of gene company. Both of them were observed under Leica-DMRA2 optical microscope. Replace first antibodies with PBS as the negative control. Determination index for the results: type II collagen presented positive if chondrocyte was surrounded with brown anachromasis, also with visible brown anachromasis in cytoplasm and wavy or irregular-shaped brown anachromasis in ECM.

Western blot analysis

Here we set beta-actin as internal reference (products of Santa Cruz). Take tissue samples which were kept in liquid nitrogen and added extracting solution of total sample protein, homogenate were iced bath and total protein was extracted. After centrifugation, detected protein density in supernatant fluid under ultraviolet spectrometer. Take 300 μg of total protein and added buffer solution of samples then degenerate 3 min with 100°C. Apply protein electrophoresis separately on 10% (beta-actin) and 8% (type II collagen) PAGE gelatin. After electrophoresis was done, protein transferred to cellulose acetate membrane and sealed with TTBS which include 0.5% skimmed milk powder. Then added first antibodies (working concentration of type II collagen was 1:200; working concentration of beta-actin antibody was 1:200) and react 2 hours at room temperature.

Figure 3. Clustered chondrocytes were seen in experimental group with H&E staining (indicated with arrows). Magnification × 200.
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Washed the extra first antibodies and added second antibodies (Gene Company, working concentration of 1:1000) then react for 1 hour at room temperature and visualized with luminescence substrate. Then the film was scanned and the strap analyzed with photodensitometry. Divided grey scale value of interest protein by grey scale value of internal reference and used the result to correct the error, the results embodied relative amount of interest protein.

Statistical analyses

All data was expressed with mean ± standard error, and processed the collected experimental data with SPSS12.0 statistical package and run t-test of two groups samples. The difference has statistical significance if \( P < 0.05 \).

Results

Morphologic observation under microscope

Experimental group: Fibrosis were visible in many chondrocytes of surface layer with H&E staining; arrangement of cells in middle layer was in disorder; cells in columnar and cell layer were decreased with irregular arrangement and a few clustered chondrocytes (Figure 3); ECM stained by toluidine blue suffered with uneven staining on columnar cell layer (Figure 4A).

Control group: Cell sizes were even with H&E staining, cells in columnar cell layer were arranged in column and perpendicular to the chondrocytes surface, no clustered chondro-
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Figure 6. Olisthrozone with blooming-effect surrounding of chondrocytes (indicated with arrow). Magnification × 400.

Immunohistochemical staining

Experimental group: Type II collagen expressed unevenly in ECM of cartilage layer, cells were surrounded with hyperchromatic, wavy or irregular-shaped collagen in ECM, and obvious hyperchromatic cytoplasm showed in chondrocyte. The positive cells were higher than those of control group (Figure 5A). Some cells suffered olisthrozone with blooming-effect (Figure 6).

Control group: Type II collagen expresses evenly in ECM of normal acetabular cartilage and staining level was low and no staining in cytoplasm of chondrocytes generally (Figure 5B).

Western blot

Visualization of expression of type II collagen protein: Strap of relative molecular mass of 129 × 103 was type II collagen, strap of relative molecular mass of 43 × 103 was β-actin protein (Figure 7).

Relative quantity of protein expression: Divide gray value of interest protein of type II collagen protein by gray value of internal reference β-actin protein and used the result to correct the error and it also presented the relative amount of interest protein.

The result showed that there were type II collagen expression in control group, meanwhile expression of type II collagen in the experimental group were higher, so there was a significant difference between two groups (t = 2.18, P = 0.047).

Discussion

In early period, Wilkinson [16] pointed out that human acetabulum shares similarity with rabbit. Originally, people would drag lower extremity with excessive physical exertion to get DDH animal models while now it has been evolved into using plastic cannula, intramedullary needle and percutaneous fixation to straighten and fix the knee joint [8]. In this experiment, these young rabbits were fixed with plaster fixation and with extended knee joint, so their hips were under mechanical stress of muscle skeleton under a long time and finally got the DDH models successfully, this has been provided a theoretical foundation on the importance of the acquired environment effect on this disease as well as proved that DDH is a “developmental” disease to some extent. 21 rabbits were involved and 16 rabbit models were obtained (8 were applied for this experiment), so the teratogenic rate was 76.2% (16/21) and 3 got dead (two were infected; one accidently failed). To sum up, special attention should be paid to improve success ratio of DDH model: 1) Plaster fixation should be long enough to wrap the whole lower extremity. Because there are large connected aliform skin between the hind limb and gaster-skin, so plaster could not be fixed in groin region completely but just fixed at the middle and lower part of thigh, meanwhile rabbits grow very quick thus make plaster easily peeled off and then knee joint will recover to the flexion mode, so the model establish will be a failure. Therefore, we should frequently observe during model establish and add plaster length at proximal to keep knee joint extended continuously. 2) Proper plaster tightness. Thigh diameter of rabbit is 3 to 4 times of crus diameter. Plaster is cone-shaped and inversed, if plaster is not fixed tightly it will get loosed easily, if it is too tight the lower extremity will be swelling and necrosis. Therefore, after fixation with plaster, open the slot is a good way to reduce stress and adjust plaster tightness. 3) Apply polyamine polymer plaster. Rabbits are rodents and common calcium sulfate plaster

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can be bitten off and destroyed easily while polyamine polymer plaster with strong intensity and high hardness can better prevent this issue.

DDH refers to children of different ages present dysplasia of hip joint in various forms, which is the most common disease of pediatric orthopedics. Mild DDH causes high morbidity of osteoarthritis for grown up. It is reported that 76% hip joint arthritis is caused by DDH, most of them have to take total hip replacement before the age of 50 [17]. Normal chondrocytes secrete proteoglycan and show prunosus after staining with toluidine blue. After compressed, the invisible layer of articular cartilage surface having immunologic barrier to type II collagen antibody will be destroyed, then collagen fibers expose in synovial and cause chondrocytes damages as well as dysfunction and secret abnormal proteoglycans, therefore toluidine blue fails staining or with unevenly staining. In this experiment, it is observed that more visible fibrosis in many chondrocytes of surface layer, chaotic arrangement of cells in the middle layer, cells in columnar cell layer reduction as well as disordered, uneven toluidine blue staining and failed staining on matrix, which indicates that chondrocytes became degenerated due to compression in early DDH and then caused ECM elements changes in quantity and quality such as proteoglycans. This may be the pathological basis of osteoarthritis in prophase.

Aigner et al [19] found that there was an obvious proliferation of chondrocytes in osteoarthritis; and presumed that cartilage degeneration in osteoarthritis started from macromolecule protein synthesis changes, which reflected as synthesis changes of type II collagen protein and proteolytic enzyme, etc. In 1990, the first genetic mutation responsible for OA was reported after researchers found a high rate of OA with a mild chondrodysplasia in the family of a 44-year-old patient with degenerative changes of both hips. The mutation responsible for OA in this family occurred in the type II collagen gene, COL2A1 (collagen, type II, alpha 1.), which encodes a protein expressed almost exclusively in cartilage [20, 21]. This mutation weakens the matrix and leads to premature degeneration of the cartilage [22].

Although a healthy adult’s articular cartilage has rich type II collagen, only little type II collagen mRNA expression in chondrocytes, which proves that anabolism of type II collagen in normal articular chondrocytes is weak. Because the expression of type II collagen mRNA in articular chondrocytes of osteoarthritis significantly increases, it indicates that injured chondrocytes may repair ECM with the extra synthesis of collagen [23]. Cellular metabolism is active in minor injured articular cartilage, gene expression of type II collagen and proteoglycans is higher than that in the control group. With the increment of gene expression, the synthesis of type II collagen increase also and meanwhile the deliquescence and denaturation of collagen increase obviously. Functions of chondrocytes will change and compound more matrix metalloproteinases to participate in deliquescence and denaturation of type II collagen, which will destroy the balance of ECM maintained by matrix components including type II collagen. Collagen total amount in cartilage during early and middle stage of osteoarthritis is up; the increment of synthesis of type II collagen in early stage shows that type II collagen is stained darker than the normal situation.
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especially around the chondrocyte cells and cytoplasm, the increment of collagen is larger than that of destroyed before cartilage reach the certain damage degree [24].

In this experiment, anachromasis of type II collagen in ECM of dysplastic acetabular cartilage suggested that more collagen in ECM; anachromasis of collagen in chondrocytes suggested that stronger synthesis of collagen. The Western-blot detected that the expression of type II collagen protein rose significantly, further explained that when DDH acetabular cartilage under continuous abnormal stress, chondrocytes were activated and cartilaginous degeneration started then synthesized many extra cellular matrix like collagen and other macromolecule protein to adapt this kind of stress changes. Olistherozones with blooming-effect were also visible around chondrocytes in this experiment, the cause of it lay in the changes of chondrocytes function and synthesis of matrix metalloproteinases would destroy matrix components like type II collagen around cells.

Morphology change of acetabular cartilage and abnormal expression of type II collagen in early stage of DDH model which generated under mechanical stress were observed in this experiment, suggesting that these degeneration changes may be a compensatory repair reaction of chondrocytes to environmental factors (such as the mechanical changes, cytokine, etc.). Once this compensatory repair reaction is decompensated, cartilage may appear inconvertible degeneration changes and present the syndrome of osteoarthritis. Therefore, the amount and intensity of II type collagen are probably interrelated with different cartilage degeneration degrees, furthermore, as a biological parameter that reflects cartilaginous degeneration to which degree osteoarthritis will occur has not yet studied.

Acknowledgements

We thank the pathological staff of Children’s Hospital of Fudan University, especially Pro. Chen Lain, for all their help completing this project.

Disclosure of conflict of interest

None.

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