Epidemiological studies have demonstrated that primary osteoarthritis (OA) has a major genetic component. A genome wide linkage scan conducted by our group disclosed suggestive linkage to six loci in a cohort of sibling pairs with OA ascertained by joint replacement of the hip or the knee. Stratification by sex and the replaced joint showed that most of the loci were particularly relevant to hip OA in women. Finer linkage mapping of our chromosome 4 linkage interval for plausible functional candidate genes. Three candidates were selected based upon a reported or suspected role in skeletal joint tissue biology. Three candidates were identified and chosen for further investigation (table 1).

Candidate genes
We were conscious of the fact that the chromosome 4q susceptibility gene could be positioned at some distance from the point of maximum linkage. We therefore searched a broad region of 4q for candidates, between markers D4S2971 (4q12, 53.6 Mb) and D4S3042 (4q21.1, 77.3 Mb). We searched the Ensembl (http://www.ensembl.org/, accessed 28 December 2004) and UCSC (http://genome.ucsc.edu/, accessed 28 December 2004) genome browsers. Three candidates were identified and chosen for further investigation (table 1).

PATIENTS AND METHODS
Patients for association analysis
Full clinical details of the 146 female-THR families have been published previously. Association was also tested in a second cohort comprising 244 female patients with hip OA. The probands and cases were ascertained through the Nuffield Orthopaedic Centre and had undergone total joint replacement of the hip only (mono- or bilateral) for primary OA. The primary status was supported by clinical, radiological, operative, and histological findings and we excluded any secondary forms of the disease or non-OA cases. The 375 female controls were from the general population. All patients and controls were aged 55 or over and were of UK Caucasian origin. Ethical approval for the study was obtained from the Oxfordshire Clinical Research Ethics Committee and informed consent was obtained from all subjects.

Reverse transcription-polymerase chain reaction (RT-PCR)
Articular cartilage biopsy specimens were collected from patients who had undergone joint replacement for OA. The nucleic acid was then extracted as described previously. cDNA was synthesised using random hexamers and served as the template for PCR using primers located in separate exons.

DNA variant detection and genotyping
The exons, the intron-exon boundaries, and the 5' and 3' untranslated regions of IGFBP7 (5 exons), ADAMTS3 (22 exons), and IL8 (4 exons) were screened for common DNA variants (rare allele frequency >0.05) by the direct sequencing of 48 female-THR patients. The variants were genotyped by PCR-restriction enzyme analysis. Further details can be obtained from the corresponding author.

Abbreviations:
- ADAMTS, a disintegrin and metalloproteinase domain with thrombospondin motifs; CI, confidence interval; IGF, insulin-like growth factor; IGFBP, insulin-like growth factor binding protein; IL, interleukin; indel, insertion/deletion; OA, osteoarthritis; OR, odds ratio; RT-PCR, reverse transcription-polymerase chain reaction; SNP, single nucleotide polymorphism; THR, total hip replacement
Statistical analysis
Genetic association, and Hardy-Weinberg equilibrium for the distribution of genotypes, was tested by $\chi^2$ analysis with Yates's correction. Odds ratios were calculated with 95% confidence intervals. The pairwise linkage disequilibrium coefficient ($r^2$) was calculated using the GOLaD program (http://www.sph.umich.edu/csg/abecasis/GOLD/, accessed 28 December 2004). Haplotype frequencies between variants showing evidence of linkage disequilibrium at $r^2 > 0.2$ were estimated using the EH-PLUS program (http://www.iop.kcl.ac.uk/IOP/Departments/PsychMed/EPiBIST/software.stm, accessed 28 December 2004). Haplotype frequency differences were then compared using $\chi^2$.

RESULTS
Expression by OA chondrocytes and DNA variant detection
RT-PCR showed that the IGFBP7, ADAMTS3, and IL8 genes were expressed by OA articular cartilage chondrocytes (data not shown). Fourteen common DNA variants were identified in the three genes, comprising 12 single nucleotide polymorphisms (SNPs) and two insertion/deletion (indel) polymorphisms (table 2). Only one of the variants was exonic, the ADAMTS3 exon 3 SNP. This is an A to G transition coding for the substitution of the arginine at amino acid residue 138 by a lysine (Arg138Lys). This SNP is a member of the dbsNP database, with reference number rs788908 (http://www.ncbi.nlm.nih.gov/SNP/, accessed 28 December 2004). The six IL8 SNPs were all previously identified by Hull et al. 

Genotyping and association analysis
Only the ADAMTS3 SNP (−19) showed evidence for association in the 146 female-THR probands, with a $p$ value of 0.015. The rare allele of this SNP had a raised frequency in the probands (0.07) versus the controls (0.03). The odds ratio (OR) for the minor allele was 2.29 (95% confidence interval (CI) 1.22 to 4.29). Because this result could represent a false positive we genotyped the SNP in an independent cohort of 244 female-THR cases. We also genotyped these cases for the ADAMTS3 indel, which had approached significance in the female-THR probands, and for the ADAMTS3 exon 3 SNP, the only exonic variant. None of the three variants was associated in the 244 cases (table 3). The minor allele of the (−19) SNP was raised in the cases (0.05) compared with the controls (0.03), a pattern observed for the probands. When the probands and cases were combined and compared with the controls, there was a significant difference for the (−19) SNP. However, the $p$ value of 0.022 and the OR of 1.91 (95% CI 1.13 to 3.23) were not as significant as those obtained for the probands alone ($p = 0.015$, OR = 2.29). Increasing the sample size had reduced the significance of the association.

Of the six ADAMTS3 variants genotyped, three variant combinations had a pairwise $r^2$ value $> 0.20$: (−19)/(−244), with an $r^2$ of 0.23; (−7)/(+316), with an $r^2$ of 0.60; and (−244)/(+316), with an $r^2$ of 0.27. We estimated the frequency of the eight possible haplotypes for these three variants in the 146 female-THR probands and in the 375 female controls (data not shown). There was no significant difference ($p = 0.17$).

The three IL8 variants that we genotyped are all in strong linkage disequilibrium with each other: (−251)/(1633) have an $r^2$ of 0.82; (−251)/(2767) have an $r^2$ of 0.84; (1633)/(2767) have an $r^2$ of 0.93. When we estimated the frequency of the eight possible IL8 haplotypes we noted that two haplotypes were extremely common and accounted for over 93% of the chromosomes for this gene (data not shown). Hull et al.

### Table 1 Candidate genes

<table>
<thead>
<tr>
<th>Gene</th>
<th>Location</th>
<th>Position</th>
<th>Nucleotide change</th>
<th>Amino acid substitution</th>
<th>Female controls</th>
<th>Female probands</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>IGFBP7</td>
<td>Intron 2</td>
<td>+4†</td>
<td>T/A</td>
<td></td>
<td>0.38 (279/465)</td>
<td>0.41 (116/169)</td>
<td>0.37</td>
</tr>
<tr>
<td>ADAMTS3</td>
<td>Intron 1</td>
<td>−7‡</td>
<td>Indel**</td>
<td></td>
<td>0.45 (351/397)</td>
<td>0.39 (108/172)</td>
<td>0.059</td>
</tr>
<tr>
<td>ADAMTS3</td>
<td>Intron 2</td>
<td>−244‡</td>
<td>A/T</td>
<td></td>
<td>0.31 (229/511)</td>
<td>0.33 (94/190)</td>
<td>0.56</td>
</tr>
<tr>
<td>ADAMTS3</td>
<td>Intron 2</td>
<td>−128‡</td>
<td>T/C</td>
<td></td>
<td>0.38 (284/458)</td>
<td>0.33 (97/195)</td>
<td>0.16</td>
</tr>
<tr>
<td>ADAMTS3</td>
<td>Exon 3</td>
<td>+316‡</td>
<td>A/G</td>
<td>Arg138Lys</td>
<td>0.03 (22/712)</td>
<td>0.07 (19/269)</td>
<td>0.015</td>
</tr>
<tr>
<td>ADAMTS3</td>
<td>Intron 15</td>
<td>−19‡</td>
<td>A/C</td>
<td></td>
<td>0.19 (140/598)</td>
<td>0.18 (51/239)</td>
<td>0.67</td>
</tr>
<tr>
<td>ADAMTS3</td>
<td>3’ UTR</td>
<td>+937‡</td>
<td>G/A</td>
<td></td>
<td>0.08 (63/681)</td>
<td>0.10 (29/259)</td>
<td>0.49</td>
</tr>
</tbody>
</table>
previously reported this phenomenon in a European population (haplotypes 1 and 12 of that publication). There was no significant difference in the frequency of the ILS haplotypes between our OA probands and our controls (p = 0.20).

**DISCUSSION**

Selecting candidate genes for an association analysis is a subjective venture influenced by current, and probably limited, knowledge of the biological basis of trait variability. As a first step toward identifying the chromosome 4q dip OA susceptibility we investigated three genes that map within the linkage interval, that have a role in skeletal biology, and which are expressed in OA articular cartilage chondrocytes: IGFBP7, ADAMTS3, and ILS.

Insulin-like growth factors (IGFs) maintain the steady state metabolism of proteoglycans in articular cartilage, and a number of studies have examined the role of the IGFs and the insulin-like growth factor binding proteins (IGFBPs) in OA. Two interesting recent findings are that IGF-I can reverse the insulin-like growth factor 1 (IL1) mediated destruction of articular cartilage and that increasing concentrations of IGFBPs 3 and 5 can lead to improvements in the OA joint. IGFBP7 has not been subjected to the same intensive investigation as other IGFBPs. However, its expression in OA articular cartilage prompted us to investigate the IGFBP7 gene. We detected no common coding variants and only one common intronic variant, which was not associated with OA.

The ADAMTS (a disintegrin and metalloproteinase domain with thrombospondin motifs) metalloproteinase family includes proteinases which mediate cartilage aggrecan degradation as well as procollagen peptidases involved in collagen biosynthesis. It has been proposed that ADAMTS3 has a role in the biosynthesis of type II procollagen, the principal collagen of articular cartilage. The ADAMTS3 gene is large, with 22 exons, a transcript length of 5.8 kb and a coding region length of 2262 bp, a transcript size of 5296 bp. We identified seven common ADAMTS3 variants. The significant p value observed in the 146 probands for the intron 12 SNP was not replicated in the additional 244 cases. It seems likely, therefore, that this initial positive result was a false positive. The genotyping of this SNP in additional cohorts will support or refute this conclusion.

Cytokines have a role in regulating the catabolic/anabolic balance of articular cartilage, and an increase in catabolic activity mediated by IL1β or tumour necrosis factor α (TNFα) can induce an OA phenotype in model systems. IL8 appears to be a mediator of cartilage catabolic responses. None of the ILS variant alleles or haplotypes showed any association with OA in our study.

Overall, we did not generate convincing evidence to support our hypothesis that IGFBP7, ADAMTS3, or IL8编码 for the hip OA susceptibility that we have linkage to chromosome 4q. We cannot, however, exclude these candidates because variants within regulatory elements of the genes that affect gene expression rather than variants that alter amino acid sequence might predispose to OA.

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The authors declare that they have no competing interests.

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