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DNA METHYLATION IN OSTEOARTHRITIS

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ABSTRACT

Osteoarthritis (OA) is a prevalent disease of articular joints and primarily characterized by degradation and calcification of articular cartilage. Presently, no effective treatment other than pain relief exists and patients ultimately need to undergo replacement surgery of the affected joint. During disease progression articular chondrocytes, the single cell type present in articular cartilage, show altered transcriptional profiles and undergo phenotypic changes that resemble the terminal differentiation route apparent in growth plate chondrocytes. Hence, given its prominent function in both regulating gene expression and maintaining cellular phenotypes, DNA methylation of CpG dinucleotides is intensively studied in the context of OA. An increasing number of studies have been published that employed a targeted approach on genes known to play a role in OA pathophysiology. As of such, it has become clear that OA responsive DNA methylation changes seem to mediate disease associated aberrant gene expression. Furthermore, established OA susceptibility alleles such as *GDF5* and *DIO2* appear to confer OA risk via DNA methylation and respective pathophysiological expression changes. In more recent years, genome wide profiling of DNA methylation in OA affected articular cartilage has emerged as a powerful tool to address the epigenetic changes in their entirety, which has resulted in the identification of putative patient subgroups as well as generic OA associated pathways.

Box 1. List of abbreviations.

OA	Osteoarthritis
ECM	Extracellular matrix
CpG	Cytosine-guanine dinucleotide
SNP	Single nucleotide polymorphism
eQTM	Expression quantitative trait methylation
eQTL	Expression quantitative trait locus
mQTL	Methylation quantitative trait locus
AZA	5-aza-2'-deoxycytidine
DMR	Differentially methylated region
HOX	Homeobox

INTRODUCTION

Osteoarthritis (OA) is currently the most prevalent arthritic disease among the elderly population (1). Patients are subject to impaired mobility, joint stiffness, pain and a significant decrease in quality of life. Presently, no cure exists and patients with end-stage disease ultimately need to undergo a joint replacement surgery of the affected joint. Subsequently, as disease incidence is increasing with the ongoing ageing population, the societal burden both in terms of disabling patients and economic concerns will continue to rise (2). Pathophysiologically, the disease is primarily characterized by progressive degradation and calcification of cartilage in the articular joints, although in recent years it has become apparent that other tissues such as subchondral bone (3-5) and synovium (6-8) play substantial roles in OA pathology as well. The articular cartilage contains a hyaline type extracellular matrix (ECM) made up of collagens, proteoglycans and other structural proteins (9). Articular chondrocytes, the single cell type present in articular cartilage, reside here and maintain tissue homeostasis by remodelling the ECM upon stresses and microtraumas (10). To ensure cartilage integrity and cope with the challenges throughout life, the maturational arrested articular chondrocytes need to continuously employ, possibly error prone, adaptations to changes in the environment (11). In order to facilitate these adaptations, articular chondrocytes are required to dynamically adjust expression of the appropriate genes, while maintaining their specific cellular phenotype (11). In this regard, articular chondrocytes present in OA affected cartilage however, show increased expression of catabolic enzymes (12-14) and have seemingly lost their maturational arrested state (15-17), as they proliferate and regain growth-plate morphology (17), while degrading and calcifying the ECM of the articular cartilage (18,19).

A substantial number of mechanisms are known that regulate gene expression (20) and cell fate persistence (21-23), commonly referred to as epigenetics. While there exists a considerable number of epigenetic layers, such as histone modifications, microRNAs and long non-coding RNAs, the most studied in OA is decidedly DNA methylation. Partly due to its seemingly, relatively stable epigenetic mark on gene expression and partly due to the readily available techniques to measure it. DNA methylation is the phenomenon in which the cytosine nucleotide that is located in cytosine-guanine (CpG) residue pairs gets methylated. Whether CpGs get methylated depend on the local genetic sequence (24), the cellular requirements (21) and environmental factors such the putative pathophysiological state of the respective tissue (24,25). The presence of methyl groups on CpGs is believed to interfere with binding of proteins to the DNA and subsequently affects transcription (26), which is marked by the correlation between the fraction of CpG methylation and gene expression (27), commonly referred to as expression quantitative trait methylation (eQTM) (24,27). As of such, tissue specific methylation profiles are assumed to maintain the respective transcriptional character and identity

of distinct cell types. Interestingly, disease associated single nucleotide polymorphisms (SNPs) that influence aberrant expression, so called expression quantitative trait loci (eQTL), frequently seem to modulate their transcriptomic properties via DNA methylation quantitative trait loci (mQTL) (28,29).

DNA METHYLATION IN OSTEOARTHRITIS

The involvement of DNA methylation in OA pathophysiology is becoming increasingly evident, reflected by the growing body of literature on the subject (13,28-40). However, given the cross-sectional nature of the reports that studied DNA methylation in OA affected articular cartilage, it is currently unclear whether the observed epigenetic differences precede disease onset or are merely consequence of the environmental changes that articular chondrocytes are subject to in OA. As of such, we here discuss two hypotheses that have been proposed to attribute distinct roles of epigenetics in OA pathophysiology.

Firstly, it is hypothesized that individuals with unfavourable epigenetic profiles will be more prone to develop OA and/or progress faster (41,42). As the local genetic sequence significantly influences the DNA methylation state of CpGs, these putative profiles likely arise due to the presence of OA susceptibility SNPs (43). This would be in concordance with the identification of susceptibility genes in complex diseases such as OA, which not uncommonly assert their susceptibility via altered DNA methylation, depending on the presence of associated risk alleles (28,29). For example, OA risk alleles of SNPs in *DIO2* and *GDF5* appear to generate OA predisposing epigenetic profiles and modulate disease associated, epigenetically associated gene expression. The OA risk allele C of the rs225014 T>C SNP, which has been identified in a combined genome wide linkage and association study by our own group (44), marked not only allele specific expression of *DIO2* (45), it additionally mediated epigenetic regulation of the gene (29). More specifically, carriers of the risk allele showed increased transcriptional activity upon hypermethylation at a CpG located in a distal regulatory element. The risk allele T of the rs143383 C>T SNP (46-48), located in the 5' untranslated region of *GDF5*, disrupts a CpG dinucleotide, of which has been shown that it modulates absolute *GDF5* expression in articular cartilage (28). Furthermore, the reported allele specific expression of *GDF5* in heterozygous carriers marked by the rs143383 alleles (49) is affected by the extent of methylation of the respective CpG. Although increasing additional evidence is reported about OA susceptibility alleles, the exact biological mechanisms that confer OA susceptibility is unclear. More specifically, it is unclear whether epigenetic regulation of aberrant gene expression brought about by genetic variation is involved in disease aetiology or disease progression. We can however, conclude that epigenetic regulation of gene expression modulates OA susceptibility, at least at the *DIO2* and *GDF5* loci.

Alternatively, it has been suggested that age-related loss of epigenetic control (50) mediates the loss of the articular chondrocyte's phenotype with ongoing OA, as life-long stresses and adaptations are expected to leave their mark on the epigenome (11). Furthermore, given the altered cellular phenotype chondrocytes acquire in OA and given the role DNA methylation fulfils in maintaining cellular phenotypes, we advocate here that loss of epigenetic control causes reactivation of developmental pathways among articular chondrocytes that are subject to OA and subsequently mediate the morphological changes that are associated with affected articular chondrocytes.

EPIGENETIC CHANGES ACCOMPANY OSTEOARTHRITIS

A growing body of literature reports on altered DNA methylation at specific genes involved in the OA disease process, commonly at precise CpG sites located in regulatory elements near the respective genes (**Table 1**). Frequently, a cross-sectional study design is utilized to study OA associated methylation differences, either between healthy and affected or pairwise between preserved and macroscopically lesioned articular cartilage. Among others, catabolic and developmentally associated genes like *MMP13*, *GDF5*, *SOX9*, *DIO2* and *ADAMTS4* were shown to be differentially expressed between control and affected tissue, presumably mediated by differences in DNA methylation (13,28-34). Although it is not quite clear to what extent DNA methylation changes in OA development contribute to disease onset or progression, a growing number of studies investigates the mechanism by which such changes may affect chondrocyte function.

Table 1. Overview of gene targeted DNA methylation studies in osteoarthritic articular cartilage.

Gene	Methylation in OA	Expression in OA	Sample size (OA, Control)	Joint	AZA	CpG vector	Reference
<i>COL9</i>	Increased	Down	12, 10	Hip	yes	Yes	(31)
<i>GDF5</i>	Decreased	Up	24, 19	Knee and hip	yes	Yes	(28)
<i>DIO2</i>	Increased	Up	52, 52	Knee and hip	yes	No	(29)
<i>IL18</i>	Decreased	Up	18, 12	Hip	yes	Yes	(30)
<i>MMP13</i>	Decreased	Up	17, 12	Hip	yes	Yes	(30)
<i>iNOS</i>	Decreased	Up	13, 15	Hip	no	Yes	(34)
<i>SOX9</i>	Increased	Down	9, 9	Hip	yes	No	(32)
<i>ADAMTS4</i>	Decreased	Up	4, 1	Hip	no	No	(33)
<i>ADAMTS4</i>	Decreased	Up	16, 10	Hip	no	No	(13)
<i>MMP13</i>	Decreased	Up	16, 10	Hip	no	No	(13)
<i>MMP3</i>	Decreased	Up	16, 10	Hip	no	No	(13)
<i>MMP9</i>	Decreased	Up	16, 10	Hip	no	No	(13)

Consequently, various experimental set ups have been applied to confirm the mechanistic relation between DNA methylation and gene expression in the chondrocyte (**Table 1**). A commonly applied experiment that aims to validate observed eQTM loci, as has our own group, is the addition of 5-aza-2'-deoxycytidine (AZA) to the medium of cultured chondrocytes, either using immortalized cell lines or primary chondrocyte cultures (28-30,32,35). AZA is a chemical agent that interferes with the addition of methyl groups on a genome wide scale to newly formed DNA strands during replication. Although valuable information can be gained from such experiments in a global sense, negative outcomes should not per se be considered as experimental falsifications of the earlier observed correlation between DNA methylation and expression, which is likely cell type and locus specific. As of such, not being able to validate earlier observed eQTM loci can very well arise from the fact that the entire genome gets demethylated upon prolonged AZA treatment, of which the transcriptomic consequences potentially overshadow the locus specific relation. Additionally, culturing cells *in-vitro* forces the cells to adapt to an artificial environment, which possibly only resembles the original tissues in a broad sense and subsequently disrupts the regulatory properties of cell type specific eQTM loci.

Another type of validation experiment being applied is cloning the genomic sequence, in which differential methylation was observed, into a CpG-free vector (51) containing a luciferase gene downstream of the multiple cloning site (30,31,34,52). Next, the vector is methylated *in-vitro* only at the cloned region and transfected into chondrocyte like cell-lines. Luciferase activity now depends on the transcriptional activity of the cloned region, which in turn depends on the methylated state of

that respective region. Although using a CpG-free vector has great advantages over AZA treatment, as it does not induce genome-wide altered methylation, the technique will potentially only work for proximal promoters, as long-distance three dimensional genomic structures, which are often seen in gene expression regulation (53-55), depend on the complex interplay of the distant and proximal regulatory elements of genes. Again, as cell culturing is required to apply the technique, measuring promoter activity using a CpG free vector might be influenced by cellular adaptations to the artificial environment.

STRATIFICATION OF OSTEOARTHRITIS PATIENTS BY GENOME WIDE DNA METHYLATION PROFILING

In more recent years, multiple studies have reported on genome wide DNA methylation profiles of articular cartilage in the context of OA (**Table 2**), not least due to development of affordable genome wide DNA methylation arrays, such as the Illumina Infinium HumanMethylation450k BeadChip array. Genome wide profiling of DNA methylation in OA affected cartilage has revealed the presence of multiple OA patient strata, which are characterized by their respective methylation profiles. Firstly, Rushton *et al.* (37) and our group (38) have reported on the distinct genome wide DNA methylation profiles of knee and hip articular cartilage. Secondly, Rushton *et al.* (37) and Fernandez-Tajes *et al.* (36) have reported on a subgroup of patients defined by altered DNA methylation at inflammatory related genes.

Table 2. Overview of current genome wide DNA methylation experiments in osteoarthritic articular cartilage.

Sample size (OA, Control)	Joint	Platform	OA associated pathways	Reference
20, 25	Knee	Illumina 27K	Inflammation, transcriptional regulation, ECM homeostasis	(36)
21, 96	Knee and hip	Illumina 450k	ECM homeostasis, ossification, inflammation, angiogenesis	(37)
24, 24	Hip	Illumina 450k	Development, angiogenesis, inflammation	(39)
16, 16	Knee	Agilent 244k	Development, catalytic activity	(40)
31, 31	Hip and knee	Illumina 450k		(38)
31, 31	Hip and knee	Illumina 450k	Development, ECM homeostasis	(56)

With respect to the first bifurcation of OA patients, the two types of joint specific articular cartilage are distinguished by large differentially methylated regions (DMRs), primarily at genes involved in development and cellular differentiation. Notably, over 10% of DMRs were observed in the four canonical homeobox (HOX) clusters while the spatiotemporal pattern present at the HOX clusters during development was absent, indicating that joint specific DMRs likely bear distinct functionality in adult articular cartilage (38). Both studies that compared knee and hip articular cartilage reported

on joint specific methylation patterns at the four HOX clusters and several homeotic cofactors, such as *IRX3*, *SIX1*, *MEIS2* and *DLX5*. By analysis of transcriptomic data our group subsequently revealed that joint specific DMRs potentially mediate distinctive regenerative capacities of articular chondrocytes residing in different joints, given the developmental and regenerative nature of joint specific expression of genes located in the developmentally associated DMRs (38). Although reports on joint specific methylation profiles might not reveal loci that are directly relevant for OA onset or progression, they undeniably expose the heterogeneity of the disease. These joint specific DNA methylation and associated transcriptomic profiles are concurrent with the reports of joint specific genetic OA susceptibility loci (57). Henceforth, we can conclude that although a substantial amount of common features are present between knee and hip OA, the disease process and subsequent mode of action of putative therapeutic interventions might be different between the two joints.

With respect to the second, another dichotomy is observed by cluster analysis of genome wide DNA methylation profiles of osteoarthritic articular cartilage (36,37). More specifically, putative patient subgroups have been defined by DNA methylation profiles of the affected cartilage that are enriched for differentially methylated CpGs located in or near genes involved in inflammation. The study of Fernandez-Tajes *et al.* used a sparse methylation array and a relatively small sample size consisting of only knee OA patients, which possibly explains why Rushton *et al.* observed substantially more differentially methylated CpGs (1,357 vs. 5,769) that separated the inflammatory cluster of patients in both knee and hip articular cartilage. While enrichment analysis of both studies revealed broadly similar enriched GO terms, being the inflammatory response and cytokine production, in the study of Fernandez-Tajes *et al.* this was only apparent among the hypomethylated CpGs and not among hypermethylated CpGs. Among the consistent inflammatory signals identified by the two studies among knee OA patients were multiple interleukin genes such as *IL10*, *IL16* and *IL19*, but also developmental and ECM related genes such as *RUNX2*, *FGFR1*, *COL6A3* and *COL18A1*. Noteworthy, in the study of Rushton *et al.* stratification by the inflammatory profile, using both hypo- and hypermethylated CpGs, of both knee and hip OA patients is observed, presumably mediated by overlapping pathways, albeit that only 3,496 out of 15,239 (23%) differentially methylated CpGs that separated OA hip patients overlapped with differentially methylated CpGs that separated OA knee patients (37). Very recently, the group of Rushton *et al.* has further investigated the hypomethylated genes that are apparent among the inflammatory profiles in their hip OA patient cohort and report on specific zinc reporter genes that potentially mediate the patient stratification. (58)

Intriguingly, when we performed GO term analysis on the genes that separated hip and knee OA patients consistently in the initial study of Rushton *et al.* and Fernandez-Tajes *et al.*, we observed significant enrichment for ECM maintenance pathways (data not shown). Thus, while one of the

reported clusters of OA patients is presumably characterized by an epigenetic inflammatory profile, a common underlying mechanism appears defined by epigenetic regulation of ECM related genes, such as *COL6A3*, *RUNX2*, *MMP13* and *ADAMTS5*. This proposition is additionally reflected by the fact that all studies report on the enrichment of ECM related pathways (**Table 2**) and by the analysis Fernandez-Tajes *et al.* performed on solely the hypermethylated CpGs, in which they also observe enrichment among ECM maintenance pathways. Finally, the study of Fernandez-Tajes *et al.* grossly compared methylation and additionally expression profiles between the OA subgroups. However, it is evident that the transcriptional consequences of these profiles need to be precisely elucidated, as modulating unfavourable epigenetic but subsequent transcriptomic profiles specifically can potentially attenuate disease onset or progression and might therefore serve as valuable therapeutic targets for the putative subtypes of OA. It should, however, be noted that of the five published genome wide DNA methylation studies in OA, only Fernandez-Tajes *et al.* and Rushton *et al.* have observed separate clustering of OA patients, while Jeffries *et al.* (39), Moazedi-Fuerst *et al.* (40) and our group (56) have not, warranting further research hereinto. Furthermore, not only discovery driven epigenomic profiling of articular cartilage is required to understand possible OA patient heterogeneity. In light of proposed OA subtypes in the literature, epigenetic interrogation of specific pathways by means of burden analyses in pathways such as those in estrogenic sensitivity (59) or apoptosis (60), might be a more powerful approach in the context of the large number of differentially methylated loci in OA reported by increasingly larger studies. In parallel, genome wide DNA methylation profiles of other joint tissues might additionally confer the proposed different OA subtypes.

CONSISTENT DNA METHYLATION DIFFERENCES BETWEEN OSTEOARTHRITIS AFFECTED AND CONTROL ARTICULAR CARTILAGE

Despite the putative segregation of distinct methylation profiles among OA affected joints (37,38) and OA patients (36,37), major communalities have been observed across the different genome wide studies (**Table 2**). Specifically, consistent enrichment of differential methylation among genes involved in development, as well as in collagen synthesis and other ECM maintenance pathways is reported by all studies, particularly among genes from the *RUNX*, *COL* and *MMP* families. Developmental processes thus appear entangled with OA associated degradation and calcification of articular cartilage during disease progression. Although our group has shown that the majority of differentially methylated CpGs do not associate *in-cis* with gene expression, enrichment analysis of OA responsive CpGs that did correlate with gene expression (87 CpGs, 70 genes) also revealed enrichment for ECM maintenance and developmental pathways, consisting of genes such as *ROR2*, *WLS*, *VIT* and *SPP1* (56). Nonetheless, genome wide DNA methylation profiles and the inherent differences between OA and control tissue need thus be interpreted with care and preferably coupled with additional molecular measurements, such as gene expression data, to clarify the biological

consequence. The 76% of genes of which expression did not correlate with *in-cis* DNA methylation could be regulated additionally by other epigenetic mechanisms, such as histon modifications or miRNA mediated silencing or alternatively, might possibly reflect remnants of early developmental or past wound healing processes.

In parallel, as reflected by OA risk alleles that affect DNA methylation mediated gene expression (28,29) and the influential role of the genomic sequence on DNA methylation, partly reflected by the 40% of OA associated CpGs that are affected by the alleles of proximal SNPs (56), it is not unlikely that the total combination of minor genetic variants among OA patient contributes to putative unfavourable epigenomic and subsequent transcriptomic profiles. In this respect, the putative OA inflammatory subtype and the tissue specific mQTLs reported by our own group, could be the consequence of an inherent unfavourable epigenomic profile preceding disease onset. In line with the observed lack of heritability explained by traditional GWA approaches, these putative minor genetic variants might make up for the larger part of heritability in OA.

CONCLUSIONS

Genome wide DNA methylation profiling of OA affected articular cartilage has revealed widespread differences between OA and control tissues. Although the majority of CpGs do not associate with gene expression, CpGs of which we can consequently not conclude whether they play an active role in disease progression, reactivation of developmental pathways due to changes in epigenetic landscape is apparent in OA affected joints. Thus, we hypothesize here that to maintain healthy articular cartilage homeostasis throughout life, the respective chondrocytes utilize epigenetic mechanisms to transcriptomically adapt to the changing environment. Furthermore, the fact that gene expression changes occur via both hyper- and hypomethylation in OA affected articular cartilage, indicates that disease associated differential methylation is unlikely to be the product of a solely passive process. Lifelong adjustment of regulatory mechanisms, such as DNA methylation, is likely subject to stochastic error and subsequent accumulation of epigenetic modifications at developmental pathways, either via inaccurate restoration of the chondrocytes' steady state or via an increasing number of epigenetic adaptations, seem to force chondrocytes towards terminal differentiation (**Figure 1**). This proposition is additionally reflected by the morphological changes that occur in articular cartilage with ongoing OA, as reflected by degradation and calcification, mechanisms also observed among growth plate chondrocytes during development.

In conclusion, studying genome wide DNA methylation in OA has proven to serve as an excellent proxy to probe the underlying pathological cellular processes in OA. In light of the inherent genetically and consequent epigenetically complex nature of OA pathology, the genome wide efforts of recent years that have reported on the DNA methylation landscape of OA affected articular cartilage have delivered valuable insight, insight that would not have become apparent by mere gene targeted approaches.

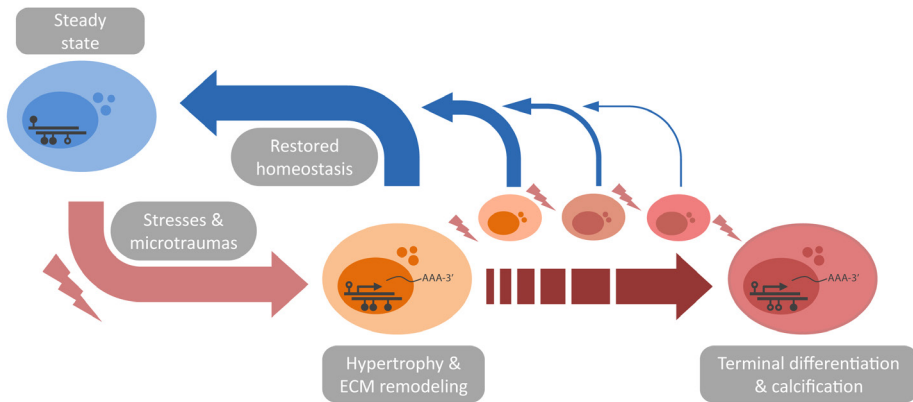


Figure 1. Schematic overview of the proposed loss of epigenetic control among articular chondrocytes. Articular cartilage is subject to lifelong challenges which requires the respective chondrocytes to dynamically adapt gene expression in order to return to and maintain homeostasis and subsequent tissue integrity. This is seemingly accomplished by using epigenetic modifications, reflected by OA associated DNA methylation differences, which are likely not reversed flawlessly upon returning to a steady state. We propose that accumulation of these epigenetic adaptations will eventually lead to altered cellular phenotypes which are unable to return to a healthy, steady state chondrocyte. In parallel, articular chondrocytes of OA patients might already bear an unfavorable epigenomic profile preceding disease onset, which implies that these chondrocytes are more prone to enter the active ECM remodeling state and/or might be less able to return to a steady state. Finally, independent of OA associated epigenetic changes in articular chondrocytes, joint and patient specific epigenomic profiles seem to modulate our proposed mechanism.

FUTURE PROSPECTS OF EPIGENETICS IN OSTEOARTHRITIS

Given that differences in the methylome of OA affected compared to healthy or preserved articular cartilage do not imply downstream effects, reflected by the relatively small number of *in-cis* eQTM loci reported by our group, stresses the need for integration of DNA methylation data with other types of molecular profiling. It is of great interest to the field that the disease relevant, tissue specific epigenomic and transcriptomic QTL profiles are characterised, as they might reveal the predisposing, complex genetic architecture that underlies OA susceptibility. Our own group has undertaken the first step herein and has reported on the effects of SNPs on OA associated DNA methylation and

transcriptional differences, as well as on the direct relation between methylation and expression on a genome wide scale (56). Moreover, the reported genetic variants will potentiate the power of GWA studies, as the multiple testing penalty is substantially reduced when only the functional SNPs, in terms of regulation of transcription, are addressed as opposed to the entire genome. Likely a larger number of SNPs are of relevance, as we have not addressed long distance effects and larger sample sizes are possibly required to gain additional, robust understanding of the reported differences and associations, a point also raised by others (61). In order to test whether indeed loss of epigenetic control confers the pathophysiological changes are apparent with OA progression, longitudinal studies that address the contribution of ageing in altered DNA methylation are required, preferably specifically at genes of which transcriptional changes potentially modulate the disease process. Confirmation of this hypothesis might potentially reveal the genetic drivers of OA. _

In light of the systemic, but not unidirectional differences in DNA methylation on a genome wide scale that are apparent with OA progression in affected cartilage, it seems that systemically targeting DNA methylation for clinical purposes in OA is farfetched. Moreover, the widespread epigenetic differences that accompany OA suggestively affect expression of genes primarily involved in developmental processes, such as endochondral ossification. Aside from the substantial scientific challenge to locus specifically modulate DNA methylation, it seems more pragmatic to directly address the mRNA or protein molecules of the respective genes and pathways. While this is a difficult task on its own, specifically and locus specifically targeting DNA methylation in our view will unlikely serve a clinical purpose for OA in the near future.

However, aside from the putative limited role of epigenetics in curing OA, it may serve as an important biomarker when measured in clinically available tissues. As has also been shown for a number of molecular markers in blood or serum (62-64), it is apparent that transcriptomic data can purposely be used to identify symptomatic OA patients using mRNA extracted from peripheral blood mononuclear cells (60). For example, the blood transcriptomic profiles of OA patients are enriched for genes involved in apoptosis, which were subsequently shown to reflect the pathophysiological state of the articular cartilage (60). Hence, in light of the responsiveness of the epigenome to environmental changes and its relationship with gene expression it is expected epigenetics can fulfil a similar purpose. Moreover, hereby not only clinical associations can be constructed, but also important insight is given into the complex disease process. Preliminary results from our own group indicate that indeed DNA methylation might serve as a powerful biomarker for OA progression. More specifically, as little as four CpGs were needed to distinguish fast progressing from non-progressing OA patients with 76% accuracy (65).

It should be noted that we have here primarily discussed DNA methylation in articular cartilage in the context of OA, not in the last place due to the fact that degradation of articular cartilage is the primary feature of OA. However, it is clear that other joint tissues are also involved in the disease process as a whole. Other disease relevant tissues, such as the synovium (66-68) and to some extent the subchondral bone (69), have been addressed in light of other musculoskeletal or rheumatic pathologies and indicate compelling prospects for OA research. In this regard, preliminary data from a small study revealed DNA methylation differences between subchondral bone adjacent to varying degrees of damaged articular cartilage of knee OA patients, indicating that epigenetic regulation is likely involved in the pathophysiological interplay between subchondral bone and articular cartilage (69).

Considering the proposed subgroups of OA patients, the initial genome wide DNA methylation studies discussed here have reported on putative stratification of OA patients based on inflammatory profiles present in articular cartilage only. However, given the increasingly important role of secondary tissues in OA, it is possible that subgroups of patients can be defined by genome wide DNA methylation profiling of those respective tissues. Consequently, the OA research field might elucidate other proposed OA patient subgroups, such as distinct differential estrogenic responses advocated by Herrero-Beaumont *et al.* (59).

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