HLA typing in families with multiple cases of rheumatoid arthritis

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Abstract
Thirty one white patients from 14 families with multiple cases of rheumatoid arthritis (RA) and 42 of their healthy relatives were completely HLA typed. In contrast with class I antigens, the class II antigens DR1 and DR4 were significantly more common in the patients than in a group of 200 healthy local white controls (DR1: 32% v 12%; DR4: 48% v 28%, in patients and controls respectively).

Owing to the small number of cases the data from this study were combined with those of published reports. Examination of patients for DR1 and DR4 homozygosity and DR1/4 heterozygosity showed an increase of DR1 homoygous patients, which was not statistically significant. There was no striking deviation from random expectation in haplotype sharing of affected sib pairs. These results are comparable with a dominant influence of DR1 and DR4 in the mode of inheritance. The nearly random haplotype sharing and the molecular relation between DR1 and DR4 support the hypothesis of a direct influence of these antigens in the pathogenesis of RA.

Only 68% of the patients in this study possessed either DR1 or DR4, possibly indicating a subtype of RA which is independent of HLA.
family. In table 3 only the index case was compared with his affected siblings. This is important to avoid overrepresentation of families with more than two patients.

**Results**

None of the class I antigens (HLA-A, B, and C) was more prevalent in patients than in healthy controls.

**DR PREVALENCE** (table 2)

In our group both antigens DR1 and DR4 show a significant association with RA (DR1: 32% ± 12%, p<0.005; DR4: 48% ± 28%, p<0.01 for patients and controls respectively). This reflects the results of other investigators.

**NUMBER OF HOMOZYGOTES**

We analysed our data and those of other reported family studies and found 110 families with 213 patients who had been completely HLA typed. The gene prevalence was 13.6% for DR1 and 49.5% for DR4. The Hardy-Weinberg-formula suggests that four DR1 and 52 DR4 homozygous patients might be expected, whereas 10 DR1 homozygous and 51 DR4 homozygous cases were found. The increased number of DR1 homozygotes was not statistically significant. These figures indicate the dominant influence of DR1 and DR4 in the inheritance of RA.

**HAPLOTYPE SHARING**

Table 3 does not show any striking deviation from random expectation in haplotype sharing of affected siblings. The results of some studies (especially small ones) are widely differing and our own figures are surprising, but almost certainly caused by the small number of sib pairs.

Two important studies, with 26 multicase families each showed increased haplotype sharing but these are not listed in table 3 because the data have been published in a form that does not allow statistical comparison.

The data do not prove the hypothesis of an RA gene in linkage disequilibrium with DR1 and DR4, but suggest that certain HLA-DR antigens play a direct part in the pathogenetic process.

**CLINICAL AND SEROLOGICAL INVESTIGATIONS**

No significant association was found between patients with DR1 or DR4 positive RA and the following variables: rheumatoid factor titre, antinuclear factor titre, age at onset of the disease, x ray index (Larsen/Walker), and spread-severity index. The discrepancy in the clinical course and serological findings between siblings with RA was surprising, even in HLA identical sib pairs. It seems that the severity of the disease was mainly determined by factors other than the HLA antigens, though published reports have shown that the prevalence of DR4 is slightly greater in seropositive patients than in seronegative RA.

**Discussion**

The association of RA and the HLA antigens DR1 and DR4 is well established, but it is not clear how these antigens affect the development of RA. If a hypothetic susceptibility gene for RA was linked to the major histocompatibility complex this should result in an increased haplotype sharing of affected siblings. Table 3 shows that this cannot be confirmed. Bodmer tried to explain this observation as follows (a) the HLA linked disease gene is effectively dominant and has a relatively high frequency and therefore a low penetrance; (b) the disease consists of a mixture of two forms, one of which is HLA linked and the other which is not. Bodmer proved that in each of these situations a distortion in the haplotype sharing of affected siblings might be difficult to detect.

Investigations with monoclonal antibodies (MoAb 109d6) reported by Lee showed cross reactions between DR1 and DR4 positive lymphocytes. A DNA sequence analysis found the identity of the third hypervariable region of class II beta chains in DR1 and in the Dw4 subtype of DR4. This supports the common
immunological background of DR1 and DR4 positive RA. This 'shared epitope hypothesis' is an alternative to the idea of an HLA linked RA susceptibility gene.

In our patients only 68% express either DR1 or DR4 (table 1). This fact leads to the assumption of a subtype of RA which is independent of HLA and supports Bodmer's second hypothesis. It will be interesting to see whether further research verifies the existence of RA independent of the major histocompatibility complex and thus shows the genetic heterogeneity of the RA syndrome.

2 Schiff, Mizrachi Y, Orpag S, Yaron M, Gazit E. Association of HLA-DR3 and HLA-DR1 with adult rheumatoid arthritis. Ann Rheum Dis 1982; 41: 403–4.
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