Genetic analysis of CRHRA1 and CRHRA2 microsatellites and their association with rheumatoid arthritis in South Asian and Caucasian populations of the East Midlands, UK

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Abstract

Two microsatellites from corticotropin releasing hormone gene locus (CRHRA1 and CRHRA2) were reported to be associated with rheumatoid arthritis (RA) in Caucasians. This study aims to replicate the association in a South Asian (SA) and Caucasian RA sample from the East Midlands, UK. DNA from 281 South Asians (111 patients, 43 siblings, 127 controls) and 287 Caucasians (116 patients, 64 siblings, 107 controls) were genotyped. The Odds Ratio for carrying at least one copy of the CRHRA1*10 risk allele was 1.32 (Confidence Interval, CI=0.77-2.28) in South Asians and 1.55 (CI=0.92-2.65) in South Asians, indicative of a trend for association. The risk allele CRHRA2*14 was lower in South Asians compared to Caucasians (5.5-8.6% vs. 17.7-18.2%; P<0.005). Significant linkage disequilibrium was observed between CRHRA1 and CRHRA2 in both cohorts. CRHRA1*10 was the frequently transmitted allele in SA patients. A non-significant association was observed with CRHRA1*10 allele in both RA populations.

Introduction

Rheumatoid arthritis (RA) is a common autoimmune disease in which both environmental and genetic factors seem to play a role in the inception, progression, severity and episodic nature of the disease. Genome-wide and candidate association studies have suggested many genetic susceptibility loci, but HLA-DR, and more recently PTPN22, PAD4 and CTLA4, remain the validated genetic markers for RA.1-3 Panayi reported that RA is multifactorial and may be complicated by non HLA factors such as the neuroendocrine hormone, the corticotropin releasing hormone (CRH).1 The hormone is released from hypothalamus under the influence of TNFα, IL-1β and IL-6.

Centrally, CRH releases pituitary adrenocorticotropin, which in turn stimulates the adrenal cortex to release cortisol from the adrenal gland. The amount of cortisol released is reduced in RA patients.4 Lewis rats, which are susceptible to RA induction, also have low cortisol levels. Locally, the CRH levels are increased in the joints of experimental arthritis (in Lewis rats) and in the synovium and synovial fluid of RA patients. In situ CRH increases the cycloxygenase-2 derived prostaglandins E2 and E1 locally, thus promoting joint disease progression and pain perception.

Baerwald et al. (2000)5 studied the genetic polymorphisms in the promoter region of the CRH gene among Caucasian (UK) and South African black RA patients and found that haplotype A2B1 was protective against developing RA in UK patients whereas allele A1B1 was positively associated with RA in the black South African RA patients (OR=1.78; CI=1.01-3.15). Gonzalez-Gay et al.6 found that CRH-A2 allele was significantly increased only in the late onset sero-negative RA patients from the Lugo region of Spain.

Fife et al.7 investigated short tandem repeat (STR)/microsatellite genetic variation at CRHRA1 (8q12.3) in 295 Caucasian families and found a significant excess of allele sharing in affected members and significant linkage over a 22 cM region between markers D8S285 and D8S530. A further STR located 20 kb downstream of the CRH structural gene named CRHRA2 also showed strong linkage disequilibrium with CRHRA1.8 Fife and colleagues reported that the haplotype (CRHRA1*10 - CRHRA2*14) was associated with RA in 131 simplex Caucasian families.9 In a Spanish study (121 simplex RA families and 101 healthy controls) significant linkage disequilibrium between the two loci was observed but there was no association of any alleles or haplotypes with RA.10 The haplotype RA1*10 - RA2*14 was found to be undertransmitted in the study (12 observed versus 17.43 expected).

Previous studies have shown interesting genetic variation in Caucasian and other ethnic populations9 but no studies have examined CRH microsatellite variation in South Asian RA patients. The increased production of PGE1 due to polymorphic CRH, with subsequent hypersensitisation of the pain nerve endings, may be associated to the differing pain perception in the two ethnic groups as some studies on South Asians have reported lower pain thresholds in RA. This difference in pain may be readily reflected in the severity of self assessed RA. The aim of this study was to determine if the two STRs (CRHRA1 and CRHRA2) are associated with RA in South Asian and Caucasian populations of the East Midlands.

Materials and Methods

Subsequent to the approvals by local ethical committees and written consent according to the 1996 Declaration of Helsinki, DNA samples were obtained and genotyped from 111 South Asian Patients, 43 South Asian siblings, 127 South Asian controls, 111 Caucasian Patients, 64 Caucasian siblings and 107 Caucasian control. Genotyping for the two CRH microsatellite markers was performed as described by Fife et al.11 in the presence of fluorescently labelled primers (CRHRA1 - (F) 5'-FAM CCC AGT CCC CAT GAT ATC AG -3', (R) 5'-AAC TTT GCT TCT ACA G -3', and CRHRA2 - (F) 5'-HEX CAG TTT CCT TGG GCT TCT ACA G -3', (R) 5'-AAG TCC TTA TCT TCA AAG CAA T-3').

Allele and genotype frequencies were compared between cases and controls using chi-squared statistics and odds ratios were calculated by Woolf's method. Family based association analysis was carried out using Sib-TDT14.
which tests for association by comparing marker allele frequencies between affected and unaffected siblings which differ in genotype. Haplotype frequencies and linkage disequilibrium between microsatellites were estimated using EHPLUS program.13

A questionnaire was used to assess self perception of severity, pain and quality of life. Patients were asked to score three categories for pain and severity in a simple questionnaire as to how they perceived their pain due to RA - Mild, Moderate or Severe. The gradations of the duration of disease, self assessed severity and pain were assessed by two-way association analysis using cross-tab (contingency Table) method on SPSS against the CRH genotypes.

Results

Both microsatellites analysed at the CRH locus (CRHRA1 and CRHRA2) showed wide genetic variation. At CRHRA1 locus, 7 genotypes (8,10; 9,10; 9,11; 10,10; 10,11; 11,11; and 11, 12) accounted for more than 90% of the genotypic variation observed in various groupings of patients and controls. Two genotypes (12, 13 and 12, 14) accounted for 10 to 18% of the frequency variation. The risk allele CRHRA1*10 frequency was slightly higher in South Asian RA compared to controls (8.6% vs 5.5%). The frequency of CRHRA1*11 was observed to be significantly higher in South Asian RAs (9.90 % versus 3.1%, P<0.05). CRHRA2*14 was rare in South Asians (5.5%) versus. 18.2 %; P = 0.005). When Bonferroni correction (P=0.003) for multiple comparisons was applied these differences became non-significant.

Association analysis

CRHRA1 and CRHRA2 genotypes were in Hardy Weinberg equilibrium in cases and controls in both populations. Two alleles (CRHRA1*10 and *11) accounted for approximately 80% of the genetic variation at CRHRA1 (Table 1). In the South Asian cohort, the previously defined risk allele CRHRA1*10 occurred at a higher frequency in cases (34.3%) than controls (26.9%) although this difference did not reach statistical significance (P=0.11). The frequency of this allele was also slightly higher in Caucasian cases (26.6%) compared with controls (23.5%) (P=0.54). In the Asian cohort, the odds ratio associated with presence (at least one copy) of CRHRA1*10 was 1.55 (95% CI 0.92 – 2.63), compared with an odds ratio of 1.32 (95% CI 0.77-2.28) in the Caucasian cohort.

There was no significant difference in the frequency of CRHRA2*14 between either Caucasian cases (17.7%) and controls (18.2%, P=0.71), or between South Asian cases (8.6%) and controls (5.5%, P=0.26). Interestingly, this allele occurred at significantly lower frequency in the South Asian than Caucasian controls (P=3.0 ¥ 10–5). We did, however, observe a significant increase in the frequency of CRHRA2*11 in South Asian cases (9.9%) than controls (3.1%) (P=0.0045), though after multiple comparison correction, it became no significant.

Linkage disequilibrium analysis

Haplotype frequencies were estimated in 218 South Asians (102 RA Patients and 116 Random Controls) and 208 Caucasians (109 RA Patients and 99 Random Controls) using

Table 1. CRHRA1 and CRHRA2 allele frequencies in two rheumatoid arthritis populations.

<table>
<thead>
<tr>
<th>STR Group</th>
<th>South Asian (n=105)</th>
<th>CRHRA1</th>
<th>Caucasian (n=111)</th>
<th>CRHRA2</th>
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</thead>
<tbody>
<tr>
<td>Allele number</td>
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<td>RA (n=111)</td>
<td>Controls (n=119)</td>
<td>Controls (n=100)</td>
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<td>0</td>
</tr>
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<td>0.4</td>
</tr>
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<tr>
<td>8</td>
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<td>26.6*</td>
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</table>

STR, short tandem repeat; RA, rheumatoid arthritis. *Caucasian Odds Ratio = 1.32, CI 0.77-2.38; South Asians, Odds Ratio=1.55, CI 0.95-2.63; ° P <0.005; #higher, not significant.
**Discussion**

In the present study, a non-significant association was observed with CRHRA1*10 in both populations (Caucasian OR = 1.32, CI 0.77-2.28, South Asians, OR = 1.55, CI 0.95-2.63).

The transmission disequilibrium test (Sib-TDT) analysis, although limited by small numbers, also showed evidence of preferential transmission of CRHRA1*10 to South Asian RA cases. Although none of these findings result in a significant association, it is important to note that the sample sizes here are small, resulting in low power to detect a significant association. Nevertheless, the consistent association with *10 allele in both case-control and sib-TDT studies suggests that further research with a larger sample of South Asians is warranted in order to clarify whether there is an association with RA. Based on the observed frequency of CRHRA1*10 in South Asians, approximately 600 case/control pairs would be required in order to achieve 80% power (\( \alpha = 0.05 \)) to detect a significant association. There was no association with CRHRA2*14 in either population, although a second allele, CRHRA2*11, occurred at higher frequency in South Asian RA cases compared to controls (\( P=0.005 \)).

It may be worth exploring these loci in future research as there is some evidence that stress may contribute to the RA pathology. In the present study, a non-significant association was observed with CRHRA2*11, occurred at higher frequency in South Asians. Differences were not significant.

Sib- transmission disequilibrium test

A total of 43 South Asian and 64 Caucasian families with at least one affected and one unaffected sibling were included in the analysis. In the South Asian cohort, CRHRA1*10 was observed 18 times in RA-affected siblings compared to an expected value of 14.3 (\( P=0.10 \)). However, there was no increase in the frequency of the *10 allele in Caucasian affected siblings (19 observed compared with 18 expected). There was no excess in the frequency of CRHRA2*14 in affected siblings in both Caucasian and Asian groups, although CRHRA2*15 was under-represented in Caucasian affected (4 observed vs. 10 expected, \( P=0.039 \)).

Only CRHRA2 showed a significant association with self reported severity by analogue scale (\( \chi^2 = 156.4, \text{df } = 114, P<0.01 \)) of the disease in South Asians. The perception of pain assessed by analogue score followed a similar pattern to severity of RA.

**References**


2. Plenge RM, Padyukov L, Remmers EF, et al. Replication of putative candidate-gene associations with rheumatoid arthritis in >4,000 samples from North America and Sweden: association of susceptibility with


