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Thyroid hormone signalling in Osteoarthritis: early life events in late life disease

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Chapter 4

The effect of forced exercise on knee-joints in *Dio2*^{-/-}-mice; type II iodothyronine deiodinase-deficient mice are less prone to develop OA-like cartilage damage upon excessive mechanical stress

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Abstract

Objective: To further explore deiodinase iodothyronine type-2 (*DIO2*) as a therapeutic target in osteoarthritis by studying the effects of forced mechanical loading on *in vivo* joint cartilage tissue homeostasis and the modulating effect herein of *Dio2* deficiency.

Methods: Wild-type and C57BL/6-*Dio2*^{-/-}-mice were subjected to a forced running regime for one hour per day for three weeks. Severity of osteoarthritis was assessed by histological scoring for cartilage damage and synovitis. Genome wide gene expression was determined in knee-cartilage by microarray analysis (Illumina MouseWG-6 v2). STRING-db analyses were applied to determine enrichment for specific pathways and to visualize protein-protein interactions.

Results: In total, 158 probes representing 147 unique genes showed significantly differential expression with a fold-change ≥ 1.5 upon forced exercise. Among these are genes known for their association with OA (e.g. *Mef2c*, *Egfr*, *Ctgf*, *Prg4* and *Ctnnb1*), supporting the use of forced running as an OA-model in mice. *Dio2*-deficient mice showed significantly less cartilage damage and signs of synovitis. Gene expression response upon exercise between wild-type and knockout-mice was significantly different for 29 genes.

Conclusion: Mice subjected to a running regime have significant increased cartilage damage and synovitis scores. Lack of *Dio2* protected against cartilage damage in this model and was reflected in a specific gene expression profile, and either mark a favorable effect in the *Dio2*-knockout (e.g. *Gnas*) or an unfavorable effect in wild-type cartilage homeostasis (e.g. *Hmbg2* and *Calr*). These data further support *DIO2* activity as a therapeutic target in OA.

Introduction

Osteoarthritis (OA) is a prevalent, complex, chronic and disabling disease of articular joints, characterized by progressive destruction of joint cartilage, remodeling of the subchondral bone, formation of osteophytes and synovitis [1, 2]. It causes pain and disability to an increasing proportion of the population and is associated with the obesity pandemic, aging of the population and not in the least by improved survival of patients with cardiovascular or oncological health problems. All together this imposes a large and growing social and economic burden [3, 4]. The development of novel therapeutic approaches is therefore urgently needed and should be based on insights into the underlying disease mechanisms [5-7]. Several genetic studies identified robust signals for OA susceptibility [8-13], that suggest specific genes, involved in cartilage development and growth, to play a key role in the OA disease process [14, 15]. Notable example of such OA susceptibility alleles, are the C-variant of the rs225014 single nucleotide polymorphism (SNP) located in the coding region of the deiodinase iodothyronine type-2 (D2) gene (*DIO2*) [8, 16] and the rs945006 SNP in the deiodinase iodothyronine type-3 (D3) gene (*DIO3*), another deiodinase with a counter regulatory function for *DIO2* [16]. For together, D2 and D3 primarily regulate the bio-availability of intracellular thyroid hormone in specific tissues such as the growth plate, but not systemically. The deiodinase type 2 protein (D2) catalyzes the conversion of intracellular inactive thyroxine (T4) to active thyroid hormone (T3). During skeletal development, this conversion plays a critical role in the process of endochondral bone formation by facilitating terminal maturation of hypertrophic chondrocytes subsequently leading to breakdown of the cartilage matrix and replacement by bone. This process is essential in skeletal development and growth but loss of the chondrocyte's maturational arrested characteristics is considered deleterious for postnatal articular cartilage.[17, 18] *DIO2* mRNA and D2 protein levels are highly upregulated in human osteoarthritic cartilage as compared to healthy cartilage [19-21], suggesting that in disease *DIO2* contributes to the loss of the highly specialized maturational arrested state of articular chondrocytes [17]. Cartilage-specific over-expression of human *DIO2* in rats was associated with increased damage to the articular cartilage in a surgical OA model. However, this was without clear evidence that hypertrophy of chondrocytes plays an essential role and rather pointing towards increased tissue destructive enzyme activity and enhanced expression of IL-1 target genes [22]. Upregulation of *DIO2* expression in a human *in vitro* model resulted in a marked reduction of the capacity of chondrocytes to deposit ECM components, including type II and type X collagen, while inducing OA-specific markers of cartilage matrix degeneration and mineralization [18]. In contrast, pharmacological inhibition of *DIO2* increased the expression of collagens and aggrecan without clear effect on hypertrophy or tissue destructive enzymes. These accumulating data suggest that D2 inhibition and/or modulation may become a therapeutic target, but the *in vivo* impact of D2 loss of function in joint biology and disease remains largely unknown in particular at the molecular level.

In this study, we set out to study the molecular network of *Dio2* in the healthy and challenged joint. We performed genome wide expression analyses in aging wild-type

and *Dio2*^{-/-} mice, including groups exposed to a moderately strenuous running regime. Our results indicate that *Dio2* is effectively involved in specific gene networks that can be associated with osteoarthritis and provide further insights into the complex molecular interactions involved in healthy and diseases articular cartilage.

Materials and methods

Animal experiments

Dio2^{-/-} mice were a kind gift of Dr. V. Galton (Dartmouth Medical School, NH, USA)[23] and were backcrossed onto the C57Bl/6 background. All experiments were approved by the Ethics Committee for Animal Research (KU Leuven, Belgium).

Four to 6 months old male *Dio2*^{-/-} (n=22) and wild-type mice (n=30) ran for 3 weeks 1 hour/day, 5 days/week, at a speed of 11 m/min and with an inclination of 5°. For additional details, see the Online Methods.

Histological assessment of osteoarthritis

Right knees were fixed overnight at 4°C in 2% formaldehyde, decalcified for 3 weeks in 0.5M EDTA pH 7.5 and embedded in paraffin. Severity of disease was determined by histological scores on hematoxylin/eosin or Safranin O stained sections (5µm) throughout the knee (5 sections at 100µm distance). For additional details on histological assessment and statistical analyses, see the Online Methods.

RNA isolation

Snap frozen cartilage of the left knees was powderized using a Retsch Mixer Mill 200 under cryogenic conditions. RNA was isolated and washed using the RNeasy mini kit (Qia-gen, Venlo, The Netherlands) according to the manufacturer's protocol. RNA quality was assessed using a Biocore lab-on-a-chip and quantity was assessed using a Nanodrop spectrophotometer. For additional details, see the Online Methods.

Microarray analysis

Complementary DNA synthesis, amplification, biotin labeling and hybridization onto the microarrays was performed using the Ambion TotalPrep-96 RNA amplification kit (Life Technologies, Bleiswijk, The Netherlands) according to manufacturer's protocol. After hybridization on Illumina MouseWG-6 v2 BeadChip microarrays (Illumina, Eindhoven, The Netherlands) the slides were scanned with the Illumina Beadscanner 500GX. For additional details on microarray handling and data analyses, see the Online Methods.

Pathway analysis and protein-protein interaction networks

Gene enrichment among the genes with significant differential expression was performed with STRING (Search Tool for the Retrieval of Interacting Genes/Proteins 9.1) [24]. Pathways with a P-value ≤ 0.05 after FDR correction were considered significant. Enrichment in protein-protein interactions was also analyzed using the STRING database. For addi-

tional details, see the Online Methods

Quantitative RT-PCR assays (validation)

Validation of the microarray results was performed by quantitative real-time PCR. 500 ng of total RNA was processed with the First Strand cDNA Synthesis Kit according to the manufacturer's protocol (Roche Applied Science, Almere, The Netherlands) upon which cDNA was diluted 5 times. RT-qPCR measurements were performed on the Roche Lightcycler 480 II, using Fast Start Sybr Green Master reaction mix according to the manufacturer's protocol (Roche Applied Science). For additional details, see the Online Methods

Results

Reduced severity of OA after forced running in *Dio2*^{-/-} mice

To study the role of *Dio2* in joint homeostasis, *Dio2*^{-/-} and wild-type mice were studied in a forced-exercise setup and compared to non-running mice (Figure 1). No striking developmental skeletal phenotype appears present in the *Dio2*^{-/-} mice [25] and our observations in our mouse colony are in agreement with these findings. Wild-type and *Dio2*^{-/-} mice displayed a similar running behavior. As shown in Figure 1, overall group analysis indicated a significant difference between wild-type- and knockout-mice with respect to cartilage damage ($P = 0.0006$) and synovial hyperplasia ($P = 0.0536$). No specific effects within wild-type- or knockout-mice of exercise were found, nor interaction between genotype and exercise. Sidak's multiple comparison test indicated that the effect was determined by the difference in genotype in the exercise group ($P < 0.0001$ for cartilage damage – 95% CI of the difference between the means (0.1820 to 0.5820) and $P = 0.0184$ for synovial hyperplasia – 95% CI of the difference between the means (0.03661 to 0.4627)) (Figure 1).

Microarrays: Differential expression in knee-joints upon forced exercise

To identify genes responsive to the forced running regime in mice, genome wide gene expression in knee-cartilage was studied by microarray. Gene expression was detected before and after the running regime in interaction with the genetic background (wild-type and *Dio2*^{-/-}-mice). Figure 2 shows a schematic overview of the study strategy.

After quality control and normalization, 20872 of the 45281 probes of the BeadChip array were used for analyses. Microarrays on human articular cartilage samples showed corresponding numbers for specific expression [26]. In order to detect all articular cartilage genes that are responsive to the applied running regime, we performed differential expression analyses in 3 strata: total (A), *Dio2*^{-/-} (B) and wild type (C) group (Figure 2). For each strata significance was adjusted for multiple testing according to the “Benjamini and Hochberg” method. In the total and wild type strata, we independently detected respectively 1862 and 892 probes representing 1699 and 830 genes (558 overlapping) that were significantly differentially expressed between the forced exercise and the control group. In contrast, in the *Dio2*^{-/-}-stratum, this comparison did not result in any significantly differentially expressed probes after multiple testing. Among the differentially ex-

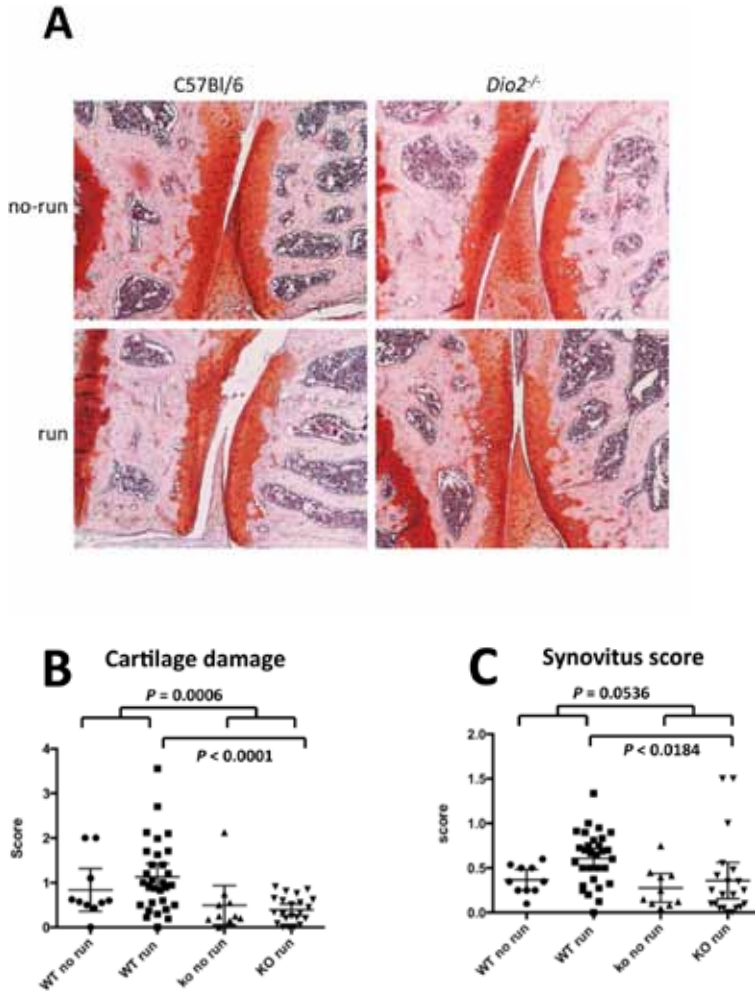


Figure 1. Histological scoring of osteoarthritis.

Comparing wild-type and *Dio2^{-/-}* mice as well as the effect of forced running. (A) Frontal Haematoxylin-Safranin O stained sections of C57Bl/6 wild-type and *Dio2^{-/-}* knees (medial) of mice subjected to a running regime (run) and control mice (no run) (magnification 10x). (B) Cartilage damage was increased in wild-type mice as compared to *Dio2^{-/-}* mice (2-way ANOVA $p = 0.0006$) and (C) a similar trend was observed for synovitis (2-way ANOVA $p = 0.0536$). None of the other comparisons (e.g. 'WT no run' versus 'KO no run') were found significantly different. Data are shown as individual values, mean and 95% confidence intervals.

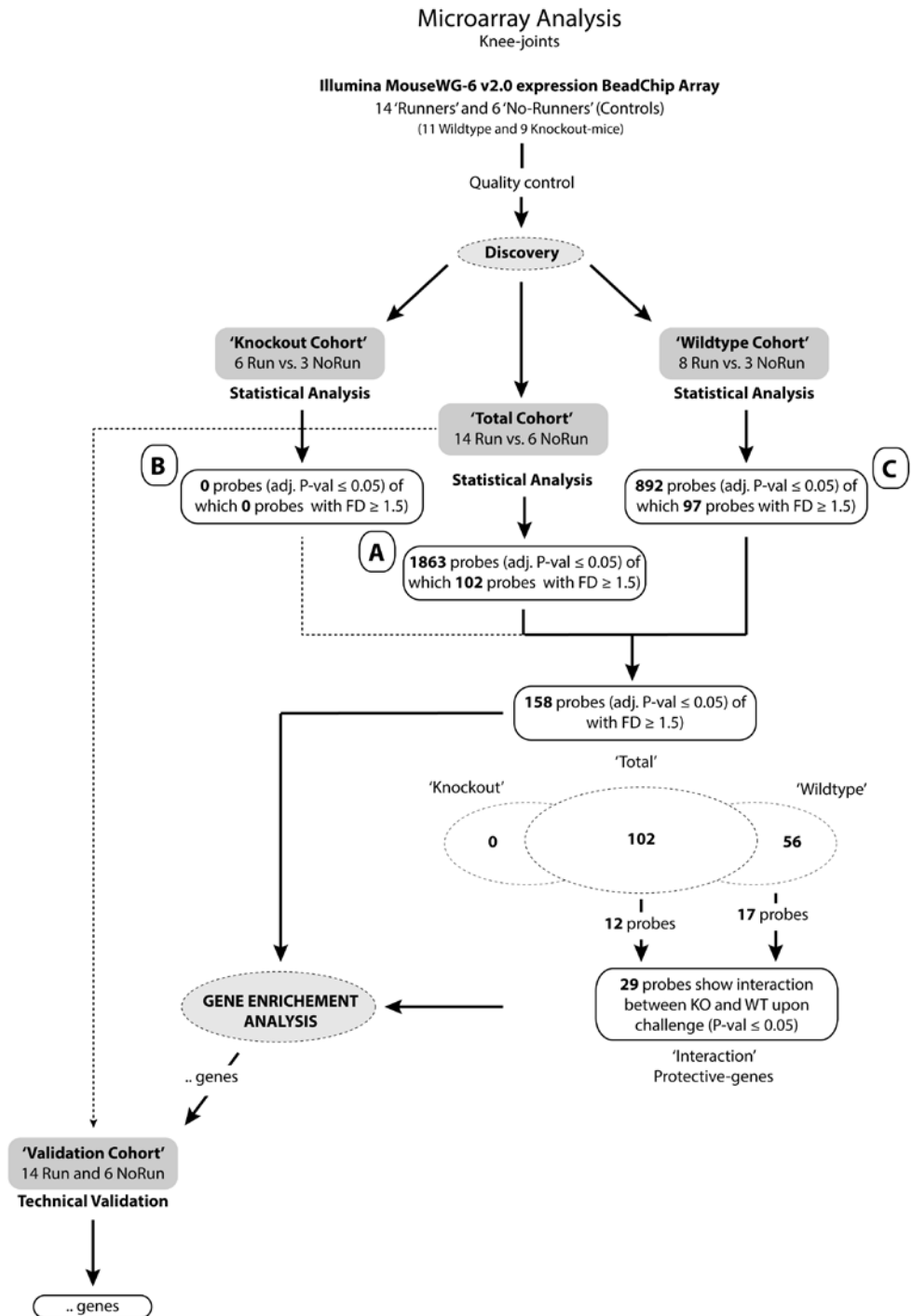


Figure 2. Overview of the study strategy.

Following microarray analysis, data were statistically analyzed. Secondary analyses using gene enrichment analysis and interaction analysis will lead to a set of genes which will be technically validated.

pressed probes in the total and wild-type groups, we observed respectively 102 and 97 probes with a fold-change of 1.5 and higher that together consisted of 158 significant unique probes, representing 147 unique transcripts that were responsive to the running regime (Figure 2 and Supplement Table S1). Notably, of the 158 differentially expressed probes, 31 were up-regulated (20%) and 127 showed down-regulation (80%). Among the 147 unique transcripts, we observed genes known for a potentially role in cartilage homeostasis and disease, such as Proteoglycan 4, also known as Lubricin (*Prg4*, 1.95-fold down; $P=4.44 \times 10^{-2}$)[27], myocyte enhancer factor 2C (*Mef2c*, 1.9-fold down; $P=3.05 \times 10^{-2}$)[28, 29] and connective tissue growth factor (*Ctgf*, also known as *Ccn2*, 1.79-fold down; $P=1.32 \times 10^{-2}$)[30].

Technical validation of our microarray results was carried out by RT-qPCR in the discovery cohort (20 samples previously included in the microarray analyses). Genes to be validated were selected based on P-value and fold difference (differential reaction on mechanical stress). Here, 19 out of the 20 genes tested showed similar effect sizes and direction as the original data, only *Pfn1* did not show similar effect of expression (see Supplementary Table S2).

Pathway analyses: Protein-protein interactions and gene enrichment analyses

To identify functional connections between the genes that were found as differentially expressed (Table S1) we determined the protein-protein interaction prediction using STRING-db. Analyzing the 147 differentially expressed genes, representing 124 unique proteins, showed enrichment of interaction in the subset of proteins ($P=1.61 \times 10^{-4}$; 63 interactions; Figure S1), indicating that the gene-products that were identified are closely interacting as a response to the forced running regime.

Furthermore, we assessed whether the differentially expressed genes occurred more frequently in a specific pathways in mice. Gene enrichment analyses revealed enrichment for biological processes (GOTERM_BP_FAT) concerning, amongst others, “regulation of metabolic process” ($P=0.0063$; GO:0019222; N=46, e.g. *Calr*, *Igfbp5*, *Sox4* and *Tcf4*), “skeletal system development” ($P=0.00395$; GO:0001501; N=12, e.g. *Ctnnb1*, *Ctgf*, *Sox4* and *Mef2c*), “anatomical structure development” ($P=0.039$; GO:0048856; N=36, e.g. *Sox4*, *Notch3*, *Egfr*, *Ctnnb1* and *Mef2c*), and “regulation of response to stress” ($P=0.0171$; GO:0080134; N=14; *Setd8*, *Ankrd1*, *Mef2c*, *Egfr* and *Ctgf*) with application of a false discovery rate algorithm.

To connect thyroid hormone signaling with the effects of forced mechanical stress, we added genes involved in intracellular thyroid hormone signaling (*Dio2*, *Thra*, *Thrb* and *Rxra*) into the protein-protein interaction prediction. The network that is formed (Figure 3) shows thyroid signaling genes/proteins to be incorporated into the large network, as previously seen (Figure S1). Thyroid signaling was found to interact directly with the differential expressed genes through, *Ctgf* (A) and *Egfr* (B), via thyroid receptor alpha (*Thra*) and retinoid x receptor (*Rxra*), both known factors in the development of OA[30-32].

Genes differentially expressed in *Dio2*^{-/-} mice compared to wild type mice upon a forced running regime

Since we found differences in differential expression patterns between knockout and wild

type mice upon the forced running regime, we assessed which of the 147 differentially expressed genes (Figure 1 and Supplementary Table S1) showed significant interaction between running regime and genetic background based on nominal P-values. In total 29 probes, representing 29 genes were found to be significantly differentially expressed in knockout-mice as compared to wild type-mice when undergoing a forced running regime (Table 1). The significant differences in effect (beta-values), as a result of the running regime, between the knockout- and the wildtype-group can be divided in 3 sub-groups. Each group shows a different effect-size based on “genotype” for the 29 genes found to show a significant interaction. We identified 4 genes that showed no differential expression upon running in wild-type mice, but are differentially expressed (down-regulated) upon running in *Dio2*^{-/-} mice (Group 1; e.g. *Gnas* and *Rhbdl2*). In total 16 genes only showed differential expression in wild-type mice and not in the *Dio2*^{-/-} mice (Group 2; e.g. *Hmgb2*, *Calr* and *Lbh*) whereas, 9 genes showed significant differential effect-sizes between wild-type and knockout mice, but having the same direction of effect (Group 3; e.g. *Sox4* and *Socs2*). Depending on the gene-expression pattern the genes in the different groups could mark a favorable effect in the *Dio2*-knockout mice (Group 1) or an unfavorable effect in wild type cartilage homeostasis (Group 2).

Discussion

Dio2-deficient mice showed less cartilage damage and reduced severity of synovitis in a treadmill running model of OA. The absence of significant differential gene expression between the running and no-exercise group in *Dio2*^{-/-} mice suggests that degenerative pathways are not activated in this knockout strain despite the biomechanical burden that is imposed. These data provide novel support for inhibition of *DIO2* as a therapeutic strategy in OA, in particular since no striking developmental skeletal phenotype appears present in the *Dio2*^{-/-} mice (results not shown). Upon forced mechanical loading, wild type mice showed clear signs of OA and differential expression of genes associated with the disease, supporting the use of forced running as an OA-model in mice. A subset of the genes was found directly interacting with thyroid signaling through *Thra*, *Rxra* and *Dio2*, depicted in Figure 3. This indicates the importance of thyroid hormone signaling as a regulatory system in the response to stress and, when suppressed, for the maintenance of cartilage homeostasis.

With the differentially expressed genes in the wild-type stratum and the combined knockout/wild-type stratum, pathway enrichment was found for expected biological processes, such as “skeletal system development”, but also for processes involved in “regulation of response to stress”. Notably, genes overlapping between these enriched processes as well as being ‘nodes’ in the recognized protein interaction networks (*Egfr* (B) and *Ctnnb1*(C); Figure 3) are well known for their association with OA [30, 33, 34]. These could, therefore, point at important modulators affecting the propensity to develop OA upon mechanical stress and as such at potential drugable targets for novel therapeutic approaches.

Tabel 1: Genes that show significant differential expression upon the running regime between wild-type- and knockout-mice (interaction)

Symbol	Run vs NoRun in Knockout			Run vs NoRun in Wildtype			Interaction
	Beta ¹	St. Error ²	P-value ³	Beta	St. Error	P-value	P-value ⁴
Group 1: Genes with an effect of the running regime in knockout- but not in wild-type-mice							
Hist1h2an	-1,024	0,2603	0,000	-0,186	0,1088	0,087	0,0030
Gnas	-0,956	0,2318	0,000	-0,258	0,1473	0,080	0,0111
Rhbdl2	-1,135	0,2176	0,000	-0,349	0,2638	0,185	0,0215
E130112E08Rik	-1,041	0,2981	0,000	-0,26	0,2171	0,231	0,0343
Group 2: Genes with an effect of the running regime in wild-type- but not in knockout-mice							
Anxa11	0,02	0,0852	0,818	-0,636	0,0668	0,000	0,0000
Setd8	-0,006	0,0864	0,943	0,725	0,1044	0,000	0,0000
Phdla1	-0,152	0,1019	0,136	-0,591	0,1011	0,000	0,0022
Hmgb2	0,131	0,1439	0,363	0,666	0,1476	0,000	0,0095
Calr	-0,086	0,1561	0,581	-0,664	0,1658	0,000	0,0111
Gdi1	0,313	0,2297	0,173	0,923	0,1446	0,000	0,0247
Lbh	-0,13	0,0682	0,057	-0,889	0,0761	0,000	0,0000
Fmo5	-0,028	0,1028	0,784	-0,622	0,0848	0,000	0,0000
Igsf4a	-0,171	0,0978	0,081	-0,789	0,1221	0,000	0,0001
Fcho1	0,132	0,1144	0,250	0,665	0,1106	0,000	0,0008
Serpinb1a	0,396	0,3687	0,283	-0,877	0,1145	0,000	0,0010
Mtpn	-0,082	0,1084	0,451	-0,594	0,1121	0,000	0,0010
Rbm47	-0,09	0,075	0,229	-0,592	0,1011	0,000	0,0013
LOC433464	-0,086	0,1638	0,599	-0,675	0,0788	0,000	0,0015
Rer1	-0,185	0,1714	0,281	-0,611	0,1121	0,000	0,0195
Hist1h4i	0,113	0,2763	0,684	-0,811	0,1011	0,000	0,0214
Group 3: Genes with an effect of the running regime in both wild-type- and knockout-mice							
Mll5	-1,378	0,2306	0,000	-0,439	0,1184	0,000	0,0003
Chd1	-1,178	0,2311	0,000	-0,307	0,1125	0,006	0,0007
Septin11	-0,28	0,0887	0,002	-0,671	0,0918	0,000	0,0022
Sox4	-0,479	0,1399	0,001	-0,969	0,1064	0,000	0,0053
Gins1	0,771	0,305	0,011	1,596	0,0735	0,000	0,0085
Socs2	-0,337	0,1132	0,003	-0,713	0,0881	0,000	0,0088
C330023M02Rik	-0,308	0,1498	0,040	-0,678	0,054	0,000	0,0202
Hgsnat	-0,29	0,1274	0,023	-0,621	0,1001	0,000	0,0410
Rap2c	-0,385	0,0743	0,000	-0,628	0,0955	0,000	0,0449

¹The beta-value is representing the difference in fold change between the running group and the non-running group. The direction of the beta-value is how the running-group is different from the non-running group. ²The standard error is showing the variation of the beta-value between samples.

³The nominal P-value is the P-value of that the single probe that was calculated to tell whether based on this single probe, the expression levels between runners is different of that of non-runners.

⁴The Interaction P-value is a summarizing value that tells whether the difference caused by exercise is significantly different between the knockout- and the wild-type-stratum

In contrast, when assessing gene expression in the knockout-group alone, we found no significant differential gene expression upon severe mechanical loading, after multiple testing adjustment. Taken together with the pathology observations these data indicate that the repression of *Dio2* is beneficial against the development of cartilage damage upon mechanical stress. This effect could theoretically also be caused by a difference in power between the different strata (wild-type and knockout). However, since the comparison made in knockout animals (6 running versus 3 controls) is comparable to the wildtype animals (8 running versus 3 controls), this is unlikely to explain the complete absence of a significant differentially expressed gene in the knockout stratum.

Looking at the effect of *Dio2*-deficiency on gene expression, we found that the gene expression response upon exercise between wild-type and knockout-mice was significantly different for 29 genes with nominal significance, of the total set of 147 differentially expressed genes. Genes categorized in group 1 (Table 1), showing no differential expression upon forced running in wild-type mice, but are differentially expressed with nominal significance in *Dio2*^{-/-} mice (*E130112E08Rik*, *Gnas*, *Hist1h2an* and *Rhbdl2*) could represent the respective favorable effects. *Gnas*, for example, is involved in skeletal development (GO:0001501) and complex skeletal disorders such as Albright Hereditary Osteodystrophy that can lead to early OA [35]. *Gnas* upregulation was found directly regulating hypertrophic differentiation of growth plate cartilage *in vivo* [36]. The knockout-specific down-regulation of *Gnas* upon forced running could be a protective mechanism within the *Dio2* knockout mouse.

We hypothesize that the sixteen genes within group 2, only showing differential expression in wild-type mice and not in the knockout-group (e.g. *Hmgb2* and *Calr*), represent the unfavorable, damaging effects of forced running in wild type mice. *Hmgb2* was shown to be expressed at higher levels in human MSCs as compared to human articular chondrocytes and the expression declined during chondrogenic differentiation of MSCs [37]. The up-regulation of *Hmgb2* seen in wild type-mice upon forced running could be explained as a marker that changes in the differentiation status of the chondrocytes are occurring. A second interesting member of this group is calreticulin (*Calr*). *Calr* interacts with the glucocorticoid receptor (GR). It may also interact with other steroid receptors or thyroid receptors in a similar way [38]. Furthermore, *Calr* was shown to be involved in cartilage thinning of mandibular cartilage in a rat model that studied the effects of compressive mechanical loading [39]. In our forced running experiment we see a similar induction of *Calr* expression upon stress, but not in the *Dio2* knockout-mice, possibly giving the knockout-mouse an advantage against the formation of OA-like degradation or cartilage thinning.

Nine additional genes showed significant differential effect-sizes between wild-type and knockout mice, albeit the same direction of effect (e.g. *Sox4* and *Socs2*). Notable within this third group (Table 1) is *Sox4*, which was previously found to be expressed very early during chondrogenesis [40], much earlier than the well defined *Sox5* and *Sox6* [41]. Furthermore, it was shown that expression of *Sox4* could be stimulated by adding physiological concentrations of human parathyroid hormone (PTH), indicating involvement of the PTH/PTHrP receptor [42]. Here we show that *Sox4* expression is influenced by the ab-

Figure 3. Gene networks in search tool for the retrieval of interacting genes.

Screenshot of the protein–protein interactions (STRING-db) for the 147 genes differing >1.5-fold between ‘runners’ and controls, including thyroid signaling members; *Dio2*, *Thra*, *Thrb* and *Rxra* (encircled by the dashed line). Mayor hubs in the network are depicted by A) *ctgf*, B) *egfr*, and C) *ctnnb1*. Disconnected proteins are hidden.

sence of *Dio2*. In the knockout-mice the expression of *Sox4* is 2 times less down-regulated upon running when compared with the control group. How this influences the structural integrity of articular cartilage remains unknown. Other genes in the list have no straight-forward connection to OA based on current literature.

In conclusion, in the current paper we find that *Dio2* deficiency has a protective effect on the homeostasis of articular cartilage in the knee-joints of mice undergoing a forced running regime. This is consistent with our earlier findings, showing that pharmacological inhibition of deiodinases in a human *in vitro* chondrogenesis model has a beneficial effect on the early formation and maintenance of articular cartilage ECM [18]. It is therefore hypothesized that control of thyroid hormone signaling, both during development and adult cartilage maintenance, is essential to ensure normal bone and cartilage homeostasis, and that it could act as the master-switch that forces maturational arrested chondrocytes to re-activate the endochondral ossification process, leading to articular cartilage destruction. Our results show that interfering with intracellular thyroid hormone levels could be a powerful way to oppose the pathological events that are occurring in OA.

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Competing interest

The authors declare no competing financial, personal, or professional interests.

Author contributions

Conceived and designed the experiments: NB; FMFC; PES; IM; RJLL

Performed the experiments: NB; FMFC; LS; RvdB; NL;

Analyzed the data: NB; FMFC; YFMR; WdH; IM; RJLL

Wrote the manuscript: NB; FMFC; IM; RJLL

Critically reviewed the manuscript: All authors

Supplemental Data

<http://ard.bmj.com/content/75/3/571/suppl/DC1>

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