Insights into the genetic architecture of osteoarthritis from stage 1 of the arcOGEN study

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Concise report

Insights into the genetic architecture of osteoarthritis from stage 1 of the arcOGEN study


ABSTRACT

Objectives The genetic aetiology of osteoarthritis has not yet been elucidated. To enable a well-powered genome-wide association study (GWAS) for osteoarthritis, the authors have formed the arcOGEN Consortium, a UK-wide collaborative effort aiming to scan genome-wide over 7500 osteoarthritis cases in a two-stage genome-wide association scan. Here the authors report the findings of the stage 1 interim analysis.

Methods The authors have performed a genome-wide association scan for knee and hip osteoarthritis in 3177 cases and 4894 population-based controls from the UK. Replication of promising signals was carried out in silico in five further scans (44449 individuals), and de novo in 14534 independent samples, all of European descent. The stage 1 genome-wide association scan.

Results None of the association signals the authors identified reached genome-wide levels of statistical significance, therefore stressing the need for corroboration in sample sets of a larger size. Application of analytical approaches to examine the allelic architecture of disease to the stage 1 genome-wide association scan data suggests that osteoarthritis is a highly polygenetic disease with multiple risk variants conferring small effects.

Conclusions Identifying loci conferring susceptibility to osteoarthritis will require large-scale sample sizes and well-defined phenotypes to minimise heterogeneity.

Osteoarthritis is the most common form of arthritis affecting 40% of people over the age of 70 years, and is associated with a substantial health economic burden. Osteoarthritis is thought to be caused by a complex interplay between environmental and genetic factors. As with many common complex disorders, the genetic architecture of osteoarthritis has not yet been characterised. Over the past decade, candidate gene association studies and genome-wide linkage scans have failed to identify robustly replicating osteoarthritis loci, with the notable exception of rs143383 in the GDF5 gene. A genome-wide association study (GWAS) recently reported a single novel locus associated with radiographically defined knee and/or hip osteoarthritis on chromosome 7q22 (rs3815148 in the COG5 gene), subsequently corroborated by a genome-wide association scan across four studies including the discovery set. Both established that osteoarthritis loci are represented by common variants (>0.2 minor-allele frequency), have small effect sizes (allelic OR ~1.15), and reach genome-wide significance (p<5×10−8). These associations have characteristics typical of common complex traits and require large sample sizes for their detection. To enable a well-powered GWAS for osteoarthritis, we formed the arcOGEN Consortium, a UK-wide collaborative effort aiming to scan genome-wide over 7500 osteoarthritis cases in a two-stage genome-wide association scan. Here we report the findings of our GWAS stage 1 analysis and replication studies, and describe the outcomes of statistical analyses designed to model the genetic architecture of osteoarthritis.

METHODS

An expanded description of the methods is provided in supplementary methods (available online only). The stage 1 genome-wide association scan included 3177 knee and/or hip osteoarthritis cases from the UK ascertained based on radiographic evidence of disease (Kellgren–Lawrence grade ≥2) or clinical evidence of disease to a level requiring joint replacement. Cases were genotyped using the Illumina Human610 platform (Illumina, San Diego, California, USA). We used 4894 early-access publicly
RESULTS

In our stage 1 genome-wide association scan analysis, we observed a slight excess of associations compared with the null distribution for the joint and gender-stratified analyses (supplementary results; supplementary figure 3, available online only). The genomic control inflation factor \(\lambda\) was 1.077, in keeping with other UK-based GWAS.\(^6\) Although there was no signal exceeding genome-wide significance (\(p<5\times10^{-8}\)), 89 SNP reached \(p\) values of less than \(10^{-4}\) (as opposed to 51 expected under the null, binomial \(p=10^{-6}\)).

The strongest statistical evidence for association with osteoarthritis was obtained for rs4512591 on chr8 (\(OR\) for allele C 1.17; 95% CI 1.10 to 1.25; \(p=1.8\times10^{-6}\)) approximately 66 kb upstream of the \(TRIB1\) gene. For knee osteoarthritis the most significant finding was also observed at rs4512891 (\(OR\) for allele C 1.25; 95% CI 1.13 to 1.38; \(p=1.1\times10^{-6}\)); for hip osteoarthritis, the strongest signal was rs4977469 (\(OR\) for allele A 1.30, 95% CI 1.17 to 1.45; \(p=1.2\times10^{-6}\)), within intron 3 of the predicted \(FAM154A\) gene (supplementary figure 1; supplementary figure 4; supplementary table 2, all available online only). Following replication studies in up to 56 917 independent European-ancestry samples, the overall statistical evidence for association was reduced, with the strongest signal for knee and/or hip osteoarthritis observed at rs2277831 on chr22 (\(OR\) for allele G 1.07, 95% CI 1.04 to 1.11; combined \(p=2.3\times10^{-5}\)), within intron 32 of the \(MICAL3\) gene. For knee osteoarthritis the most significant finding post-replication was observed at rs11280 (\(OR\) for allele C 1.10, 95% CI 1.05 to 1.16; \(p=3.2\times10^{-5}\)), within \(Cebpe\); for hip osteoarthritis, the strongest signal was at rs2615977 (\(OR\) for allele A 1.10, 95% CI 1.05 to 1.15; \(p=1.1\times10^{-5}\)) within intron 31 of the \(COL11A1\) gene (supplementary figure 1; supplementary figure 4; supplementary table 2, all available online only). Out of the 65 independent signals (composed of 89 SNP) with \(p<10^{-4}\) in all the osteoarthritis analysis, 17 are present with \(p<10^{-4}\) in the hip osteoarthritis analysis and nine are present with \(p<10^{-4}\) in the knee osteoarthritis analysis, but none reach \(p<10^{-4}\) in both.

Additional analyses, including testing for pairwise interactions, examining overlap between genome-wide association scan and linkage scan signals, association with body mass index, deviation from the additive model, genome-wide HapMap-based imputation, chromosome X association and genome-wide low frequency/rare variant analysis did not identify any additional osteoarthritis signals (supplementary methods; supplementary results, available online only).

The arcOGEN stage 1 GWAS is well powered to detect association with common variants of modest effect at the genome-wide significance level (eg, 90% power to detect an allelic \(OR\) of 1.25 at a SNP with frequency 0.55). However, it is poorly powered to detect effects at the established osteoarthritis variants (8% and 4% power for \(C6F5\) and \(7q22\), respectively), as the index SNP (rs143383, risk allele frequency 0.67; rs851548, risk allele frequency 0.23) both have small effect sizes (\(OR\) ~1.15).\(^2\)

Retaining the arcOGEN case–control ratio of 1.54, 7774 cases would be required to achieve 80% power to detect an allelic \(OR\) of 1.15 at a SNP with risk allele frequency 0.67, and 9084 cases would be required to achieve the same power to detect the same effect size at a SNP with risk allele frequency 0.23.

To evaluate the robustness of association signals, we performed sensitivity analyses using different control sets. We first used ‘supercontrols’ (hip and knee osteoarthritis-free individuals) and found high correlation between association effect estimates (>0.88) and 95% concordance in the direction of effects (supplementary results, available online only). The two established osteoarthritis loci demonstrated stronger evidence for association compared with the main analysis, even though the ‘supercontrol’ sample size was smaller. We also used a subset of the population-based controls, genotyped on a different platform, to assess robustness to the typing method. We observed 100% concordance in the direction of effect and high correlation between estimates of the \(OR\) (\(r=0.94\)) (supplementary results, available online only). Therefore, the choice of controls may have affected association strength but not the direction of effect.

The genetic architecture of osteoarthritis is likely to be polygenic with multiple variants along the spectrum of allele frequencies contributing modest and small effects. Our polygene analyses support a model of osteoarthritis in which there is a substantial genetic component comprising multiple contributing variants with small effect sizes (figure 1). SNP with \(p\) values as high as 0.25 appear to contribute to the genetic component of osteoarthritis (empirical \(p=3\times10^{-5}\), based on ~85 million permutations), with the bulk of the contribution seen in the 0–0.1 \(p\) value range (empirical \(p=3.5\times10^{-5}\), ~101 million permutations). Evaluation of discreet \(p\) value bins corroborates this observation and supports a role for SNP with \(p\) values up to 0.25 (0.10\(\leq p<0.15\) bin empirical \(p=0.06\); 0.15\(\leq p<0.20\) bin \(p=0.0072\); 0.20\(\leq p<0.25\) bin \(p=0.022\)), but with the major contribution coming from SNP with \(p<0.10\) (0\(\leq p<0.05\) bin \(p=3\times10^{-5}\); 0.05\(\leq p<0.10\) bin \(p=0.0092\)). The estimated proportion of variance in disease state explained by the osteoarthritis score alleles is 3.05% (\(p=3.3\times10^{-4}\) based on nine million permutations). This is in keeping with findings in other complex common diseases.\(^7\)

DISCUSSION

Our stage 1 arcOGEN genome-wide association scan analysis results are in agreement with other large-scale genetic studies.\(^3\)
controls in GWAS. Our 'supercontrol'-based sensitivity analyses demonstrated the utility of population-based, rather than disease-free, disease. The red line represents stage 1 genome-wide association study data and shows that SNP at the tail of the p value distribution (up to p<0.25, in the single-nucleotide polymorphism (SNP) sets. The blue line represents expectation under the null hypothesis of no genetic component to the disease. The red line represents stage 1 genome-wide association study data and shows that SNP at the tail of the p value distribution (up to p<0.25, but primarily p<0.10) have a significantly higher case–control discriminatory capacity.

and clearly indicate that common SNP with large effect sizes are not likely to underpin the aetiology of osteoarthritis. The only two established osteoarthritis loci to date have small OR and our polygene analyses on common SNP suggest that the genetic architecture of osteoarthritis is likely to consist of numerous signals of similar magnitude. The significant increase in sample size with stage 2 of the arcOGEN genome-wide association scan (~2.4 times as many cases) will increase the power to detect osteoarthritis associations. In addition, large-scale international meta-analysis efforts are underway and will ensure maximal GWAS sample size. A further important parameter in enhancing power and increasing the chances of success involves the improved definition of phenotype. One of the reasons why replication of findings has been difficult to achieve in osteoarthritis could be the inherent heterogeneity of the different osteoarthritis diagnostic and study inclusion criteria used. The field is currently active in evaluating phenotype definition differences and their effects on study power.4

The Wellcome Trust Case Control Consortium first demonstrated the utility of population-based, rather than disease-free, controls in GWAS.5 Our ‘supercontrol’-based sensitivity analyses suggest that for a highly prevalent and heterogeneous disorder such as osteoarthritis, with multiple smaller effects contributing to overall susceptibility, the brute force approach of maximising sample size to balance misclassification in controls is not as successful as it has been for other common diseases, in which ‘low-hanging fruit’ discoveries (of loci with substantial effects) were robust to subtle allele frequency fluctuations.

Osteoarthritis is a heterogeneous disease characterised by variable clinical features with conceivably different genetic aetiologies.9 Although the allele score was associated with osteoarthritis, the proportion of disease variance explained cannot be highly accurately quantified, primarily because of signal attenuation (risk alleles are unlikely to be at the actual causal locus) and sampling variation. The genetic architecture of osteoarthritis is emerging as complex. Large-scale sample sizes and well-defined phenotypes will be required to gain a better understanding of this genetic component, possibly leading to optimising treatment, developing efficacious disease-modifying interventions, improving prognosis and tailoring intervention to the individual.

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Competing interests None.
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REFERENCES


8. The Wellcome Trust Case Control Consortium. Genome-wide association study of 14,000 cases of seven common diseases and 3,000 shared controls. Nature 2007;447:661–78.