Aging is associated with progressive structural, functional, and metabolic alterations in a variety of tissues and systems. Many of these age-related alterations have been implicated in subsequent impairment of physiological and physical function. Functional changes in the musculoskeletal system are among the most prevalent health problems of middle and old age. Osteoarthritis, the most common form of joint disease, is highly correlated with increased prevalence of degenerative cartilage changes, the underlying mechanisms whereby age is involved in the development or progression of osteoarthritis are so far unknown.

Important age-dependent changes in articular cartilage—which increase in prevalence, extent, and severity with advancing age—concern matrix macromolecules (MMM) such as proteoglycans and collagens. Qualitative and quantitative variations in these components are observed, suggesting an alteration in the physiological cartilage composition and mechanical properties. Moreover, and possibly more important, an age-related reduction in total proteoglycan synthesis after skeletal maturation has been reported.

The role and influence of chondrocytes in this condition remain unclear. Several studies have suggested that the cell density of the whole thickness of the uncalcified articular cartilage declines sharply during the growth and maturation period of skeletal development, but remains relatively constant in adult life. On the other hand, there is evidence that chondrocyte numbers decrease progressively in healthy articular cartilage as a function of age.

Given the reduction in total proteoglycan synthesis with advancing age, we hypothesised that a decline in cell number was an important factor in limiting tissue maintenance. Thus we evaluated the cell numbers in healthy and osteoarthritic human articular cartilage in relation to increasing age, and investigated total MMM synthesis in healthy and osteoar-thritic cartilage to determine a possible influence of age on proteoglycan production.

**METHODS**

**Source of articular cartilage**

Human articular cartilage from 41 patients (aged 37 to 87 years, mean age 62.9 years) without macroscopic osteoarthritic lesions were obtained from surgical specimens at the time of endoprosthetic replacement for acute transcervical femoral fractures and from organ donors. The removal of cartilage from organ donors was approved by the ethics committee of the University of Vienna.

Human osteoarthritic cartilage samples were obtained from 30 individuals who underwent surgery for total hip endoprosthesis. The age of the patients ranged from 35 to 84 years with a mean age of 61.8 years.

**Cell count and cell culture**

Cartilage slices were aseptically dissected from the load bearing joint surfaces and finely minced. In osteoarthritic specimens, dissection of neo-cartilage at the joint margins was avoided. The wet weight of the samples obtained was then measured. Chondrocytes were released by overnight digestion in 0.2% collagenase B (Boehringer Mannheim, Mannheim, Germany) in a recently described, chemically defined, serum-free basal medium. Following digestion, an aliquot of the cell suspension was evaluated for chondrocyte number after Trypan blue staining in a Buerker–Tuerk chamber. The values are given in chondrocytes $\times 10^5$ g wet weight. The fraction of dead chondrocytes was 5–10%, independent of advanced age or underlying osteoarthritic disease. In both osteoarthritic and unaffected cartilage samples, cell clusters formed about 2% of all chondrocytes and could easily be counted. To ensure completeness of tissue digestion, the suspension after collagenase incubation was subsequently filtered using a cell strainer (Falcon, Becton
Dickinson Labware, Lincoln Park, New Jersey, USA), and the weight of the solid residues determined on a microgram scale. The percentage of solid residues after collagenase digestion was uniformly between 1.3% and 5.9% of the cartilage weight, with no difference between healthy and osteoarthritic samples (p = 0.46).

In an additional series of experiments, the influence of age on chondrocyte numbers was assessed histomorphometrically. We investigated histological samples from hip joints from 14 unaffected donors (aged 24 to 68 years, mean 43.5 years) and from 17 patients with osteoarthritis (aged 38 to 91 years, mean 70.4 years). Specimens of cartilage and subchondral bone were fixed in 7.5% formalin for 48 hours, decalcified with EDTA solution, and then paraffin embedded and stained with haematoxylin/eosin following standard protocols. Using the KS 300 version 3.0 software (Carl Zeiss Vision GmbH 1997) three randomly chosen full thickness cartilage areas were marked in each histological sample and the surface areas were calculated. The chondrocyte numbers were counted in the respective areas and the results are given as cell number/mm².

For cell cultures, the chondrocyte filtrate was centrifuged at 500 × g for 10 minutes. Pellets were resuspended in 1:1 mixture of Dulbecco’s modified Eagle’s medium (DMEM, 25 mM Heps + 4500 mg/l glucose + pyridoxine, without sodium pyruvate; Life Technologies, Gaithersburg, Maryland, USA) and Ham’s F12 (Ham’s F-12 +L-glutamine; Life Technologies) containing 10% fetal bovine serum (PAA Laboratories, Linz, Austria) and antibiotics/antimycotics (100 U/ml penicillin G, 100 mg/ml streptomycin, and 0.25 μg/ml amphotericin B; Life Technologies). The isolated cells were grown as monolayer cultures in 24-well plates (Costar, Cambridge, Massachusetts, USA) in quadruplicate at a density of 1 × 10⁵ cells/cm². At 90% confluence of the chondrocyte cultures (approximately 2 × 10⁵ cells/well), serum-containing medium was changed to basal medium and the cells were subsequently cultured for seven days. The medium was replaced every other day. Cultures were maintained at 37°C in humidified air and 5% CO₂.

Statistical analysis
Statistical analysis was done using Student’s t test. A normality test was carried out to determine whether the distribution of the samples was Gaussian. To examine relations between age and MMM synthesis rate, as well as between age and chondrocyte numbers, we used Pearson correlation calculations. Statistical significance was defined as a probability (p) value of <0.05.

RESULTS
Effect of age on chondrocyte numbers of healthy and osteoarthritic human articular cartilage
The evaluation of chondrocyte numbers after tissue digestion revealed an age dependent decrease of cells in the cartilage samples derived from joints without macroscopically visible defects (n = 41). In the patient age group <55 years (n = 12), a mean (SEM) of 4.2 (0.5) × 10⁶ cells/g wet weight of cartilage was found, whereas cartilage from patients >55 years old (n = 29) contained a mean of 2.7 (0.2) × 10⁶ chondrocytes/g wet weight—a significant (p < 0.002) decrease of 35% (lig 1; r = –0.69, p < 0.0001). To further evaluate whether this reduction in cartilage cellularity reflects a possible mechanism for the onset or progression of osteoarthritis, we compared the data from the healthy samples with the chondrocyte numbers in 30 tissue samples obtained from osteoarthritic joints. In contrast to the healthy specimens, the
Values were normalised to protein content and are given as counts per min/mg protein (cpm/mg protein). No age dependent decrease in total difference between these groups could be determined (NS).

seven days. In cells from healthy individuals, [35S]sulphate chondrocytes were cultured in serum-free basal medium for thritic and non-osteoarthritic patients at different ages. The cing age, we investigated cartilage specimens from osteoar-

To investigate whether the levels of biosynthetic activity in articular chondrocytes can biosynthesis in healthy articular chondrocytes between healthy and 17 osteoarthritic patients. In accordance with previous investigations we also did not observe dedifferentiation of cells under the experimental conditions employed in this study no such decrease—either with aging or with underlying osteoarthritis—was observed. On the other hand, there was a mild trend towards a decrease in macromolecular synthesis, and lack of statistical significance does not exclude the possibility that some patients with osteoarthritis may have a more severe impairment of proteoglycan biosynthesis. A possible limitation of the current study is that cell metabolic activity was assessed in monolayer cultures after initial expansion in serum-containing medium, thereby potentially overriding possible differences in metabolic activity between cells derived from osteoarthritic cartilage and controls. On the other hand, previous investigations did not show differences between explant cultures derived from osteoarthritic and healthy articular cartilage studied under serum-free conditions. Moreover, to minimise such potential effects, we exchanged the serum-containing to serum-free medium for seven days before assessing metabolic activity. Finally, in previous investigations we also did not observe dedifferentiation of cells under the experimental conditions employed.

Levels of biosynthetic activity may be decreased as a function of age or osteoarthritis, causing a breakdown of cartilage integrity. Under the experimental conditions employed in this study no such decrease—either with aging or with underlying osteoarthritis—was observed. On the other hand, there was a mild trend towards a decrease in macromolecular synthesis, and lack of statistical significance does not exclude the possibility that some patients with osteoarthritis may have a more severe impairment of proteoglycan biosynthesis. A possible limitation of the current study is that cell metabolic activity was assessed in monolayer cultures after initial expansion in serum-containing medium, thereby potentially overriding possible differences in metabolic activity between cells derived from osteoarthritic cartilage and controls. On the other hand, previous investigations did not show differences between explant cultures derived from osteoarthritic and healthy articular cartilage studied under serum-free conditions. Moreover, to minimise such potential effects, we exchanged the serum-containing to serum-free medium for seven days before assessing metabolic activity. Finally, in previous investigations we also did not observe dedifferentiation of cells under the experimental conditions employed.

Nevertheless our data are in agreement with the reports by Lafeber and Brocklehurst, who previously described no difference in proteoglycan biosynthetic rate between healthy and osteoarthritic cartilage. A constant rate of proteoglycan biosynthesis in healthy articular chondrocytes between 30 and 95 years was also shown by Bayliss and colleagues.

chondrocyte number in osteoarthritic cartilage had a mean (SEM) value of 1.9 (0.1)×10⁶ cells/g wet weight, which differed markedly—by an average of 38%—from the number found in healthy cartilaginous tissue (p<0.0001). It is noteworthy that the decrease in chondrocyte numbers was already evident in younger patients with osteoarthritis and did not change further as a function of age (fig 1; r = 0.27, p = 0.88).

To support the above data, we evaluated the chondrocyte numbers/mm² cartilage in histological sections from 14 healthy and 17 osteoarthritic patients. In accordance with the results from the cartilage digests, we found an age dependent decrease in cell numbers in healthy specimens (r = −0.89, p<0.0001) (fig 2). In osteoarthritic patients no such decrease could be seen (r = 0.12, p = 0.65). When we compared the healthy group with the osteoarthritic group we found a significant difference in cell numbers: healthy group, 91.6 (10.9) cells/mm²; osteoarthritic group, 48.6 (2.5) cells/mm², p<0.007.

Effect of increasing age on biosynthetic activity in articular chondrocytes

To investigate whether the levels of biosynthetic activity in healthy and osteoarthritic human articular chondrocytes—as determined by synthesis of MMM—decreases with advancing age, we investigated cartilage specimens from osteoar-thritic and non-osteoarthritic patients at different ages. The chondrocytes were cultured in serum-free basal medium for seven days. In cells from healthy individuals, [35S]sulphate incorporation rate ranged from 509 to 2689 cpm/mg protein, with an average of 1353 (161) cpm/mg protein (mean (SEM)). In osteoarthritic cultures, isotope uptake ranged from 118 to 2932 cpm/mg protein, with a mean of 1090 (249) cpm/mg protein. Although there was a mild trend towards decreasing MMM synthesis with increasing age, these differences were not significant, either among healthy samples (r = −0.23, p = 0.38) or among osteoarthritic samples (r = −0.19, p = 0.55) (fig 3A). Importantly, when we divided the healthy patient group into a “young” group (<55 years) and an ‘old’ group (>55 years) and compared these groups with the osteoarthritic specimens (fig 3B), no significant difference could be observed (<55 years v osteoarthritic, p = 0.24; >55 years v osteoarthritic p = 0.8).

DISCUSSION

The ability of cells to maintain metabolic homeostasis is believed to decline with advancing age.6 Age represents a major risk factor for the occurrence of osteoarthritis.7 The question arises as to whether articular chondrocytes retain their capacity to uphold tissue homeostasis by synthesising matrix proteoglycans with advancing age. In the case of reduced biosynthetic activity, the maintenance of cartilage extracellular matrix would be impaired and subsequently lead to disruption of tissue integrity, as seen in osteoarthritis. Two possible underlying mechanisms for impaired proteoglycan synthesis in the aging articular cartilage could be a decrease in total cell number or a decline in proteoglycan synthesis rate.

Levels of biosynthetic activity may be decreased as a function of age or osteoarthritis, causing a breakdown of cartilage integrity. Under the experimental conditions employed in this study no such decrease—either with aging or with underlying osteoarthritis—was observed. On the other hand, there was a mild trend towards a decrease in macromolecular synthesis, and lack of statistical significance does not exclude the possibility that some patients with osteoarthritis may have a more severe impairment of proteoglycan biosynthesis. A possible limitation of the current study is that cell metabolic activity was assessed in monolayer cultures after initial expansion in serum-containing medium, thereby potentially overriding possible differences in metabolic activity between cells derived from osteoarthritic cartilage and controls. On the other hand, previous investigations did not show differences between explant cultures derived from osteoarthritic and healthy articular cartilage studied under serum-free conditions. Moreover, to minimise such potential effects, we exchanged the serum-containing to serum-free medium for seven days before assessing metabolic activity. Finally, in previous investigations we also did not observe dedifferentiation of cells under the experimental conditions employed.

Nevertheless our data are in agreement with the reports by Lafeber and Brocklehurst, who previously described no difference in proteoglycan biosynthetic rate between healthy and osteoarthritic cartilage. A constant rate of proteoglycan biosynthesis in healthy articular chondrocytes between 30 and 95 years was also shown by Bayliss and colleagues.

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However, contrary findings were reported by Schafer et al., DeGroot et al., and Verbruggen et al., who all showed a significant reduction in proteoglycan synthesis as a result of aging. A possible explanation for these contradictory findings may be the use of different culture techniques. It must be assumed that all these results are valid, but studies employing in vivo rather than cell culture conditions may have to take mechanisms other than failure of chondrocyte metabolism into account.

As the cell counts in the monolayer cultures employed here were constant we assume that the age-dependent reduction of biosynthetic activity is caused by an impaired cartilage cellularity, because it is well known that chondrocyte numbers decrease in healthy articular cartilage as a function of age.25–28 Furthermore it was reported that cell numbers in osteoarthritic cartilage declined in both fissured and intact joint surface areas compared with healthy specimens.28–30 Our present study confirms these data on healthy cartilage, showing a decrease in chondrocyte numbers with advancing age. In addition we found that cell numbers are markedly reduced in osteoarthritic patients independently of age, even in young patients. Thus the loss in MMM from articular cartilage—which represents an early event in osteoarthriti- sis—may be caused primarily by a decrease in cartilage cellularity. Moreover, with respect to the changes in matrix composition in old age and osteoarthrosis, it is likely that a reduction in chondrocyte numbers plays a role in the formation of condition, as the production of inappropriate non-cartilage-specific matrix constituents47,48 may be caused by an exhaustion or premature aging of the remaining cells.

The mechanisms leading to a loss of chondrocytes in aging and osteoarthritis are still unknown but it may result from a loss of responsiveness to anabolic growth factors40 or possibly from cell death.41,42 Whether the state of reduced tissue cellularity reflects cause or outcome of the osteoarthritic disease process is yet to be investigated.

Conclusions

We found an age-related decrease in chondrocyte numbers in healthy cartilage, whereas cell counts in osteoarthritic tissue were reduced at all ages, even in younger patients. Chondrocytes released from cartilage samples synthesized similar amounts of MMM, regardless of their provenance—that is, whether they were from younger or older healthy cartilage or from osteoarthritic cartilage. These findings suggest that reduction in cartilage cellularity may be an important factor in impairing tissue maintenance during aging and in osteoarthritis.

Authors’ affiliations

K Bobacz, L Erlacher, J Smolen, W B Graninger, Department of Rheumatology, Internal Medicine III, University of Vienna, Vienna, Austria

A Soleiman, Department of Pathology, University of Vienna

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