Association of Killer Cell Immunoglobulin-Like Receptor Genes with Pandemic Influenza A (H1N1)pdm09 Infection in Critically Ill Macedonian Patients

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Key words: Killer immunoglobulin-like receptor (KIR) gene polymorphism; KIR genotyping; PCR-SSP; patients with (H1N1)pdm09 infection; Republic of Macedonia.

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Abstract

Background: Infection by the pandemic influenza A (H1N1)pdm09 virus results in significant pathology disease in many cases in different populations worldwide. The natural killer (NK) cells are among the major effectors important in early innate immune responses to viral infections, interacting with host cells through their activating or inhibiting receptors.

Aim: The aim of this study was to analyze Killer Ig-Like Receptor (KIR) gene polymorphisms in critically ill Macedonian patients with pandemic influenza A (H1N1)pdm09 infection.

Material and Methods: The studied sample consists of 63 critically ill Macedonian patients with pandemic influenza A (H1N1)pdm09 infection. The population genetics analysis package, Arlequin, was used for analysis of the data.

Results: We found that all 16 KIR genes were observed in the studied individuals and framework genes (KIR3DL3, KIR3DP1, KIR2DL4, and KIR3DL2) were present in all individuals. The results of tested linkage disequilibrium (LD) among KIR genes demonstrated that KIR genes present a wide range of linkage disequilibrium. Comparison of KIR gene frequencies between critically ill H1N1/09 Macedonian patients and healthy subjects reveals statistically significant difference for frequency of KIR2DL1 (F=1 in the patients group, and 0.94 in the control group, p=0.045).

Conclusion: We did not found any significant association of all 16 KIR genes or KIR genotypes with critically ill (H1N1)pdm09 Macedonian patients, except for the KIR2DL1.

Introduction

Infection with influenza A virus resulted in significant disease in many cases in different populations worldwide. The rapid global spread of the novel swine origin influenza virus A (H1N1)pdm09 and more than 620,000 cases reported worldwide as of the end of 2009 [1, 2], prompted the World Health Organization to raise the pandemic alert to the highest level, officially declaring pandemic [3]. An intriguing observation was that the highest prevalence of significant pathology was reported among youths and young adults [4-7]. It is known that innate immune cells, such as the macrophages, neutrophils, and DCs are efficient in clearing inûuenza virus via phagocytosis or promotion of adaptive responses [8-13]. In the recent years, the role of NK cells in response to inûuenza infection has also been studied extensively, and their involvement in the early control of
The aim of this study was to examine KIR gene polymorphisms by determining the frequencies of 16 KIR genes and pseudogenes (KIR2DL1, KIR2DL2, KIR2DL3, KIR2DL4, KIR2DL5, KIR3DL1, KIR3DL2, KIR3DL3, KIR2DS1, KIR2DS2, KIR2DS3, KIR2DS4, KIR2DS5, KIR3DS1, KIR2DP1, and KIR3DP1) and KIR genotypes in Macedonian patients with pandemic influenza infection, and to compare the gene content between patients with very severe disease including fatalities and patients with less severe disease.

To our knowledge, this is the first study of the diversity of KIR genes in patients with pandemic influenza A (H1N1)pdm09 infection in the Republic of Macedonia and among the few in the world.

**Material and Methods**

**Population samples.** The study included 63 unrelated patients (male 39 and female 25) with laboratory PCR confirmation of Influenza (H1N1)pdm09 pandemic flu hospitalized at the University Clinic of Infective Diseases in Skopje, Republic of Macedonia, between December 2009 and March 2010. Only consecutive patients with severe disease course (with need of ventilation support) were selected.

After signing of written consent, genomic DNA was extracted from the peripheral blood leukocytes using standard phenol/chloroform procedure, described elsewhere [34], and stored in the Macedonian Human DNA Bank (hDNAMKD) [35] until processing.

**PCR amplification.** For KIR genotyping, commercially available PEL-FREEZ KIR genotyping SSP kit (Dynal Biotech, Brown Deer, WI) was used. It is a PCR-based method (using sequence-specific priming approach) designed to detect the presence or absence of 16 KIR genes and pseudogenes defined by the International nomenclature committee of WHO [36, 37]. Briefly, locus specific primer sets, dispensed in a 96 well thermal tray were used for amplification of genomic DNA. After the amplification, the PCR products are loaded and separated by electrophoresis onto a 2% agarose gel stained with ethidium bromide, after which the results are interpreted using a worksheet for the specific amplification patterns. The presence of each KIR gene was determined by the presence of a band of DNA of the expected size.

All PCRs contained an internal positive control consisting of an additional pair of primers specific for the growth hormone (GH) gene and a negative control [38]. Individuals were determined negative for a particular KIR gene when a band of expected size was absent in
the presence of a band for the GH gene. We have used external quality control consisting of cell lines from Immunogenetics and Histocompatibility Workshop Conferences and Centre d’ Etude du Polymorphisme Humain.

**Statistical analysis.** The occurrence of KIR genes in individuals (frequency = F) was obtained by direct counting. Gene frequencies (GF) were calculated using the formula $GF = 1 - \sqrt{1 - F}$, being aware of the limitation in its ability to detect KIR genes present at low frequency. For analysis of the molecular polymorphism of the locus studied, the Arlequin software version 3.0 (Genetics and Biometry Laboratory, University of Geneva, Switzerland) [39] was used.

Linkage disequilibrium (LD) values for two locus associations were calculated using 2×2 tables [40]. Because LD is not independent of allele frequencies, normalized LD was calculated as described previously [41, 42]. Comparisons of different genotypes for two groups were tested by the $\chi^2$ test. Crude odds ratios (OR) were calculated within 95% CI.

**Results**

**KIR gene frequencies.** The frequencies of the 16 KIR genes (14 genes and 2 pseudogenes) determined in the 63 Macedonian (H1N1)pdm09 patients, is shown in Table 1 along with the corresponding frequencies of the 214 healthy Macedonian controls. All 16 KIR genes were observed in both groups of the studied population and framework genes (KIR3DL3, KIR3DP1, KIR2DL4, and KIR3DL2) were present in all individuals.

Comparison of KIR gene frequencies between critically ill (H1N1)pdm09 Macedonian patients and healthy Macedonians reveals statistically significant difference for KIR2DL1 (frequency of 1 in the patients group, and 0.94 in the control group, p=0.045) (Table 1).

**Linkage Disequilibrium.** The classical linkage disequilibrium coefficient (D), linkage disequilibrium coefficient D standardized by the maximum value it can take ($D_{max}$), given the allele frequencies (D'), standardised simple measure of linkage disequilibrium ($\rho$), and statistical significance (P) for KIR genes are shown in Table 2. The genes present in all individuals (KIR2DL1, KIR2DL4, KIR3DL2, KIR3DL3, KIR2DP1 and KIR3DP1) were excluded from the analysis.

Pairs of KIR loci that displayed significant (P<0.05) LD in critically ill Macedonian patients with pandemic influenza A (H1N1)pdm09 infection are given in Table 3. Positive LD was observed between pairs KIR3DL1 and KIR2DS4, KIR2DL2 with KIR2DS2 and KIR2DS3, KIR2DL5 and KIR3DS1, KIR3DS1 with KIR2DS1 and KIR2DS5, KIR2DS1 and KIR2DS5, and between KIR2DS2 with KIR2DS3. Negative LD was found between KIR3DL1 and KIR2DS5, KIR2DL3 and KIR2DS3, and KIR2DS4 and KIR2DS5. Other KIR genes were not in significant LD.

**Genotype frequencies.** KIR groups, genotype ID, KIR genotypes, number of individuals displaying certain genotype, and the frequency of genotypes are shown in Table 4.

If any of the genes 2DL2, 2DL5, 3DS1, 2DS1, 2DS2, 2DS3, or 2DS5 was present, the genotype was

### Table 1: Comparison of the observed and estimated KIR gene frequencies for critically ill Macedonian patients with pandemic influenza A (H1N1)pdm09 infection (N = 63) and healthy Macedonians (N=214).

<table>
<thead>
<tr>
<th>Pseudogenes</th>
<th>Inhibitory KIR</th>
<th>Non inhibitory KIR</th>
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<tr>
<td></td>
<td>H1N1 Infection (N)</td>
<td></td>
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<tr>
<td></td>
<td>63</td>
<td>63</td>
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<tr>
<td></td>
<td>H1N1 Infection (F)</td>
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<td></td>
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<td>1</td>
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<td></td>
<td>H1N1 Infection (GP)</td>
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<tr>
<td></td>
<td>1</td>
<td>1</td>
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<tr>
<td>Healthy Macedonians (N)</td>
<td>210</td>
<td>214</td>
</tr>
<tr>
<td>Healthy Macedonians (F)</td>
<td>0.980</td>
<td>1</td>
</tr>
<tr>
<td>Healthy Macedonians (GP)</td>
<td>0.670</td>
<td>1</td>
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<tr>
<td>Pearson’s p</td>
<td>0.274</td>
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<td>OR</td>
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<td>Wald 95% CI</td>
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N, number of individuals; F, observed frequency was obtained by direct counting; GF, gene frequencies were calculated using the formula $GF = 1 - \sqrt{1 - F}$; p, statistical significance; &, cannot be calculated because expected <5, c2 test; OR, Odds ratio; CI, confidence interval.
considered as B. If none of these were present, genotype is considered as AA. We have not attempted to distinguish between AB and BB genotypes and called any of this Bx.

KIR genotypes were numerated according to the Allelefrequencies Database [43]. Total of 29 different KIR genotypes were found to be present in the studied sample, based on the presence of 16 KIR genes. We have found two AA genotypes, AA1 and AA180 with frequencies of 0.127 and 0.016, respectively. The most frequent genotypes in the Bx group were genotypes Bx2 (F=0.095), Bx5 (F=0.079) and Bx4 (F=0.064). One new genotype of the Bx group was found and is being referred to Allelefrequencies.net (Table 4).

There is not statistically significant difference in distribution of AA and Bx KIR genotypes between Macedonian patients with (H1N1)pdm09 infection with severe course compared to healthy Macedonians (P=0.207, OR=0.609, Wald 95% CI=0.280-1.324) (Table 5).

Discussion

Influenza A has been an important threat to global public health and remains in the focus of clinical diagnosis, treatment, and basic research [44, 45].
hundreds of thousands of patients have been reported during the last outbreak of the (H1N1)pdm09 virus, we have witnessed that some subgroups of patients have a poorer outcome – the Mexico outbreak suggested that younger patients, especially those with co-morbidities and morbidly increased BMI were more susceptible to respiratory failure [4, 7].

We present the KIR genes distribution in Macedonian critically ill patients infected with (H1N1)pdm09. The studied group of patients and the healthy control subjects belonging to the same, Macedonian population used for comparison, showed similar frequencies for most KIR genes. The only statistically significant difference was noted for the inhibiting KIR2DL1 gene, which was present in all patients (F=1) and in 94% of the controls (P=0.045). Several statistically significant differences were found between the two populations when comparing the frequencies of AA and Bx KIR genotypes, the most notable being for Bx94, which was present in 3 patients, but not present at all in the control group (P=0.001). Two other genotypes with significant differences in frequencies were Bx19 (P=0.012) and Bx63 (P=0.045), both more frequent in the patients group. Comparable predominance of group Bx genotypes has been observed in many different populations, such as North Indians, Palestinians, South Asians, Afro-Caribbeans and also, general Macedonian population [46-49]. In a similar recent study analyzing the KIR polymorphisms in patients infected with Ebola virus [50], significantly higher frequencies of activating KIR2DS1 and KIR2DS3 genes were found in the group of fatally ill patients, when compared to the group of survivors from the infection. This finding led the authors to a conclusion that proposed overactivation of the NK cells was responsible for their rapid depletion. Although for the KIR2DS1 gene, we find "inverse" situation with these gene being more frequent in the control group, we still find this hypothesis tempting and possible, since as much as four activating genes (KIR2DS3, KIR2DS4, KIR2DS5 and KIR3DS1) were present in higher frequency (not statistically significant) in the critically ill patients.

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patients when compared to the healthy subjects. This finding is in concordance with another recent study [51], where predominance of activating KIR genes (KIR2DS5 and KIR3DS1) was noted in severely ill patients with influenza (H1N1)pdm09 infection. It would be interesting to take into account the allelic polymorphism, which might allow different alleles to be expressed differentially and thus influence ligand binding and consecutive cytolysis [50]. Unfortunately, at present, we are not able to perform this analysis.

The results of tested linkage disequilibrium among KIR genes demonstrated that KIR genes present a wide range of linkage disequilibrium. Again, we cannot assume an absolute correlation between the KIR loci, as we only detect a certain percentage of alleles at a locus.

In conclusion, we have determined the distribution of KIR genes in patients with confirmed (H1N1)pdm09 infection and compared it to healthy control subjects, all living in Republic of Macedonia. While there is evident predominance of KIR activating genes in the group of patients with very severe disease, these differences do not reach statistical significance. Our results suggest that activating KIR genes might be a predisposing factor for more severe viral disease and are in agreement with earlier proposed theories [50, 51].

Our next step should be HLA genotyping of the samples in order to explore the ligands for the KIRs in the critically ill Macedonian patients with pandemic influenza A (H1N1)pdm09 infection. Finally, KIR typing at allelic level might be more helpful in elucidating the different efficacy of NK cells and different disease course in virally infected patients.

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References


