Surfactant protein A genetic variants associate with severe respiratory insufficiency in pandemic influenza A virus infection

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Abstract

Introduction: Inherited variability in host immune responses influences susceptibility and outcome of Influenza A virus (IAV) infection, but these factors remain largely unknown. Components of the innate immune response may be crucial in the first days of the infection. The collectins surfactant protein (SP)-A1, −A2, and -D and mannose-binding lectin (MBL) neutralize IAV infectivity, although only SP-A2 can establish an efficient neutralization of poorly glycosylated pandemic IAV strains.

Methods: We studied the role of polymorphic variants at the genes of MBL (MBL2), SP-A1 (SFTPA1), SP-A2 (SFTPA2), and SP-D (SFTPD) in 93 patients with H1N1 pandemic 2009 (H1N1pdm) infection.

Results: Multivariate analysis showed that two frequent SFTPA2 missense alleles (rs1965708-C and rs1059046-A) and the SFTPA2 haplotype 1A0 were associated with a need for mechanical ventilation, acute respiratory failure, and acute respiratory distress syndrome. The SFTPA2 haplotype 1A1 was a protective variant. Kaplan-Meier analysis and Cox regression also showed that diplotypes not containing the 1A1 haplotype were associated with a significantly shorter time to ICU admission in hospitalized patients. In addition, rs1965708-C (P = 0.0007), rs1059046-A (P = 0.0007), and haplotype 1A0 (P = 0.0004) were associated, in a dose-dependent fashion, with lower PaO2/FiO2 ratio, whereas haplotype 1A1 was associated with a higher PaO2/FiO2 ratio (P = 0.001).

Conclusions: Our data suggest an effect of genetic variants of SFTPA2 on the severity of H1N1pdm infection and could pave the way for a potential treatment with haplotype-specific (1A1) SP-A2 for future IAV pandemics.

Introduction

Influenza A virus (IAV) infection is usually a mild and self-limited condition. Likewise, infection with the H1N1 pandemic 2009 (H1N1pdm) IAV often results in an uncomplicated flu, but, in a small subset of patients, it may rapidly evolve to primary viral pneumonia, acute respiratory failure (ARF), and acute respiratory distress syndrome (ARDS) [1]. Inherited and acquired variability in host immune responses may influence susceptibility and outcome of IAV infection, although these factors remain largely unknown [2-4]. Before exposure to H1N1pdm IAV, most individuals, particularly those born after 1957, lack serum antibodies capable of neutralizing the virus [1]. Adaptive immune responses, which are needed for ultimate viral clearance, take several days to develop. Therefore, components of the innate immunity that are able to neutralize IAV infection with minor inflammation may be crucial for host defense in the first few days after infection. Different soluble pattern-recognition molecules of the innate immunity

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can neutralize IAV infection. Several secreted human C-type lectins of the collectin family, the serum mannoselbinding lectin (MBL), the pulmonary surfactant proteins (SP) –A1, –A2, and –D, and collectin 11 (CL-11, alias collectin kidney 1, CL-K1) may neutralize IAV infectivity in vitro [5-8]. Among these collectins, the SPs have been shown to exert an important role against IAV infection in animal models: mice lacking SP-A or SP-D have increased susceptibility to IAV infection, and their role seem to depend on IAV strains, particularly pandemic versus seasonal strains [9-14]. In addition, variability at the collectin genes, MBL2 (Ensembl: ENSG00000165471), SFTPA1 (Ensembl: ENSG00000122852), SFTPA2 (Ensembl: ENSG00000185303), and SFTPD (Ensembl: ENSG00000133661) have been found to be associated with susceptibility to and/or severity of several bacterial and viral infectious diseases [6,15].

It hence follows that these collectins are firm candidates to explain, at least in part, the role of host genetic variability in the defense against IAV infection. The human SP-A locus consists of two similar genes, SFTPA1 and SFTPA2, localized within a cluster (10q21-24) that includes the SP-D gene (SFTPD) [16]. MBL2 was reported not to be in physical linkage with the genes of these SPs [17], but linkage disequilibrium of MBL2 with SFTPA1 and SFTPA2 has been shown; and LD among variants at these genes could influence the results of genetic-association studies [6,18,19].

In the present study, we assessed the role of variants at the SFTPA1, SFTPA2, and SFTPD genes in H1N1pdm IAV infection in humans. Variants at the neighbor collectin gene MBL2 were also analyzed.

Materials and methods
H1N1pdm-infected patients
We recruited 124 patients with H1N1pdm infection between July 2009 and November 2011. Thirty-one individuals with ancestors other than Spanish were excluded. Of 93 unrelated white Spanish patients, 70 were hospitalized at five tertiary Spanish hospitals, and 23 were attended at primary care centers (Figure 1). Data and samples from all ambulatory patients and from 30% of hospitalized individuals were retrospectively obtained; in the remaining patients, data were obtained prospectively. All patients were treated with oseltamivir, and only one patient had been previously vaccinated against H1N1pdm.

Diabetes, previous lung disease, solid-organ transplantation, immunosuppression, body mass index (BMI) ≥30, human immunodeficiency virus (HIV) infection, and pregnancy were considered risk factors. Sepsis, septic shock, and multiorgan dysfunction syndrome (MODS) were defined by using the American College of Chest Physicians/Society of Critical Care Medicine criteria [20]. ARF and ARDS were diagnosed according to the American European Consensus Conference Definition [21]. Functional parameters of gas exchange were calculated on the basis of the ratio of oxygen arterial pressure to oxygen inspiratory fraction (PaO2/FiO2). In hospitalized patients without arterial blood gas analysis (n = 18), ARF was established when hemoglobin oxygen saturation, breathing room air, was lower than 90%. In patients with intensive care unit (ICU) admission, severity of illness was evaluated with the Acute Physiology and Chronic Health Evaluation II score, taking the worst reading of the first 24 hours in the ICU.
All steps were performed in complete accordance to the Helsinki declaration. The protocol was approved by Clinical Research Ethics Committees (CEIC) of hospitals involved (CEIC Hospital Universitario de Gran Canaria Doctor Negrín, Las Palmas de Gran Canaria; CEIC Hospital San Jorge, Huesca and CEIC Hospital Clínico y Universitario de Valencia, Valencia). Informed consent was obtained of all patients included.

Detection of H1N1pdm by real-time polymerase chain reaction

Influenza A H1N1 virus in the 124 patients was detected in nasopharyngeal swabs by using the Real-Time ready Influenza A (H1N1) Detection Set (Roche Diagnostics GmbH, Mannheim, Germany) according to manufacturer’s protocol.

General Spanish-population subjects

The general population group consisted of white unrelated Spanish healthy volunteers (blood and bone marrow donors as well as hospital staff) from four tertiary Spanish hospitals. The control group was analyzed for MBL2 (<i>n</i> = 1,736) and SFTPA1 and SFTPA2 (<i>n</i> = 769) polymorphisms in previous studies [18,19]. For the nine SFTPD variants under study, 963 individuals from the general population group recruited at the same hospitals were analyzed in this study. The protocol was approved by Clinical Research Ethics Committees (CEIC) of hospitals involved (CEIC Hospital Universitario de Gran Canaria Doctor Negrín, Las Palmas de Gran Canaria; CEIC Hospital de la Princesa, Madrid; CEIC Hospital San Jorge, Huesca and CEIC Hospital Clínico y Universitario de Valencia, Valencia). Informed consent was obtained of the general-population individuals included. No data about whether individuals from the general Spanish population group were or were not infected with H1N1pdm were available. Foreigners and individuals with ancestors other than Spanish were previously excluded.

Selection of single-nucleotide polymorphisms

Deficient and low MBL serum levels are mainly due to the presence of three common point mutations in the exon 1 of the MBL2 gene: alleles <i>B</i> (rs1800450), <i>C</i> (rs1800451), and <i>D</i> (rs5030737) are termed <i>O</i> alleles, <i>A</i> being the wild-type. Serum MBL levels are very low or absent in individuals homozygous for <i>O</i> alleles. In addition, the presence of the promoter allele <i>X</i> (rs7096206) has an important down-regulating effect, and individuals with <i>XA</i>/<i>O</i> also have very low MBL serum levels. <i>O/O</i> together with <i>XA/O</i> genotypes are considered MBL-deficient genotypes, which are common in most populations.

The human SP-A locus consists of two similar genes, SFTPA1 and SFTPA2, located on chromosome 10q22.3, within a cluster that includes the SP-D gene (<i>SFTP</i>D). In addition, a certain degree of linkage disequilibrium (LD) exists among SP genes and the MBL gene (10q21-24) [18,19]. On the basis of the existence of several SNPs, SP-A haplotypes are conventionally denoted as 6<i>A</i> for the SFTPA1 gene (V19A, rs1059047; L50V, rs1136450; R219W, rs4253527) and 1<i>A</i> for the SFTPA2 gene (T9N, rs1059046; A91P, rs17886395; Q223K, rs1965708) [22]. These missense single-nucleotide polymorphisms (SNPs) at SFTPA1 and SFTPA2 were analyzed in our study. The most frequent conventional haplotypes of these genes, except 1<i>A</i> and 1<i>A</i>5, can be unambiguously identified.

For the analysis of SFTP<i>D</i>, genotypic data of individuals of European ancestry (CEU) from the International HapMap Project [23] were used to select LD tagging SNPs. Pairwise LD-tagging was achieved with Haploview v. 4.2 [24] for SNPs with a minimum minor allele frequency of 0.05 and <i>r</i>2 of 0.8. As result, three intronic SNPs (rs10887199, rs7078012, and rs723192) and one synonymous SNP (rs6413520) were selected. Three missense SNPs were also analyzed: rs3088308 (S290T), rs2243639 (T180A), and rs721917 (M31T). Additionally, we studied an SNP in the SFTP<i>D</i> promoter region (rs1885551) that was recently reported to have a profound impact on serum SP-D levels [25] and an intronic SNP (rs17886286) associated with susceptibility to invasive pneumococcal disease [26].

DNA extraction and genotyping

Genomic DNA was extracted from 400 μl of peripheral blood by using iPrep PureLink gDNA Blood kit in the iPrep Purification Instrument (Invitrogen by Life Technologies, Carlsbad, CA, USA). Extracted DNA integrity was checked by NanoDrop ND-1000 (NanoDrop Technologies, Wilmington, DE, USA). In total, 19 polymorphisms at MBL2, SFTPA1, SFTPA2 and SFTP<i>D</i> were studied.

MBL2, SFTPA1, SFTPA2, and SFTP<i>D</i> rs721917 SNPs were analyzed by means of PCR-RFLP and PCR-SSP techniques, as previously described [18,19]. The other eight SNPs of SFTP<i>D</i> were determined by predesigned Taqman SNP genotyping assays (Assays IDs: C__26726209_10, C__26726205_10, C__31530298_10, C__63652102_10, C__29213175_10, C__1362981_20, C__630297_10, and C__12145257_20), according to the manufacturer’s protocol, with commercially available reagents by means of ViIA™ 7 Real-time PCR System (Applied Biosystems, Foster City, CA, USA).

Data analysis

The Hardy-Weinberg equilibrium was analyzed with Haploview v. 4.2. Haplotypes were estimated in silico with PHASE v. 2.1.1 under 1,000 permutations. Statistical analyses were performed by using SPSS 20.0 (SPSS, Inc., Chicago, IL, USA). The comparison of all genotype
distributions based on susceptibility and severity was performed by using the \( \chi^2 \) test or Fisher Exact test when needed, and odds ratios with 95% confidence intervals were calculated. The relation between severity in hospitalized patients and genotypes was evaluated by binary logistic regression models: age, gender, risk factors, and secondary infection (either bacteremia or secondary bacterial pneumonia) were included as independent variables.

Comparison of the \( \text{PaO}_2/\text{FiO}_2 \) ratio according to genetic variants was performed by a using linear regression model adjusted by age, gender, risk factors, and secondary infection (either bacteremia or secondary bacterial pneumonia).

To assess the need for ICU admission, a multivariate analysis was performed, including the aforementioned variables, by conditional logistic regression stratified by hospital of origin.

Log-rank (LR) \( \chi^2 \) tests were calculated to compare ICU admission according to the distribution of genetic variants. Cox proportional hazard ratios adjusted for the independent variables age, gender, risk factors, development of secondary bacterial infection (pneumonia or bacteremia), and hospital of origin were also performed.

**Results**

Demographic and clinical characteristics of patients are shown in Table 1. The genotype distribution of SNPs did not differ significantly under conditions of Hardy-Weinberg equilibrium in all of the groups studied (data not shown). Frequencies of the genetic variants under analysis were not found to be significantly different between H1N1pdm-infected patients and the general population (Table 2).

When severity of infection was evaluated in hospitalized patients (n = 70) (Table 3), we observed that two missense variants at \( \text{SFTPA2} \) (rs1965708-C and rs1059046-A) were significantly associated with the need for mechanical ventilation (MV) and with development of ARF and ARDS. Most of these associations remained significant after multivariate analysis adjusted for age, gender, risk factors, and secondary infection (either bacteremia or secondary bacterial pneumonia). The alleles rs1965708-C and rs1059046-A were overrepresented in patients requiring MV (\( P = 0.003; \text{OR}, 2.43; \) and \( P = 0.016; \text{OR}, 3.71, \) respectively), and in those who developed ARF (\( P = 0.006; \text{OR}, 4.09; \) and \( P = 0.005; \text{OR}, 3.70, \) respectively) or ARDS (\( P = 0.006; \text{OR}, 17.68; \) and \( P = 0.016; \text{OR}, 3.71, \) respectively). The variant rs1965708-C was also associated with septic shock (\( P = 0.007; \text{OR}, 17.16. \) The \( \text{SFTPA1} \) allele rs1136450-G was also associated, although to a lower extent, with the development of ARF (\( P = 0.038; \text{OR}, 2.64 \) and the use of MV (\( P = 0.040; \text{OR}, 2.55; \) these associations may be secondary to the existence of LD between \( \text{SFTPA1} \) and \( \text{SFTPA2} \) variants (Figure 2).

| Characteristics | Subjects |
|-----------------|---------|
| Age (years)     | 43.5 ± 18.0* |
| Gender (male)   | 52 (55.9) |
| Hospital admission |
| No              | 23 (24.7) |
| Yes             | 70 (75.3) |
| PVP             |
| No              | 40 (43.0) |
| Yes             | 53 (57.0) |
| ICU admission |
| No              | 63 (67.7) |
| Yes             | 30 (32.3) |
| ARF             |
| No              | 45 (48.4) |
| Yes             | 48 (51.6) |
| Shock           |
| No              | 77 (82.8) |
| Yes             | 16 (17.2) |
| ARDS            |
| No              | 77 (82.8) |
| Yes             | 16 (17.2) |
| MODS            |
| No              | 87 (93.5) |
| Yes             | 6 (0.65) |
| MV              |
| No              | 69 (74.2) |
| Yes             | 24 (25.8) |
| Hospital mortality |
| No              | 89 (95.7) |
| Yes             | 4 (0.43) |
| Risk factor     |
| No              | 32 (34.4) |
| Yes             | 61 (65.6) |
| Secondary bacterial infectionb |
| No              | 80 (86.0) |
| Yes             | 13 (14.0) |

ARDS, acute respiratory distress syndrome; ARF, acute respiratory failure; ICU, intensive care unit; MODS, multiorgan dysfunction syndrome; MV, mechanical ventilation; PVP, Primary viral pneumonia.

aData are presented as mean ± SD or number of individuals (%).bTen patients had secondary bacterial pneumonia, and seven patients had bacteremia (four of them with secondary bacterial pneumonia).

When haplotypes, based on combinations of missense SNP, were analyzed (Table 4), the \( \text{SFTPA2} \) haplotype \( I A^b \) was found to be associated with the need for MV and with the development of ARF and ARDS in hospitalized patients. These differences were found to be independent of age,
gender, risk factors, and secondary infection (either bacteremia or secondary bacterial pneumonia) in multivariate analysis: $P = 0.003$ (data in Table 4), OR 3.73; $P = 0.022$; OR, 2.81; and $P = 0.016$; OR, 3.41, for the need for MV, ARF, and ARDS comparisons, respectively. By contrast, the $SFTPA2$ haplotype 1A1 was found to protect against the requirement for MV and the development of ARF and ARDS in the multivariate analysis ($P = 0.015$; OR, 0.16; $P = 0.005$; OR, 0.21; and $P = 0.021$; OR, 0.12).

As an additional measurement of severity, we analyzed the need for ICU admission in hospitalized patients. The variants rs1965708-C and rs1059046-A were associated with a higher rate of ICU admission, although only rs1965708-C remained significant after conditional logistic regression adjustment for the previously mentioned variables and stratified by hospital of origin ($P = 0.031$; OR, 4.33) (Table 3). The same analysis remained significant at $SFTPA2$ haplotype 1A1 in a dominant model ($P = 0.029$; OR, 0.32) (Table 4). It is noteworthy that Kaplan-Meier analysis and the log-rank test showed that the variants rs1965708-C and rs1059046-A were associated with a significantly shorter time to ICU admission in hospitalized patients (Figure 3). The same analysis also showed that ICU admission was less frequently required among patients with the haplotype 1A1, and this difference was readily detected in the first days after hospitalization. The effect of these variants at the time of ICU admission remained significant in Cox regression.

We finally evaluated the influence of genetic variants of $SFTPA1$ and $SFTPA2$ on respiratory gas exchange in 52 hospitalized patients. $SFTPA2$ alleles (rs1965708-C and rs1059046-A) and haplotypes (1A$^e$) predisposing to increased severity were found to be associated with significantly lower PaO$_2$/FiO$_2$ ratios, independent of age, gender, risk factors, secondary bacterial pneumonia, and bacteremia (Figure 4). Interestingly, the effect of these alleles or haplotypes was found to be dependent on the

### Table 2 Distribution of genotype frequencies of collectin genes in general Spanish population and H1N1pdm-infected patients

| Variants | Genotypes | General Spanish population | H1N1pdm-infected patients |
|----------|-----------|----------------------------|----------------------------|
| **MBL2** |           | n = 1736                   | n = 93                     |
| rs1800451 (G57E) | AA/AO/OO | 1,032 (59.4)/615(35.4)/89 (5.1) | 54 (58.1)/36 (38.7)/3 (3.2) |
| rs1800450 (G54D) | AA/AO/OO | 1,032 (59.4)/615(35.4)/89 (5.1) | 54 (58.1)/36 (38.7)/3 (3.2) |
| rs5030737 (RS2C) | AA/AO/OO | 1,032 (59.4)/615(35.4)/89 (5.1) | 54 (58.1)/36 (38.7)/3 (3.2) |
| rs7096206 (Prom) | AA/AO/OO | 1,032 (59.4)/615(35.4)/89 (5.1) | 54 (58.1)/36 (38.7)/3 (3.2) |
| MBL deficiency | AA + YAO/XXAO + YAO | 1,475 (85.0)/261 (15.0) | 77 (82.8)/16 (17.2) |
| **SFTPA2** |           | n = 769                    | n = 93                     |
| rs1965708 (Q223K) | TT/TC/CC | 22 (2.9)/244 (31.7)/503 (65.4) | 4 (4.3)/33 (35.5)/66 (60.2) |
| rs17886395 (A91P) | TT/TC/CC | 22 (2.9)/244 (31.7)/503 (65.4) | 4 (4.3)/33 (35.5)/66 (60.2) |
| rs1059046 (T9N) | TT/TC/CC | 22 (2.9)/244 (31.7)/503 (65.4) | 4 (4.3)/33 (35.5)/66 (60.2) |
| **SFTPA1** |           | n = 769                    | n = 93                     |
| rs1965708 (Q223K) | TT/TC/CC | 22 (2.9)/244 (31.7)/503 (65.4) | 4 (4.3)/33 (35.5)/66 (60.2) |
| rs17886395 (A91P) | TT/TC/CC | 22 (2.9)/244 (31.7)/503 (65.4) | 4 (4.3)/33 (35.5)/66 (60.2) |
| rs1059046 (T9N) | TT/TC/CC | 22 (2.9)/244 (31.7)/503 (65.4) | 4 (4.3)/33 (35.5)/66 (60.2) |
| **SFTPD** |           | n = 963                    | n = 93                     |
| rs3088308 (S200T) | AA/AT/TT | 829 (86.1)/129 (13.4)/5 (0.5) | 77 (82.8)/14 (15.1)/2 (0.2) |
| rs2243639 (T180A) | TT/TC/CC | 373 (38.7)/449 (46.6)/141 (14.6) | 42 (45.2)/45 (42.5)/9 (9.6) |
| rs10887199 (Intr) | TT/TC/CC | 373 (38.7)/449 (46.6)/141 (14.6) | 42 (45.2)/45 (42.5)/9 (9.6) |
| rs17886268 (Intr) | CC/CG/G | 759 (78.8)/189 (19.6)/15 (1.6) | 71 (76.3)/20 (21.5)/2 (0.2) |
| rs7078012 (Intr) | CC/CT/TT | 759 (78.8)/189 (19.6)/15 (1.6) | 71 (76.3)/20 (21.5)/2 (0.2) |
| rs6413520 (S455) | AA/AG/GG | 824 (85.6)/134 (13.9)/5 (0.5) | 82 (88.2)/11 (11.8)/0 (0.0) |
| rs721917 (M31T) | TT/TC/CC | 356 (37.0)/438 (45.5)/169 (17.5) | 36 (38.7)/38 (40.9)/19 (20.4) |
| rs723192 (Intr) | CC/CT/TT | 356 (37.0)/438 (45.5)/169 (17.5) | 36 (38.7)/38 (40.9)/19 (20.4) |
| rs1885531 (Prom) | AA/AG/GG | 766 (79.5)/182 (18.9)/15 (1.6) | 72 (77.4)/19 (20.4)/2 (0.2) |

*O/O together with XA/O genotypes are considered MBL-deficient genotypes. Intr, intronic region; Prom, promoter region. SNPs were added on the basis of chromosome position.
number of alleles present in a genotype or diplotype: \( P = 0.0007, P = 0.0007 \) and \( P = 0.0004 \) for rs1965708-C, rs1059046-A, and \( I A^Q \) respectively. The \( I A^Q \) haplotype, which was associated with lesser severity, was also associated with higher \( PaO_2/F_iO_2 \) ratios (\( P = 0.0007 \)), and this effect was also found to be dose-dependent in a linear regression model adjusted for the same independent variables (\( P = 0.0014 \)) (Figure 4).

**Discussion**

Glycosylation of hemagglutinins may be an important mechanism by which IAV can evade recognition by antibodies in an immune population. By contrast, glycosylation of hemagglutinins is important in determining sensitivity of IAV to recognition by collectins [7,27]. SP-D and MBL bind to mannose-rich glycans on the hemagglutinins and neuraminidase glycoproteins of IAV, agglutinating viral particles and inhibiting infectivity and neuraminidase activity [7]. Among the known human collectins, SP-D has the strongest in vitro capability of aggregating and neutralizing activity of IAV [7,12,27]. Hemagglutinins from all available strains of pandemic influenza viruses show significantly lower glycosylation sites compared with seasonal strains; and pandemic viruses, particularly H1N1pdm, were found

### Table 3 Severity of H1N1pdm infection in hospitalized patients regarding missense single-nucleotide polymorphisms of SFTPA2 and SFTPA1

| Polymorphism       | Genotype frequencies | Statistical significance | Genotype comparisons | Allele comparisons |
|--------------------|----------------------|--------------------------|----------------------|-------------------|
|                    | CC                   | CA                       | AA                   | CC vs CA + AA     | C vs A            |
|                    | OR (95% CI)          | OR (95% CI)              | OR (95% CI)          | OR (95% CI)       |
| rs1965708 (Q223K)  |                      |                          |                      |                   |
| ARF                | 48                   | 33 (0.69)                | 15 (0.31)            | 0 (0.00)          | 0.027             |
|                   |                      |                          |                      |                   | 0.018             |
| No ARF             | 22                   | 9 (0.41)                 | 10 (0.46)            | 3 (0.14)          | 3.18 (1.12-9.01)  |
|                   |                      |                          |                      |                   | 5.40 (1.34-21.71) |
| ARDS               | 16                   | 14 (0.88)                | 2 (0.13)             | 0 (0.00)          | 6.45 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.81 (1.45-42.05) |
| No ARDS            | 54                   | 28 (0.52)                | 23 (0.43)            | 3 (0.06)          | 4.86 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.64 (1.41-41.44) |
| Shock              | 16                   | 14 (0.88)                | 2 (0.13)             | 0 (0.00)          | 6.45 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.81 (1.45-42.05) |
| No shock           | 54                   | 28 (0.52)                | 23 (0.43)            | 3 (0.06)          | 4.86 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.64 (1.41-41.44) |
| MV                 | 24                   | 20 (0.83)                | 4 (0.17)             | 0 (0.00)          | 6.45 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.81 (1.45-42.05) |
| No MV              | 46                   | 22 (0.48)                | 21 (0.46)            | 3 (0.07)          | 5.46 (1.61-18.47) |
|                   |                      |                          |                      |                   | 18.60 (2.23-155.24) |
| ICU                | 30                   | 23 (0.77)                | 7 (0.23)             | 0 (0.00)          | 6.45 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.81 (1.45-42.05) |
| No ICU             | 40                   | 19 (0.48)                | 18 (0.45)            | 3 (0.08)          | 3.64 (1.27-10.42) |
|                   |                      |                          |                      |                   | 2.98 (1.19-7.46)  |
| rs1059046 (T9N)    |                      |                          |                      |                   |
| ARF                | 48                   | 20 (0.42)                | 24 (0.50)            | 4 (0.08)          | 6.45 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.81 (1.45-42.05) |
| No ARF             | 22                   | 3 (0.14)                 | 14 (0.64)            | 5 (0.23)          | 4.52 (1.18-17.24) |
|                   |                      |                          |                      |                   | 8.02 (1.61-39.98) |
| ARDS               | 16                   | 9 (0.56)                 | 6 (0.38)             | 1 (0.06)          | 6.45 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.81 (1.45-42.05) |
| No ARDS            | 54                   | 14 (0.26)                | 32 (0.59)            | 8 (0.15)          | 5.46 (1.61-18.47) |
|                   |                      |                          |                      |                   | 18.60 (2.23-155.24) |
| Shock              | 16                   | 11 (0.46)                | 12 (0.50)            | 1 (0.04)          | 6.45 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.81 (1.45-42.05) |
| No shock           | 54                   | 12 (0.26)                | 26 (0.57)            | 8 (0.17)          | 6.45 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.81 (1.45-42.05) |
| MV                 | 24                   | 11 (0.46)                | 12 (0.50)            | 1 (0.04)          | 6.45 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.81 (1.45-42.05) |
| No MV              | 46                   | 14 (0.47)                | 14 (0.47)            | 2 (0.07)          | 3.64 (1.27-10.42) |
| ICU                | 30                   | 14 (0.47)                | 14 (0.47)            | 2 (0.07)          | 3.64 (1.27-10.42) |
| No ICU             | 40                   | 9 (0.23)                 | 24 (0.60)            | 7 (0.18)          | 3.64 (1.27-10.42) |
| rs1136450 (L50V)   |                      |                          |                      |                   |
| ARF                | 48                   | 21 (0.44)                | 25 (0.52)            | 2 (0.04)          | 6.45 (1.35-31.25) |
|                   |                      |                          |                      |                   | 7.81 (1.45-42.05) |
| No ARF             | 22                   | 5 (0.23)                 | 12 (0.55)            | 5 (0.23)          | 6.77 (1.20-38.22) |
|                   |                      |                          |                      |                   | 19.80 (1.64-238.70) |
| ARDS               | 16                   | 11 (0.46)                | 13 (0.54)            | 0 (0.00)          | 6.77 (1.20-38.22) |
|                   |                      |                          |                      |                   | 19.80 (1.64-238.70) |
| No ARDS            | 46                   | 15 (0.33)                | 24 (0.52)            | 7 (0.15)          | 6.77 (1.20-38.22) |
|                   |                      |                          |                      |                   | 19.80 (1.64-238.70) |

\( ^a \)P value for the bivariate comparison calculated with the \( \chi^2 \) test. \( ^b \)P value for the multivariate analysis calculated with binary logistic regression, including the variables age, gender, risk factors, secondary bacterial pneumonia, and bacteremia. \( ^c \)P value by Fisher exact test. \( ^d \)P value for the multivariate analysis calculated with binary logistic regression, including the variables age, gender, risk factors, and secondary bacterial pneumonia (bacteremia variable was excluded because it shows a co-linear relation with the variable mechanical ventilation). \( ^e \)Patients who required (ICU) or did not require (No ICU) ICU admission, \( ^f \)P value for the multivariate analysis calculated with conditional logistic regression stratified by hospital of origin, including the co variables age, gender, risk factors, secondary bacterial pneumonia, and bacteremia. Only those comparisons with \( P < 0.05 \) and significant odds ratios were included. OR (95% CI), Odds ratio (95% confidence interval); ARF, acute respiratory failure; ARDS, acute respiratory distress syndrome; MV, mechanical ventilation; ICU, intensive care unit.
to be resistant to the antiviral activities of SP-D, MBL, and the pentraxin PTX3 [7,13,27]. We have not observed any association with severity when deficient-, low-, or high-MBL producer genotypes and SFTPD SNPs or haplotypes were compared (data not shown). Interestingly, SP-A binding to hemagglutinins and SP-A-dependent IAV neutralization in vitro are not influenced by the extent of hemagglutinins glycosylation. SP-A is slightly more effective than SP-D at neutralizing nonglycosylated IAV strains, and it neutralizes IAV strains resistant to SP-D [12].

Accordingly, our results point toward a role of SFTPA2 SNPs and haplotypes, particularly haplotypes IA0 and IA4, in the severity of H1N1pdm infection. The rationale for such an association seems to be the involvement of SFTPA2 variants in gas-exchange parameters, which, in the case of pulmonary IAV infection, is largely dependent on IAV-induced lung inflammation.

Besides its role in IAV neutralization, the hydrophilic SP-A and -D have been shown to have an antiinflammatory function. Binding of the carbohydrate-binding recognition domains (CRDs) to signal-inhibitory regulatory protein α (SIRPα) on alveolar macrophages suppresses NF-κB activation and inflammation, allowing the healthy lung to remain in a quiescent state. SP-A and SP-D can also inhibit inflammation, through the CRD, blocking Toll-like receptors 2 and 4. Pulmonary clearance of IAV was reduced, and pulmonary inflammation and severity were increased in Sftpa−/− mice compared with wild-type mice [10,11]. SP-A was also found to disturb the host inflammatory response to IAV infection in mouse models without directly influencing viral growth and spread, and even without demonstrable viral binding, at least when the virus was resistant to neutralization by SP-D [9]. Insufficient amounts of surfactant, particularly SP-A, have been observed in prematurely born infants with respiratory distress syndrome (pRDS). The haplotype IA0 or SFTPA1-SFTPA2 haplotypes containing IA0 have been repeatedly associated with the development of pRDS, whereas SP haplotypes containing IA4 were found to be protective against that condition [5,15]. These findings parallel those observed in our study, suggesting that IA0 and IA4 might influence the inflammatory response and the severity of H1N1pdm infection without binding to IAV.

Human SP-A1 and −A2 have similar in vitro hemagglutination-inhibition activity of IAV strains exhibiting non- or poorly glycosylated hemagglutinin heads [14]. However, in all functional assays, SP-A2 is more active than SP-A1 [5,15]. So it is not surprising that among all the genetic variants of collectins analyzed in our study, only a few alleles and haplotypes of SFTPA2 were associated with H1N1pdm severity in hospitalized patients. Furthermore,
the effect of the SFTPA2 variants on the need for ICU admission was detected in the first few days of hospitalization, as would be expected for components of the innate immunity involved in inflammation and defense against IAV infection.

The significance of the functional differences between variants at SFTPA1 and SFTPA2 is poorly understood [5,15]. The variant rs1965708 produces an amino-acid change at residue 223 (Q223K), located in the CRD of SP-A2, and might directly influence the binding properties to either IAV or the antiinflammatory receptor SIRPa. Residue 9 (rs1059046, T9N) is located in the signal peptide, and it is unknown whether this variant may affect the protein. It is, however, worth noting that haplotypes 1A0 and 1A1 differ precisely at residues 9 and 223. A role of SNPs at regulatory regions in haplotypes 1A0 and 1A1 in translation and/or RNA stability of SFTPA2 cannot be ruled out [5,15]. Interestingly, SP-A2 1A1 variants were shown to have a lower activity for enhancement of TNF-α in macrophage-derived THP-1 cells than other variants, such as 1A and 1A0, particularly after oxidative stress [28,29]. These data suggest that the protective effect of the SP-A2 1A1 variant in the severity of H1N1pdm infection could be due to its lower proinflammatory activity.

Irrespective of the causal SNP(s), our data suggest that the haplotype 1A0 of SFTPA2 is associated with a higher severity after H1N1pdm infection, whereas the SFTPA2 haplotype 1A1 was associated with a dominant protective effect against severe forms of H1N1pdm infection.

We acknowledge several limitations of our study. First, the overrepresentation of hospitalized patients, could bias us to analyze susceptibility to H1N1pdm infection. In addition, our control group could be considered a representative sample from the Spanish population rather than a representation of non-H1N1pdm-infected individuals.

Second, our study is underpowered to detect the role of some variants on severity of H1N1pdm infection. Nevertheless, considering odds ratios and a significance level of 5%, the power of the associations of rs1965708-C with the development of ARDS and septic shock was

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**Table 4 Severity of H1N1pdm infection in hospitalized patients regarding haplotypes of SFTPA2**

| Haplotype | Diplotype frequencies | Diplotype comparisons | Haplotype comparisons |
|-----------|-----------------------|-----------------------|-----------------------|
|           | 1A0/1A0 | 1A0/rest | Rest/rest | 1A0/1A0 vs 1A0/rest + rest/rest | OR (95% CI) | OR (95% CI) | OR (95% CI) | OR (95% CI) |
| ARF       | 48      | 16 (0.33) | 27 (0.56) | 5 (0.10) | 0.031 | 0.045 | 0.023 | 0.022 |
| No ARF    | 22      | 2 (0.09)  | 14 (0.64) | 6 (0.27) | 5.00 (1.04-24.10) | 6.38 (1.04-38.97) | 2.30 (1.11-4.76) | 2.81 (1.16-6.77) |
| ARDS      | 16      | 8 (0.50)  | 7 (0.44)  | 1 (0.06) | 0.020 | 0.013 | 0.029 | 0.016 |
| No ARDS   | 54      | 10 (0.19) | 34 (0.63) | 10 (0.19) | 4.40 (1.33-14.56) | 5.44 (1.43-20.77) | 2.56 (1.08-6.02) | 3.41 (1.26-9.22) |
| MV        | 24      | 11 (0.46) | 12 (0.50) | 1 (0.04) | 0.005 | 0.004 | 0.007 | 0.003 |
| No MV     | 46      | 7 (0.15)  | 29 (0.63) | 10 (0.22) | 4.72 (1.51-14.69) | 7.03 (1.83-26.95) | 2.78 (1.31-5.83) | 3.73 (1.55-9.00) |
| 1A1 (CGA) | 1A1/1A1 | 1A1/rest | rest/rest | 1A1/1A1 + 1A1/rest vs rest/rest | 1A1 vs rest |
|ARF       | 48      | 0 (0.00)  | 11 (0.23) | 37 (0.77) | 0.009 | 0.0017 | 0.004 | 0.005 |
|No ARF    | 22      | 2 (0.09)  | 10 (0.46) | 10 (0.46) | 0.25 (0.08-0.73) | 0.08 (0.02-0.39) | 0.23 (0.11-0.68) | 0.21 (0.07-0.62) |
| ARDS     | 16      | 0 (0.00)  | 2 (0.12)  | 14 (0.88) | - | 0.040 | - | 0.021 |
| No ARDS  | 54      | 2 (0.04)  | 19 (0.35) | 33 (0.61) | - | 0.16 (0.03-0.92) | - | 0.12 (0.02-0.73) |
| MV       | 24      | 0 (0.00)  | 4 (0.17)  | 20 (0.83) | 0.037 | 0.019 | 0.034 | 0.015 |
| No MV    | 46      | 2 (0.04)  | 17 (0.37) | 27 (0.59) | 0.28 (0.08-0.97) | 0.08 (0.01-0.66) | 0.31 (0.1-0.96) | 0.16 (0.04-0.70) |
| ICU      | 30      | 0 (0.00)  | 4 (0.13)  | 26 (0.87) | 0.003 | 0.015 | 0.003 | 0.029 |
| No ICU   | 40      | 2 (0.05)  | 17 (0.43) | 21 (0.53) | 0.17 (0.05-0.58) | 0.25 (0.08-0.76) | 0.20 (0.07-0.62) | 0.32 (0.11-0.89) |

Only those comparisons with P < 0.05 and significant odds ratio were included. OR (95% CI), odds ratio (95% confidence interval); ARF, acute respiratory failure; ARDS, acute respiratory distress syndrome; MV, mechanical ventilation; ICU, intensive care unit.

*P value for the bivariate comparison calculated with the χ² test.  + + value for the multivariate analysis calculated with binary logistic regression, including the variables age, gender, risk factors, secondary bacterial pneumonia, and bacteremia. P value by Fisher Exact test. *P value for the multivariate analysis calculated with binary logistic regression, including the variables age, gender, risk factors, and secondary bacterial pneumonia (bacteremia variable was excluded because it shows a co-linear relation with the variable mechanical ventilation). Patients who required ICU or did not require (No ICU) ICU admission, P value for the multivariate analysis calculated with conditional logistic regression stratified by hospital of origin, including the covariables age, gender, risk factors, secondary bacterial pneumonia, and bacteremia. Conventional haplotypes at SFTPA2 were identified on the basis of combinations of the polymorphism rs1059046 (T9N), rs17886395 (A91P), and rs1965708 (Q223K). Rest refers to the other haplotypes for each comparison.
84%, and more than 88% for the need for MV. In the haplotype analysis, statistical power was 88.56% for the effect of the haplotype 1A1 on ICU admission (90.62% for the effect in a dominant model). No correction for multiple comparisons was performed in these comparisons, but, as expected by in vivo and in vitro studies among the human collectins, only SP-A would be expected to influence H1N1pdm infectivity, and significant associations were repeatedly observed with several clinical phenotypes. Furthermore, the observed associations were found to be independent of age, gender, risk factors predisposing to severe H1N1 infection, and development

Figure 3 Kaplan-Meier estimation of days until ICU admission in hospitalized H1N1pdm-infected patients according to SFTPA2 variants. Only those comparisons with P < 0.05 were included. Solid curves represent the most frequent variant under study, and the dotted curves, the rest of variants. Significance levels calculated by means of log-rank test and Cox regression stratified by hospital of origin and adjusted for the variables age, gender, risk factors, secondary bacterial pneumonia and bacteremia are shown at the bottom of the figure. HR (95% CI), hazard ratio (95% confidence interval).
of either secondary bacterial pneumonia or bacteremia. Variants of \textit{SFTPA2} were clearly associated with functional parameters of gas exchange, underscoring their role in the severity of H1N1pdm-induced lung injury.

Third, criteria for ICU admission may vary between different centers. To avoid these differences, multivariate and Cox regression analysis to evaluate the need for ICU admission were stratified by hospital of origin.

\textbf{Conclusion}

Our study suggests that variants at \textit{SFTPA2} influence the severity of H1N1pdm infection in hospitalized patients. The potential of collectins as therapeutic agents for the treatment of IAV-mediated disease is now being explored \cite{7,30}. In \textit{Sftpa}−/− and \textit{Sftpd}−/− mice, intratracheally administered SP-A or SP-D can restore microbial clearance and inflammation \cite{5}, and exogenous surfactant preparations containing the hydrophobic SP-B and -C are widely used for replacement therapies in pRDS. The data from our study, together with a better knowledge of the functional significance of the genetic variability at \textit{SFTPA2} on IAV-associated disease, could pave the way for a potential treatment with haplotype-specific (1A\textsuperscript{1}) SP-A2 for patients with the most severe forms of the disease in future IAV pandemics.

\textbf{Key messages}

- Genetic variation in the \textit{SFTPA2} gene influences the outcome of patients infected with the 2009 pandemic H1N1 influenza A virus.
- Data from our study may help to identify patients at higher risk of severe pandemic (nonglycosylated) IAV infection, who may eventually benefit from more-personalized and targeted therapies.

\textbf{Abbreviations}

ARDS: Acute respiratory distress syndrome; ARF: acute respiratory failure; BMI: body mass index; CEIC: Comité Ético de Investigación Clínica (Clinical Research Ethics Committee); CRD: carbohydrate recognition domain; FiO\textsubscript{2}: fraction of inspired oxygen; H1N1pdm: virus influenza A H1N1 pandemic; HIV: human immunodeficiency virus; HR: hazard ratio; IAV: influenza A virus; ICU: intensive care unit; MBL: mannose-binding lectin; MODS: multigang dysfunction syndrome; MV: mechanical ventilation; NF-\kappa B: nuclear factor kappa B; OR: odds ratio; PaO\textsubscript{2}: partial pressure of oxygen;
pRDS: respiratory distress syndrome in prematurely born infant; PVP: primary viral pneumonia; SaO2: arterial oxygen saturation; SIRPα: signal inhibitory regulatory protein α; SNP: single-nucleotide polymorphism; SP: surfactant protein.

Competing interests
The author(s) declare that they have no competing interests.

Authors’ contributions
EHR: acquisition, analysis, and interpretation of molecular genetic data, and statistical analysis and collaboration in the writing of the manuscript. MLR: acquisition, analysis, and interpretation of clinical data and critical review of the manuscript. JPH: acquisition, analysis, and interpretation of clinical data and critical review of the manuscript. JPP: acquisition, analysis, and interpretation of clinical data and critical review of the manuscript. JB: acquisition, analysis, and interpretation of clinical data and critical review of the manuscript. MCPG: acquisition, analysis, and interpretation of H1N1pdm infection data and critical review of the manuscript. MB: molecular genetic data acquisition and critical review of the manuscript. VMB: molecular genetic data acquisition and critical review of the manuscript. WMB: acquisition, analysis, and interpretation of H1N1pdm infection data and critical review of the manuscript. MM: acquisition, analysis, and interpretation of clinical data and critical review of the manuscript. JM: acquisition, analysis, and interpretation of clinical data and critical review of the manuscript. LS: acquisition, analysis, and interpretation of clinical data and critical review of the manuscript. CV: acquisition, analysis, and interpretation of H1N1pdm infection data and critical review of the manuscript. OR: acquisition, analysis, and interpretation of clinical data and critical review of the manuscript. MB: acquisition, analysis, and interpretation of clinical data and critical review of the manuscript. JA: acquisition, analysis, and interpretation of clinical data and critical review of the manuscript. ELC: acquisition, analysis, and interpretation of clinical data and critical review of the manuscript. JSV: acquisition, analysis, and interpretation of clinical data and collaboration in the writing of the manuscript. FRC: financial support, acquisition, analysis, and interpretation of clinical data and collaboration in the writing of the manuscript. CRG: coordination, conception and design, financial support, and manuscript writing. All authors read and approved the final version of the manuscript.

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