KIR haplotypes are associated with late-onset type 1 diabetes in European–American families

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Classical human leukocyte antigens (HLA) genes confer the strongest, but not the only, genetic susceptibility to type 1 diabetes. Killer cell immunoglobulin-like receptors (KIR), on natural killer (NK) cells, bind ligands including class I HLA. We examined presence or absence, with copy number, of KIR loci in 1698 individuals, from 339 multiplex type 1 diabetes families, from the Human Biological Data Interchange, previously genotyped for HLA. Combining family data with KIR copy number information allowed assignment of haplotypes using identity by descent. This is the first disease study to use KIR copy number typing and unambiguously define haplotypes by gene transmission. KIR A1 haplotypes were positively associated with T1D in the subset of patients without the high T1D risk HLA genotype, DR3/DR4 (odds ratio = 1.29, P = 0.0096). The data point to a role for KIR in type 1 diabetes risk in late-onset patients. In the top quartile (age of onset > 14), KIR A2 haplotype was overtransmitted (63.4%, odds ratio = 1.73, P = 0.024) and KIR B haplotypes were undertransmitted (41.1%, odds ratio = 0.70, P = 0.0052) to patients. The data suggest that inhibitory ‘A’ haplotypes are predisposing and stimulatory ‘B’ haplotypes confer protection in both DR3/DR4-negative and late-onset patient groups.

INTRODUCTION

Type 1 diabetes is an autoimmune disorder in which the insulin-producing cells of the pancreas are destroyed, resulting in a requirement for exogenous insulin for survival. Although > 40 genetic loci are implicated in type 1 diabetes susceptibility, genes encoding the human leukocyte antigens (HLA) are the strongest genetic contributors.1 Killer cell immunoglobulin-like receptors (KIR) are found on the surface and regulate the function of Natural Killer (NK) cells, CD8+ and CD4+ γδ T cells as well as γδ T cells.2 Inhibitory KIR (KIR2DL and KIR3DL groups) have immunoreceptor tyrosine-based motifs within their long cytoplasmic tail. Activating KIR (KIR2DS and KIR3DS groups) have shorter cytoplasmic tails that are coupled via a charged residue in the transmembrane region to adaptor proteins, such as DAP12, which contain immunoreceptor tyrosine-based activating motifs. Ligands for KIR include class I HLA. HLA class I alleles are associated with type 1 diabetes, thus providing a biological rationale to investigate KIR association with type 1 diabetes.3 KIR haplotypes fall into two types: ‘A,’ containing mostly inhibitory genes; and ‘B,’ containing one or more activating KIR genes. Common haplotypes include centromeric (cen) and telomeric (tel) KIR gene groups, exhibiting strong linkage disequilibrium within, but not between, groups (Figure 1). The telomeric A (tA) group includes KIR2DS4, which is the only potentially short-tailed activating KIR on the A haplotype. However, KIR2DS4 is disabled by a 22 bp frameshift deletion in exon 5 on the majority of Caucasian A haplotypes. This distinction defines tA01 (KIR2DS4 full length) and tA02 (KIR2DS4 deletion variant). A small number of studies have reported KIR association with type 1 diabetes; however, most are small case–control studies that focus on individual KIR genes and test only the presence or absence of KIR loci.4 A recent, larger study reports a method for imputation of KIR copy number for one KIR locus using GWAS SNP data but reports no significant type 1 diabetes association.5 In the current study, we performed KIR genotyping, with copy number determination, on a set of type 1 diabetes multiplex families that we previously genotyped for HLA class I and class II loci.6 This unique combination of family-based data with complete HLA and KIR copy number genotyping allows inference of individual haplotypes, based on identity by descent, for both KIR and HLA. We tested association of KIR haplotypes with type 1 diabetes in samples where full HLA context was known. Further, we tested those patients not carrying the highest-risk HLA genotype, that is, heterozygotes for DRB1*03:01-DQA1*05:01-DQB1*02:01 and DRB1*04:01/02/04/05/08-DQA1*03:01-DQB1*03:02 (referred to as ‘DR3/DR4’) separately from the patients who do carry the high-risk genotype. Finally, given that HLA is associated with age of onset for type 1 diabetes,7 we examined the KIR genotyping data for association with age of disease onset.

RESULTS

All families in the Human Biological Data Interchange (HBDI) data set were complete, with two parents and at least two affected siblings. All individuals were genotyped for KIR presence or absence and copy number for each locus.8 All samples were previously genotyped for the loci HLA-DRB1, -DQA1, -DQB1, -DPB1, −A, −B and −C and exhibited Mendelian inheritance. Both HLA and KIR haplotypes were inferred using identity by descent. Common KIR haplotypes are illustrated schematically in Figure 1.

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Full haplotypes, containing both centromeric and telomeric portions, showed marginally significant overtransmission of KIR A1 haplotypes to affected individuals \((P = 0.0494)\). Separate analysis of centromeric and telomeric components of the KIR A1 haplotype (cA01 and tA01) revealed overtransmission of both to affected individuals, although neither result reached statistical significance. T1D patients were grouped into those who carry the very high-risk HLA class II genotype known as ‘DR3/4’ \((DRB1^*03:01-DQA1^*05:01-DQB1^*02:01\) and \(DRB1^*04:01/02/04/05/08-DQA1^*03:01-DQB1^*03:02/04\) or \(*02:01)\) and those who do not (‘non-3/4’). Transmission Disequilibrium Test (TDT) analyses revealed no significant transmission distortion in the high-risk DR3/4 group for any KIR haplotype tested. However, KIR A1-positive association with T1D was observed in the non-3/4 group, in which the HLA class II attributable risk is lower \((P = 0.0096, \text{Table} \ 1)\). Analysis of the data with Unphased 3.1.6 software also revealed this association \((P = 0.006, \text{not shown})\). Separate analysis of telomeric and centromeric haplotypes showed significant overtransmission of the tA01 \((P = 0.0169, \text{Table} \ 1; \text{Unphased:} \ P = 0.003, \text{not shown})\). tA01 is included in ~55% of common KIR haplotypes in European populations (Figure 1). To test the hypothesis that KIR3DL1 may contribute to the T1D predisposing effect observed for tA01, the data were stratified by the presence or absence of HLA-Bw4, the ligand for KIR3DL1. Both the Bw4-positive and Bw4-negative subsets had overtransmission of tA01 (54.5% and 55.2%, odds ratio = 1.23 and 1.20, respectively), with the Bw4-negative result reaching the threshold for statistical significance \((P = 0.05; \text{data not shown})\). The fact that tA01 overtransmission is observed either in the presence or absence of the ligand for KIR3DL1 suggests that the KIR3DL1-Bw4 interaction is not the basis for the disease association of the tA01 haplotype.

HLA-C alleles can be divided into two groups, C1 and C2, depending on the identity of the amino-acid residues at position 77 and 80. HLA C2 alleles, which are ligands for 2DS1 and 2DL1, are underrepresented in T1D, whereas C1 alleles, which are ligands for 2DL2, 2DL3 and 2DS2 (weak interaction), are overrepresented. Data from the current study were consistent with that report; C2 was undertransmitted \((\text{frequency transmitted} = 0.291, \text{frequency not transmitted} = 0.336)\) to affected individuals, whereas C1 was overtransmitted \((f\text{-trans} = 0.709, f\text{-not trans} = 0.66; P = 0.000139)\) (data not shown). However, testing for gene–gene interaction between HLA-C allele groups and KIR loci showed no evidence of interaction for any KIR locus with C1 or C2 (data not shown).

**HLA** genes, particularly class I, are associated with age of onset for T1D, suggesting that KIR loci may also be associated in this manner. Mean age of onset for T1D patients was 11.22 years \((\text{range} \ 0–36 \text{ years, s.d.} = 7.57; \text{95%CI} = 10.66–11.79)\); median age of onset was 10 years. Patients were grouped into four age of onset categories: (1) \(>0 \text{ and} \leq 5\) \((n = 181)\); (2) \(>5 \text{ and }< 10\) \((n = 156)\); (3) \(\geq 10 \text{ and }< 14\) \((n = 197)\), (4) \(>14\) \((n = 167)\). Analysis of patients in the top age of onset quartile \((>14 \text{ years})\) revealed significant transmission distortion for several KIR haplotypes, including A2 \((63.4%, p = 0.024; \text{Unphased} = 0.044)\) and B \((41.1%, p = 0.0052; \text{Unphased} = 0.001)\), Table 2). No significant over- or undertransmission of KIR A and B haplotypes was observed in patients with age of onset 14 or younger. Notably, the KIR haplotypes that are associated in the over 14 age group \((A2 \text{ and } B, \text{Table} \ 2)\) differ from those that are associated in the non-DR3/4 group \((A1, \text{and more specifically, } tA01, \text{Table} \ 1)\), although all data were consistent with overtransmission of KIR A haplotypes and undertransmission of KIR B haplotypes.

**DISCUSSION**

The fact that some HLA class I molecules are ligands for KIR leads to the notion that KIR may have a role in T1D susceptibility. Published reports of KIR association studies with T1D have little consistency among findings for particular loci \((\text{see allelefrequencies.net/diseases}^4 \text{ and references therein})\). Most are small case–control studies based only on presence or absence of individual KIR loci. The results reported in the current study are based on KIR haplotypes, inferred using identity by descent from family-based data, rather than on individual loci.

Our finding that the predisposing effect of the tA01 haplotype is more pronounced in the subset of T1D patients not carrying the high-risk DR3/DR4 genotype increases our confidence that the observed predisposing effect is real. This is consistent with previous studies of T1D risk loci other than HLA-DR- and DQ-encoding genes, including TCF7, IL-4 R and PTPN22, where...
Thus, our finding that TA01 haplotypes are predisposing for T1D is consistent with the NOD result. If the dominance of the inhibitory or educative activity of KIR3DL1 is important then presumably this would only be apparent when an appropriate HLA ligand (Bw4+) or TA02 molecules that have threonine at position 80, which show stronger interaction with KIR3DL1 receptors than BW4 molecules that have threonine at this position (54.7% (odds ratio = 1.21, 95% CI 0.92–1.59). Using a combination of KIR copy number typing and SNP imputation of KIR copy number, Pontikos et al. report no association of 3DL1/3DS1 with KIR and T1D susceptibility appears to be modest. This does not rule out the presence of the KIR2DS4v gene that encodes a soluble and potentially secreted form of the protein. This may be biologically relevant if it antagonizes the effects of other KIRs. These haplotype-based data suggest that KIR genes have a role in T1D susceptibility. This study was powered with the assumption that KIR haplotype effects might be similar in magnitude to those on T1D can be made, the combination of evidence from genetic and animal models warrants further study of this locus. An alternative explanation for the TA01 association could relate to the presence of the KIR2DS4v gene that encodes a soluble and potentially secreted form of the protein. This may be biologically relevant if it antagonizes the effects of other KIRs. The effect of the risk allele was stronger in the non-DR3/4 patients than in the DR3/4 group. One report of functional studies in the NOD mouse model of T1D suggests that expression of the inhibitory receptor KIR3DL1 predisposes NOD mice to diabetes by downregulating function of regulatory T cells. KIR3DL1 is found on telomeric KIR A haplotypes; thus, our finding that TA01 haplotypes are predisposing for T1D is consistent with the NOD result. If the dominance of the inhibitory or educative activity of KIR3DL1 is important then presumably this would only be apparent when an appropriate HLA ligand (Bw4+) is also present. Although no statistically significant transmission distortion was observed in the HBBDI data set for KIR in the context of HLA class I ligands, the combination of KIR3DL1 with the gene for Bw4-801 (Bw4 molecules that have isoleucine at dimorphic position 80, which show stronger interaction with KIR3DL1 receptors than Bw4 molecules that have threonine this position (Bw4-80 T)) exhibited the highest transmission percentage at 54.7% (odds ratio = 1.21, 95% CI 0.92–1.59). Using a combination of KIR copy number typing and SNP imputation of KIR copy number

Table 1. TDT analysis of full, centromeric and telomeric KIR haplotypes for type 1 diabetes association with stratification by DR3/4 status

| KIR Haplotype^a | Trans | N-trans | TDTTest^b | P-value^c | Trans % | OR (95% CI) |
|-----------------|-------|---------|-----------|-----------|---------|-------------|
| A B full, non-DR3/4 |       |         |           |           |         |             |
| A1              | 236   | 183     | 6.70      | **0.0096**| 56.3%   | 1.29 (1.06–1.56) |
| A2              | 91    | 91      | 0.00      | 1.00      | 50.0%   | 1.00 (0.75–1.34) |
| B               | 283   | 314     | 1.61      | 0.20      | 47.4%   | 0.90 (0.77–1.06) |
| Others          | 37    | 59      | 5.04      | **0.0247**| 38.5%   | 0.63 (0.42–0.95) |
| A B full, DR3/4 |       |         |           |           |         |             |
| A1              | 128   | 121     | 0.20      | 0.66      | 51.4%   | 1.06 (0.83–1.36) |
| A2              | 53    | 53      | 0.00      | 1.00      | 50.0%   | 1.00 (0.68–1.46) |
| B               | 185   | 198     | 0.44      | 0.51      | 48.3%   | 0.93 (0.76–1.14) |
| Others          | 31    | 25      | 0.64      | 0.42      | 55.4%   | 1.24 (0.73–2.10) |
| Cen haplotypes, non-DR3/4 |       |         |           |           |         |             |
| cA01            | 421   | 381     | 2.00      | 0.16      | 52.5%   | 1.10 (0.96–1.27) |
| cB01            | 109   | 117     | 0.28      | 0.59      | 48.2%   | 0.93 (0.72–1.21) |
| cB02            | 80    | 90      | 0.59      | 0.44      | 47.1%   | 0.89 (0.66–1.20) |
| Others          | 37    | 59      | 5.04      | **0.0247**| 38.5%   | 0.63 (0.42–0.95) |
| Cen haplotypes, DR3/4 |       |         |           |           |         |             |
| cA01            | 232   | 220     | 0.32      | 0.57      | 51.3%   | 1.05 (0.88–1.27) |
| cB01            | 67    | 87      | 2.60      | 0.11      | 43.5%   | 0.77 (0.56–1.06) |
| cB02            | 67    | 65      | 0.03      | 0.86      | 50.8%   | 1.03 (0.73–1.45) |
| Others          | 31    | 25      | 0.64      | 0.42      | 55.4%   | 1.24 (0.73–2.10) |
| Tel haplotypes, non-DR3/4 |       |         |           |           |         |             |
| TA01            | 324   | 266     | 5.70      | **0.0169**| 54.9%   | 1.22 (1.04–1.43) |
| TA02            | 142   | 149     | 0.17      | 0.68      | 48.8%   | 0.95 (0.76–1.20) |
| TB01            | 144   | 173     | 2.65      | 0.10      | 45.4%   | 0.83 (0.67–1.04) |
| Others          | 37    | 59      | 5.04      | **0.0247**| 38.5%   | 0.63 (0.42–0.95) |
| Tel haplotypes, DR3/4 |       |         |           |           |         |             |
| TA01            | 181   | 189     | 0.17      | 0.68      | 48.9%   | 0.96 (0.78–1.17) |
| TA02            | 93    | 98      | 0.13      | 0.72      | 48.7%   | 0.95 (0.71–1.26) |
| TB01            | 92    | 85      | 0.28      | 0.60      | 52.0%   | 1.08 (0.81–1.45) |
| Others          | 31    | 25      | 0.64      | 0.42      | 55.4%   | 1.24 (0.73–2.10) |

Abbreviations: CI, confidence interval; OR, odds ratio. ^A B Full = haplotypes containing both the centromeric and telomeric groups. A1 = 3DL3-2DL3-2DP1-2DL1-3DP1-2DL4-3DL1-3DS3FL-3DL2; A2 = 3DL3-2DL3-2DP1-2DL1-3DP1-2DL4-3DL1-2DS4FL-3DL2; B = haplotypes with at least one activating KIR gene; cA01 = 3DL3-2DL3-2DP1-2DL1-3DP1; cB01 = 3DL3-2DS2-2DL2-2DP1; cB02 = 3DL3-2DS2-2DL2-2DS3-2DS5-2DS5-2DS5-2DP1-2DL1-3DP1; tA01 = 2DL4-3DL1-2DS4FL-3DL2; tA02 = 2DL4-3DL1-2DS4FL-3DL2; tB01 = 2DL4-3DS1-2DL5-2DS35T-2DS1-2DL2; others: haplotypes with frequency < 1.0%. *TDT tests were performed separately for each affected sibling in a family. Testing using an average of the two siblings for each family gave similar results; however, the P-values did not reach statistical significance. Significant P-values (> 0.05) are indicated with bold type. P-values are uncorrected based on prior hypothesis for KIR-T1D association. Trans, number of transmitted haplotypes; N-trans, number of untransmitted haplotypes.
report support the notion that KIR A haplotypes, which contain mostly inhibitory KIR genes, are predisposing for T1D, whereas KIR B haplotypes, which contain more activating KIR loci, are protective. Activating KIR could act by stimulating regulatory T cells to control disease, or by influencing NK killing of effector T cells. On the other hand, inhibitory KIR could constrain the action of T1D-specific regulatory T cells, or fail to control cytotoxic T cells, allowing disease progression. In contrast with autoimmune diabetes in children, where HLA predominates, the genetics of autoimmune diabetes in adults are not well understood, despite its relatively high prevalence. Our finding will potentially be useful to better categorize patients in the future and to probe the influence of KIR on NK/T cells and autointer-genic-specific regulatory T cells. Given the complexity of both the HLA and KIR genetic loci, and extreme variability of both among populations, a thorough understanding of the role of KIR genes and haplotypes in T1D will require very large sample sets, where individual subsets are large enough to uncover potential highly significant results.

SUBJECTS AND METHODS

The HBDI is a repository for families including children with T1D (National Disease Research Interchange; http://ndiresource.org/Donor-Programs/Family-Genetics-HBDI/36/). These families were previously genotyped for the loci HLA-DRB1, -DQA1, -DQB1, -DPB1, A, -B and -C and were analyzed extensively for T1D association.8,13 The 339 families in this study are multiplex, that is, include more than one affected child. Parents are unaffected. All families exhibit Mendelian inheritance. HBDI families were genotyped for HLA class I and class II at two-field (four digit) resolution by polymerase chain reaction sequence-specific oligonucleotide probe (PCR-SSO) technology as previously described.6,11 KIR genotype and copy number were measured using a quantitative PCR comparative Ct method developed from a protocol for determining genomic copy number as described earlier.8 Reactions were carried out in quadruplicate to ensure accuracy of the copy number scoring. Identity by descent-inferred KIR haplotypes were determined using the Merlin program as previously described.8 A TDT was applied to the data (two-sided test). Each family was analyzed as trios for each affected sibling. Odds ratios and 95% confidence intervals were estimated using the proportion T of transmitted risk allele as described.14 The estimate of the odds ratio is given by: T/(1−T). In addition, Unphased software was used to test gene–gene interaction between HLA allele groups and KIR loci. P-values were not corrected based on prior hypothesis for KIR-T1D association.

CONFLICT OF INTEREST

The authors declare no conflict of interest.
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