Evidence of New Risk Genetic Factor to Systemic Lupus Erythematosus: The UBASH3A Gene

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Abstract

The ubiquitin associated and Src-homology 3 (SH3) domain containing A (UBASH3a) is a suppressor of T-cell receptor signaling, underscoring antigen presentation to T-cells as a critical shared mechanism of diseases pathogenesis. The aim of the present study was to determine whether the UBASH3a gene influence the susceptibility to systemic lupus erythematosus (SLE) in Caucasian populations. We evaluated five UBASH3a polymorphisms (rs2277798, rs2277800, rs9976767, rs13048049 and rs17114930), using TaqMan® allelic discrimination assays, in a discovery cohort that included 906 SLE patients and 1165 healthy controls from Spain. The SNPs that exhibit statistical significance difference were evaluated in a German replication cohort of 360 SLE patients and 379 healthy controls. The case-control analysis in the Spanish population showed a significant association between the rs9976767 and SLE (Pc = 9.9E-03 OR = 1.21 95%CI = 1.07–1.37) and a trend of association for the rs2277798 analysis (P = 0.09 OR = 0.99 95%CI = 0.79–1.02). The replication in a German cohort and the meta-analysis confirmed that the rs9976767 (Pc = 0.02; Pc = 4.7E-03, for German cohort and meta-analysis, respectively) and rs2277798 (Pc = 0.013; Pc = 4.7E-03, for German cohort and meta-analysis, respectively) UBASH3a variants are susceptibility factors for SLE. Finally, a conditional regression analysis suggested that the most likely genetic variation responsible for the association was the rs9976767 polymorphism. Our results suggest that UBASH3a gene plays a role in the susceptibility to SLE. Moreover, our study indicates that UBASH3a can be considered as a common genetic factor in autoimmune diseases.

Introduction

The T cell ubiquitin ligand proteins (TULA) family is characterized by function as suppressors of T cell receptor signalling. One of the members of the TULA family proteins is the ubiquitin associated and Src-homology 3 (SH3) domain containing A (UBASH3a) which is expressed only in lymphoid cells and facilitates apoptosis induced in T cells by certain stimuli, such as growth factor withdrawal [1]. UBASH3a gene spans 40 kb, contains 15 exons and is located on human chromosome 21q22.3 [2]. The lack of TULA proteins resulted in hyper-reactivity of T cells [1]. Evidence for both B and T lymphocyte hyper-reactivity is typically observed in autoimmune disorders [2]. These disorders are characterized by an inappropriate, ultimately excessive, inflammatory response against self, resulting in tissue destruction. Although many individuals affected by autoimmune diseases demonstrate multorgan involvement, the primary end-organ target (e.g., autoimmune destruction of pancreatic islet cells in type 1 diabetes mellitus) typically drives the clinical presentation and disease definition. Recent studies have showed that single nucleotide polymorphisms (SNPs) of the UBASH3a gene are associated with some autoimmune diseases, like type 1 diabetes (T1D), celiac disease (CD), rheumatoid arthritis (RA) and vitiligo, suggesting that this gene could play an important role in the pathogenesis of autoimmune disorders [3–8]. Systemic lupus erythematosus (SLE) is a prototypic autoimmune diseases characterized by the production of autoantibodies, immune-complex deposition, and subsequent multiple organ damage. The complex aetiology of autoimmune diseases includes environmental, hormonal and genetic factors. Some of those factors remained to be defined [3,4]. Based on these insights, the aim of the present study was to evaluate the role of five UBASH3a polymorphism in SLE.
Materials and Methods

Ethics Statement

Written informed consent was obtained from all participants and the respectively ethics committee approved the study according to the principles expressed in the Declaration of Helsinki.

The case-control study included 906 SLE patients and 1165 healthy controls from a white Spanish population. The replication cohort from white Germans comprehends 360 SLE patients and 379 healthy controls. All the patients met the American College of Rheumatology criteria for classification of SLE [5]. Written informed consent was obtained from all participants and the respectively ethics committee approved the study. DNA was obtained from peripheral blood using standard methods. The samples were genotyped for the UBASH3a rs2277798, rs2277800, rs9976767, rs13048049 and rs17114930 polymorphisms via TaqMan® allelic discrimination technology using a predesigned SNPs genotyping assays provided by Applied Biosystems (assay ID: C___1724055_10, C__15885522_20, C___1724067_10, C__1724073_20 and C___25622591_10, respectively; Figure S1). At the moment of the design of the study the only confirmed case-control associated SNP with autoimmune diseases was the rs9976767 [6]. The other four SNPs were selected because they were not included in previous SLE genetic studies and they are non-synonymous changes located in different exons of the UBASH3a gene. Moreover, the minor allele frequency (MAF) of those SNPs was reported in Caucasian populations and they exhibited moderated LD with at least one SNP in the loci. Deviation from Hardy-Weinberg equilibrium (HWE) was tested by standard chi-square analysis. The differences in genotype distribution and allele frequency among cases and controls were calculated by contingency tables and when necessary by Fisher’s exact test. Odds ratios (OR), and 95% confidence intervals (CI), were calculated according to Woolf’s method. Combined data were analysed by Mantel-Haenszel tests under fixed effect model and the Breslow-Day (BD) test was used to estimate the OR heterogeneity amongst the two cohorts. An association was considered statistically significant if \( P < 0.05 \). Benjamini & Hochberg (1995) step-up false discovery rate (FDR) control correction

Table 1. Genotype and minor allele frequencies of UBASH3a SNPs located in Caucasian SLE patients and healthy controls from Spain, the discovery cohort.

| SNP     | Allele 1/2 | Genotype, N (%) | Alleles, N(%) | P-value* | OR [CI 95%]**** |
|---------|------------|-----------------|---------------|----------|-----------------|
|        |            | 1/1 1/2 2/2     | 1 2           |          |                 |
| rs2277798 G/A | Controls (n = 1165) | 477 (40.94) 529 (45.41) 159 (13.65) | 1483 (63.6) 847 (36.4) | 0.0993** | 0.90 [0.79–1.02] |
|        |            | SLE (n = 906)   | 402 (44.37) 394 (43.49) 110 (12.14) | 1198 (66.1) 614 (33.9) | 0.4592 | 1.13 [0.82–1.55] |
| rs2277800 C/T | Controls (n = 1165) | 1080 (92.70) 84 (7.21) | 1 (0.09) 2244 (96.3) 86 (3.7) | 0.4592 | 1.13 [0.82–1.55] |
|        |            | SLE (n = 906)   | 832 (91.83) 73 (8.06) | 1 (0.11) 1737 (95.9) 75 (4.1) | 0.4592 | 1.13 [0.82–1.55] |
| rs9976767 A/G | Controls (n = 1165) | 363 (31.16) 558 (47.90) 244 (20.94) | 1284 (55.1) 1046 (44.9) | 1.99E-03*** | 1.21 [1.07–1.37] |
|        |            | SLE (n = 906)   | 230 (25.39) 451 (49.78) 225 (24.83) | 911 (503) 901 (49.7) | 1.99E-03*** | 1.21 [1.07–1.37] |
| rs13048049 G/A | Controls (n = 1165) | 1038 (89.10) 126 (10.82) 1 (0.09) | 2202 (94.5) 128 (5.5) | 0.9719 | 1.01 [0.77–1.32] |
|        |            | SLE (n = 906)   | 808 (89.18) 96 (10.60) 2 (0.22) 1712 (94.5) 100 (5.5) | 0.9719 | 1.01 [0.77–1.32] |
| rs17114930 C/G | Controls (n = 1165) | 1066 (91.50) 95 (8.15) 4 (0.34) 2227 (95.6) 103 (4.4) | 0.1649 | 1.22 [0.92–1.63] |
|        |            | SLE (n = 906)   | 811 (89.51) 93 (10.26) 2 (0.22) 1715 (94.6) 97 (5.4) | 0.1649 | 1.22 [0.92–1.63] |

*All P-values have been calculated for the allelic model. ** \( PC = 0.248 \) Benjamini & Hochberg (1995). ***\( PC = 9.9E-03 \) Benjamini & Hochberg (1995) step-up FDR control. ****Odds ratio for the minor allele.

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Figure 1. Graphical representation of the meta-analysis (A) Forest plot for the meta-analysis of the UBASH3a rs2277798 polymorphism in SLE in two Caucasian cohorts. (B) Forest plot for the meta-analysis of the UBASH3a rs9976767 polymorphism in SLE in two Caucasian cohorts.

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Table 2. Genotype and minor allele frequencies of UBASH3a SNPs located in Caucasian SLE patients and healthy controls from Germany.

| SNP       | Genotype, N (%) | Alleles, N(%) | Allele test |
|-----------|----------------|---------------|-------------|
|           | 1/2 Subgroup (N) | 1/1 | 1/2 | 2/2 | 1 | 2 | P-value* | Pren** | OR [CI 95%]*** |
| rs2277798 | G/A Controls (n = 379) | 184 (48.55) | 132 (34.83) | 63 (16.62) | 448 (59.1) | 310 (40.9) | 0.0064 | 0.0128 | 0.75 [0.60–0.92] |
| rs9976767 | A/G Controls (n = 379) | 186 (49.08) | 136 (35.88) | 57 (15.04) | 458 (60.4) | 300 (39.4) | 0.0201 | 0.0201 | 1.28 [1.04–1.57] |

*All P-values have been calculated for the allelic model. **Benjamini & Hochberg (1995) step-up FDR control. ***Odds ratio for the minor allele.

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Table 3. Meta-analysis of two UBASH3a genetic variants within Spanish and German SLE populations.

| SNP       | Genotype, N (%) | Alleles, N(%) | Allele test |
|-----------|----------------|---------------|-------------|
|           | 1/2 Subgroup (N) | 1/1 | 1/2 | 2/2 | 1 | 2 | P-value* | Pren** | OR [CI 95%]*** |
| rs2277798 | G/A Controls (n = 1544) | 565 (44.63) | 543 (42.89) | 158 (12.48) | 1673 (66.1) | 859 (33.9) | 0.0047 | 4.7E-03 | 0.85 [0.76–0.95] |
| rs9976767 | A/G Controls (n = 1544) | 499 (32.32) | 744 (48.19) | 301 (19.49) | 1742 (56.4) | 1344 (43.6) | 1.2E-04 | 2.4E-04 | 1.23 [1.11–1.37] |

*All P-values have been calculated for the allelic model. **Benjamini & Hochberg (1995) step-up FDR control. ***Odds ratio for the minor allele.

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[7] for multiple testing was applied to the P-values in both the independent analysis and the combined meta-analysis (Pc). Linkage disequilibrium (LD) measurement (r²) between the studied SNPs was estimated by expectation-maximization algorithm using HAPLOVIEW (version 4.2; Broad Institute of MIT and Harvard). Finally, the dependency of the association between each SNP and every studied genetic variant was determined by a conditional logistic regression analysis (considering the different cohorts as covariate). The analyses were performed using PLINK (version 1.07) [8].

Results

The distributions of genotypic and allelic frequencies of the five UBASH3a evaluated polymorphisms were in HWE at 5% significance level. Additionally, MAFs of the studied SNPs were evaluated polymorphisms were in HWE at 5%. The distributions were similar to those reported by the HapMap project for the CEU population (http://hapmap.ncbi.nlm.nih.gov/) in both, Spanish and German cohorts. The LD structure of the five UBASH3a SNPs in the Spanish cohort is shown in (Figure S1). The Table 1 summarizes the results of the association analysis for the discovery cohort. The minor allele of the rs9976767 polymorphism exhibited a statistical significant association with SLE in the Spanish cohort (P = 9.9E-03, OR = 1.21, 95%CI = 1.07–1.37). In addition we observed a trend of association with the rs2277798 with lower risk to SLE (Pc = 2.4–04, OR = 0.85, 95%CI = 0.76–0.95).

Finally, we prompted out to evaluate whether one of both polymorphisms is responsible for the associations detected using a logistic regression analysis. Pair-wise conditional analysis showed that the association of the rs2277798 SNP was explained by the rs9976767 effect, because only the coefficient for the test of rs9976767 remained significant (model conditioned by rs2277798P = 0.76; model conditioned by rs9976767 P=9E-03, Table 4).

Discussion

UBASH3a is implicated in the regulation of tyrosine phosphorylation levels within T cells and is involved in facilitates the apoptosis induced in these cells. UBASH3a binds to the apoptosis-inducing protein AIF, which has previously been shown to function as a key factor of caspase-independent apoptosis [9]. It has also been reported that SLE T cells, compared with control T cells, undergo an increased rate of apoptosis, which contribute to SLE pathogenesis [4]. Changes in the UBASH3a structure or expression levels can affect the binding with AIF leading to an alteration in the apoptosis level.

Herein, we described for the first time the influence of five UBASH3a genetic variants in SLE susceptibility. Interestingly, the
rs9976767 polymorphism is located in the intronic region between the exons 5 and 6 while the other four studied SNPs (rs2277798, rs2277800, rs13048049 and rs17114930) are non-synonymous changes located in three different exons. The intronic regions flanking constitutive exons contain potential splicing regulatory sequences. Moreover, a study restricted to analysis of the canonical splice signals reported that 15% of point mutations disrupted splicing, a likely gross underestimate of the impact of splicing on human disease [10]. This suggests that the rs9976767 polymorphism could be affecting the expression of different UBAH3a isoforms consequently affecting the binding to AIF. Concerning to this we checked if there is any relation between the rs9976767 and expression of UBAH3a gene using expression quantitative trait loci (eQTL) databases. Interestingly, there is a significant statistical correlation between the increase of UBAH3a expression in lymphoblastoid cell lines and the homozygotes for the minor allele of rs9976767 (rho = 0.483, P = 1.3E-05; Figure S2A) in one of the two groups of twins studied (this observation was done using Genevar 3.2.0 software) [11,12]. Furthermore the eQTL studies in asthma showed that the SNPs (rs9784215, rs3746923, rs2277797) with highest LOD score (LOD > 4.5, P < 1E-05) in the UBAH3a locus are in moderate to high LD with rs9976767 (Figure S2B and C; this observation was done using mRNA by SNP Browser 1.0.1 http://www.sph.umich.edu/csg/liang/asthma/) [13,14]. This evidence suggested that rs9976767 could have a functional role in the regulation of the expression of UBAH3a. However, and according with HapMap project (http://hapmap.ncbi.nlm.nih.gov/), this SNP tags other six variants in this region (rs7278547, rs1702374, rs9976479, rs3746923, rs3761378, rs7283281; r2 > 0.95) and considering the present study and the previous GWAS [15,16] we have studied approximately 15% of the genetic variation of UBAH3a locus. In order to cover all the genetic variation of this gene, it is necessary to genotype 181 SNPs (calculated through an aggressive tagging with 2-marker haplotypes in Haploview 4.2 software using CEPh population from HapMap project). All these together suggest that the rs9976767 is a good functional candidate risk factor to SLE, but it could be more than one variant related to SLE.

No previous reports have associated the rs9976767 UBAH3a polymorphism with SLE. Nevertheless, it is worth noting that the rs9976767 SNP or its six tags variants were not included in previous genome wide association studies (GWAS) in Caucasian SLE cohorts [15,16]. Although the statistical power is 96% for our meta-analysis (calculate using a p value = 0.05; OR = 1.2; MAF = 0.4), the results found in study should be replicated in different Caucasian cohorts and other populations. Furthermore there is a need to determine whether the statistical associations are related with the involvement of UBAH3a in the pathogenesis of SLE and other autoimmune diseases. Regarding to this, the UBAH3a gene seems to be a common genetic factor in autoimmune diseases because different polymorphism of this locus has been associated with autoimmune diseases like T1D, CD, RA and vitiligo [6,17–21]. Our results showed that the minor allele of the rs9976767 UBAH3a polymorphism is a risk factor to SLE, as similarly observed with T1D [6]. Nevertheless, there is no evidence of association between this variant and other autoimmune diseases. This can be linked with the suggestion that common genetic factors in autoimmune diseases could match a regional level but differ in the specific genetic variant associated to each disease, like the associations observed with IL2–IL21 and MHC loci [22]. Based on the concept of quantitative thresholds for immune-cell signalling, the effect of the rs976767 UBAH3a variant could diversely affect the range of values for the stimulus-response selection of the immune cells in different autoimmune pathologies, making it more or less relevant in different diseases [3].

In conclusion, our study showed the first evidence of association of the UBAH3a gene with the genetic background of SLE. Together the functional role of the protein encoded by this gene, the reported data in the eQTLs databases and our results point to the UBAH3a gene as a new element in the pathogenic mechanism of autoimmune diseases.

### Supporting Information

**Figure S1** Pattern of linkage disequilibrium of the five studied SNPs and their location in the UBAH3a gene.

The values correspond to r2 calculated for the Spanish cohort. The rs2277798 polymorphism [G/A] is located in exon 1 of UBAH3a gene. It’s a no-synonymous change in the position 18 of the protein [S[Ser]/G[Glu]]. The rs2277800 polymorphism [C/T] is also located in exon 1 of UBAH3a gene and generate a change in the position 28 of the protein [L[Leu]/F[Phe]]. In the other hand, the rs9976767 [A/G] is an intronic variant located between the exons 5 and 6 of the UBAH3a gene. Both variants rs13048049 [G/A] and rs17114930 [C/G] are no-synonymous changes in exons 7 and 11, respectively. The first one produce a change from arginine (R[Arg]) to glutamine (Q[Gln]) in position 296; while the rs17114930 polymorphism generates a change from aspartic acid (D[Asp]) to glutamic acid (E[Glu]) in position 428 in Caucasian population.

**Figure S2** Results observed using different expression quantitative trait loci (eQTL) tools to evaluate if there is any relationship between the rs9976767 variant and the UBAH3a expression (A) SNP-gene association plot for the rs9976767 and the UBAH3a gene based on Spearman’s rank correlation coefficient (rho) using the Genevar 3.2 software (http://www.sanger.ac.uk/resources/software/genevar/). The eQTL analysis was performed in lymphoblastoid cell lines from peripheral donors.

| Group of analysis | SNP     | MAF Cases | MAF Controls | p Value: add to rs9976767 | rs9976767 p value: add to SNP | r2 with rs9976767 |
|------------------|---------|-----------|--------------|---------------------------|-------------------------------|------------------|
| SLE              | rs2277798 | 0.34      | 0.38         | 0.758                     | 0.0087                        | 0.45             | 0.41             |

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Table 4. Conditional logistic regression analysis for two UBAH3a SNPs located in SLE considering the two European populations as covariate.
blood sample (n = 74). The plot corresponds to one of the two twins groups studied [2]. (B) Linkage disequilibrium (LD) plot performed in Haploview 4.2 [3]. LD plot between rs9976767 and the rs9784215, rs3746923, rs2277797 SNPs which exhibited the highest LOD score (LOD = 4.5, P < 1E-05) in the UBASH3a locus showed in (C) Snapshot of observed eQTLs related with UBASH3a gene from the mRNA by SNP Browser 1.0.1 software [http://www.sph.umich.edu/csg/liang/asthma/] based on eQTL studies in asthma [4,5]. The LOD scores and P values for those SNPs are: rs9784215, LOD = 4.909 P = 2E-06; rs3746923, LOD = 4.905 P = 2E-06; rs2277797, LOD = 4.68 P = 3.4E-06. They are signalled as red dots in the LOD plot. 1. Yang TP, Beazley C, Montgomery SB, Dimas AS, Gutierrez-Arcelus M, et al. (2010) Genevar: a database and Java application for the analysis and visualization of SNP-gene associations in eQTL studies. Bioinformatics 26: 2474-2476. 2. Nica AC, Parts L, Glass D, Nisbet J, Barrett A, et al. (2011) The architecture of gene regulatory variation across multiple human tissues: the MuTHER study. PLoS Genet 7: e1002003. 3. Barrett JC, Fry B, Maller J, Daly MJ (2005) Haploview: analysis and visualization of LD and haplotype maps. Bioinformatics 21: 263-265. 4. Dixon AL, Liang L, Moffatt MF, Chen W, Heath S, et al. (2007) A genome-wide association study of global gene expression. Nat Genet 39: 1202-1207. 5. Moffatt MF, Kabesch M, Liang L, Dixon AL, Strachan D, et al. (2007) Genetic variants regulating ORMDL3 expression contribute to the risk of childhood asthma. Nature 448: 470-473. (TIF)

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