To the Editor,
SARS-CoV-2 infection may induce a broad spectrum of consequences ranging from asymptomatic infection to fatal pneumonia.1 The most severe complication is the acute respiratory distress syndrome (ARDS) which is often fatal. The so-called "IL-6 cytokine storm" and a disseminated intravascular cascade in the lung characterize most severe cases of COVID-19. The coincidence of these events with the rise of the adaptive immune response suggests that the response itself may play a role. In effect, COVID-19 patients with agammaglobulinemia recovered without lung complications.2 Atopic status is the genetic predisposition to produce a Type 2 immune response to environmental antigens. Such response relies on some key cytokines, including interleukin (IL) 4, IL-5, IL-9, and IL-13.3 In infection, the Th2 response counteracts the microbicidal Th1 response, which could limit the tissue damage induced by Th1-mediated inflammation4 on one hand, but also cause a less efficient anti-virus response, as shown in a study on experimental Coronavirus 229E infection in healthy volunteers, where atopy appeared to be associated with a more severe rhinitis score.5 Further, atopic subjects show a reduced expression of ACE2, the SARS-CoV-2 receptor, which could be associated with reduced susceptibility to the virus.6 We therefore hypothesized that atopic subjects infected by SARS-CoV-2 might have a milder clinical course than nonatopic subjects, and tested this hypothesis in a large cohort of hospitalized COVID-19 patients.

We performed a retrospective study on patients with SARS-CoV-2-induced pneumonia, as confirmed by the detection of viral nucleic acid in nasal and/or pharyngeal clinical specimens, hospitalized in several Italian hospitals. Doctors recorded clinical data (age, sex, smoking habits, diabetes, hypertension, coronary heart disease, and thrombosis) along with respiratory allergy and graded the severity of the respiratory disease at the end of the hospital stay as mild, severe, or very severe based on no need for respiratory assistance, need for noninvasive respiratory assistance or need for invasive respiratory assistance or death, respectively. Patients were considered “atopic” in the presence of an unequivocal history of respiratory allergy to airborne allergens confirmed by positive skin prick tests performed by a specialist allergy center and/or by elevated specific IgE. All allergy investigations had been performed before hospitalization. Patients’ data were anonymized, and the internal review board of the promoting center approved the study. The association between severity of COVID-19 and both the atopy status and the clinical co-factors recorded was studied in both univariate and multivariate analyses (details in the Appendix S1). Type I statistic error probability values<5% were considered significant.

REFERENCES
1. Paules CI, Marston HD, Fauci AS. Coronavirus infections—more than just the common cold. JAMA. 2020;323(8):707.
2. Kissler SM, Tedijanto C, Goldstein E, Grad YH, Lipsitch M. Projecting the transmission dynamics of SARS-CoV-2 through the postpandemic period. Science. 2020;368:860–868.
3. Azkur AK, Akdis M, Azkur D, et al. Immune response to SARS-CoV-2 and mechanisms of immunopathological changes in COVID-19. Allergy. 2020;75(7):1564-1581.
4. Health SMo. Distribución geográfica de los casos totales y los casos de las últimas 24h (obtenidos a partir de la declaraciónagregada de casos del Ministerio de Sanidad). 2020. https://cneovid.isciii.es/covid19/
5. Li Z, Yi Y, Luo X, et al. Development and clinical application of a rapid IgM-IgG combined antibody test for SARS-CoV-2 infection diagnosis. Infect Ecol Epidemiol. 2020;10(1):175438.
6. Kramer F, Simon V. Serology assays to manage COVID-19. Science. 2020;368(6495):1060-1061.

SUPPORTING INFORMATION
Additional supporting information may be found online in the Supporting Information section.

Atopic status protects from severe complications of COVID-19

Domingo Barber, Departamento de Ciencias Médicas Básicas, Facultad de Medicina, Instituto de Medicina Molecular Aplicada (IMMA), Campus Montepríncipe, Universidad San Pablo-CEU, 28925 Alcorcón, Madrid, Spain.

Correspondence
Domingo Barber, Department of Basic Medical Sciences, Faculty of Medicine, Institute of Applied Molecular Medicine, Universidad San Pablo CEU, CEU Universities, Madrid, Spain.

Supporting Information
Additional supporting information may be found online in the Supporting Information section.

DOI: 10.1111/all.14551

1 Department of Basic Medical Sciences, Faculty of Medicine, Institute of Applied Molecular Medicine, Universidad San Pablo CEU, CEU Universities, Madrid, Spain
2 Microbiology Section, Departamento Ciencias Farmacéuticas y de la Salud, Facultad de Farmacia, Universidad San Pablo CEU, CEU Universities, Boadilla del Monte, Spain
3 HM Hospitales, Instituto de Investigación Sanitaria HM Hospitales, Hospital Universitario HM Montepríncipe, Madrid, Spain
4 Fundación de Investigación HM Hospitales, CEU Universities, Universidad San Pablo-CEU, Madrid, Spain

ORCID
María M. Escribese https://orcid.org/0000-0001-5057-5150
Domingo Barber https://orcid.org/0000-0002-5488-5700

SUPPORTING INFORMATION
Additional supporting information may be found online in the Supporting Information section.

DOI: 10.1111/all.14551
Five hundred and thirty one adult individuals (214 females, 40%) aged 68 ± 14 years (range 25-100 years) were studied. Their disease was classified as mild in 191 cases (36%) and severe or very severe in 340 cases (64%). The clinical features are summarized in Table 1. Male gender was significantly associated with severe disease (P = .001), and men were more likely to be admitted to intensive care units (P = .01). Smokers had a significantly higher prevalence of coronary heart disease (P = .023), and intensive care admission (P = .032) than nonsmokers. Male smokers showed a higher prevalence of coronary heart disease (39.7% vs 21.6%, P = .002) and thrombosis (P = .027) than female smokers (Table 1).

The multiple logistic regression analysis including age, sex, smoking, and all the comorbidities shows a significant association between nonatopic status and a significantly higher risk of having severe COVID-19-related pneumonia.

Fifty-seven participants (10.7%) were atopic. 32/57 (56.1%) were females (mean age 62 ± 14, range 28-93 years). Atopic subjects were younger (69 ± 14 years, P < .050) and, notably, showed a much lower occurrence of severe or very severe COVID-19 pneumonia (33.3% vs 67.7%, P < .0001). The protective effect of atopic status against severe lung disease was evident throughout all the age subsets (P = .001; Figure 1). The multiple logistic regression analysis (details in Appendix S1) confirmed a significant association between atopic status and milder COVID-19; nonatopic patients had a significantly higher risk of having severe COVID-19 (OR\textsubscript{adj} 3.0, 95% CI 1.6-5.7, P = .001) (Table 1). The unsupervised two-way hierarchical clustering analysis identified a clear cluster including mild COVID-19-related pneumonia and atopy status (P < .0001; OR\textsubscript{a} = 4.523, 95% CI = 2.221-9.221) (Figure S1).

In severe SARS-CoV-2 infection hyper-expressed cytokines include IFN-gamma, TNF-alpha, and IL-6, which cause fever, fatigue, flu-like symptoms, vascular leakage due to endothelial dysfunction, cardiomyopathy, hypotension, lung injury, activation of the coagulation cascade, and diffuse intravascular coagulation. The cytokine storm, which does not occur in patients with uncomplicated SARS-CoV-2 infection, leads to the rapid proliferation and activation of T cells, macrophages, and natural killer cells that eventually secrete >150 inflammatory cytokines, chemokines, and chemical mediators. Atopic patients are genetically predisposed to mount Th2 type immune-mediated responses; these responses do not imply the expression of the main cytokines involved in the ARDS. We hypothesized that allergic patients might be both less prone to SARS-CoV-2 infection and/or might have a less severe SARS-CoV-2 infection than nonatopic individuals. In the impossibility to carry out a field epidemiological study, we tried to address the second hypothesis focusing on more than 500 patients with confirmed COVID-19 infection.

### Table 1

Demographic and clinical features of the study population

| Variable | Level | Covid-19 | Severe | Mild | ORc | 95% CI | P-value | ORadj | 95% CI | P-value |
|----------|-------|----------|--------|------|-----|--------|---------|--------|--------|---------|
| Overall  |       | 531     | 340    | 191  |     |        |         |        |        |         |
| Atopy    | Yes   | 57      | 19     | 38   | 4.2 | 2.3-7.6 | <.001   | 3.0    | 1.6-5.7 | <.001   |
|          | No    | 474     | 321    | 153  |     |        |         |        |        |         |
| Sex      | Female | 214     | 115    | 99   | 2.1 | 1.5-3.0 | <.001   | 1.9    | 1.3-2.9 | <.002   |
|          | Male   | 317     | 225    | 92   |     |        |         |        |        |         |
| Age group (years) |       |        |        |      |     |        |         |        |        |         |
| <60     | No    | 140     | 88     | 52   |     |        |         |        |        |         |
|         | Yes   | 188     | 121    | 67   | 1.1 | 0.7-1.7 | .168    | 1.1    | 0.6-1.8 | .784    |
| 60-74   | No    | 203     | 131    | 72   | 1.2 | 0.8-2.0 | .444    | 1.2    | 0.7-2.0 | .530    |
|         | Yes   | 0       | 0      | 0    |     |        |         |        |        |         |
| 75+     | No    | 361     | 221    | 140  |     |        |         |        |        |         |
|         | Yes   | 105     | 56     | 49   | .724| 0.4-1.1 | .174    | 0.57   | 0.4-10.9 | .018    |
| Smoking | No    | 441     | 82     | 163  |     |        |         |        |        |         |
|          | Yes   | 90      | 18     | 28   | 0.8 | 0.5-1.2 | .292    |        |        |         |
| Diabetes | No    | 232     | 157    | 75   |     |        |         |        |        |         |
|          | Yes   | 299     | 183    | 116  | 0.8 | 0.5-1.1 | .123    |        |        |         |
| Hypertension | No | 395     | 253    | 142  |     |        |         |        |        |         |
|          | Yes   | 156     | 87     | 69   | 1.0 | 0.7-1.5 | .987    |        |        |         |
| Coronaropathy | No | 470     | 296    | 174  |     |        |         |        |        |         |
|          | Yes   | 61      | 44      | 17   | 1.5 | 0.8-2.7 | .161    |        |        |         |
| Thrombosis | No | 428     | 238    | 190  |     |        |         |        |        |         |
|          | Yes   | 93      | 92     | 1    | 73.4 | 10.1-531.8 | <.001  |        |        |         |

Note: The multiple logistic regression analysis including age, sex, smoking, and all the comorbidities shows a significant association between nonatopic status and a significantly higher risk of having severe COVID-19-related pneumonia.
Letters to the Editor

severe enough to warrant hospital admission. Thus, our study population lacks both asymptomatic and symptomatic SARS-CoV-2 infected patients with very mild disease, as these patients had to spend their quarantine period at home. Despite these obvious limitations, our study demonstrates that among hospitalized patients with severe COVID-19, atopic patients have less severe disease. Interestingly, the prevalence of atopic patients in our population (10.7%) appeared lower than that in the age-matched general population, although recent epidemiological data on allergy in Italy are missing. On the other hand, we cannot exclude that also other of our patients were atopic, although undiagnosed, as they had not undergone allergy investigations before. Notably, the "protective" effect of atopic status did not depend on the age or sex of patients nor the presence of other co-factors, such as cigarette smoke, coronary heart disease, diabetes, thrombosis, or hypertension. We are not in the position to ascertain whether the immunological scenario that we have hypothesized and that prompted us to perform this study is correct, as this would require an immune-histochemical analysis of patients' sputum. Nonetheless, our clinical findings make our initial hypothesis believable suggesting that the genetic predisposition to a Th2-oriented immune response might help to avoid the cytokine storm. Altogether, our findings indirectly confirm the observed low prevalence of asthmatics among patients admitted due to COVID-19 pneumonia. Of course, we cannot exclude that other factors such as treatment with inhaled corticosteroids might have played a role in our observation of a milder disease among atopic patients, particularly in the light of the recent finding of effective dexamethasone protection against severe pneumonia. However, all our patients, both atopic and nonatopic underwent systemic corticosteroid treatment during their hospital stay, and most atopic patients were not asthmatic and hence did not use inhaled corticosteroids before their admission.

In conclusion, atopic status seems to protect against the most severe, often fatal consequences of SARS-CoV-2 infection. Such finding may be of help for future studies investigating how to limit the clinical consequences of this infection.

Keywords

allergy, COVID-19, immune response, SARS-CoV-2

Conflict of Interest

No author has conflicts of interest to declare.

Enrico Scala
Damiano Abeni
Alberto Tedeschi
Giuseppina Manzotti
Baoran Yang
Paolo Borrelli
Alessandro Marra
Mauro Giani
Antonio Spadari
Francesca Saltalamacchia
Riccardo Asero

1Istituto Dermopatico dell’Immacolata, IRCCS FLMM, Roma, Italia
2UO Medicina Generale, Ospedale Bolognini ASST Bergamo Est, Seriate, Italia
3Casa di Cura Palazzolo, Bergamo, Italia
4ASST Carlo Poma, Mantova, Italy
5SSD Dermatologia e Allergologia, Ospedale Beauregard, Aosta, Italia
6UOC Medicina, ASST Rhodense – P.O. Rho (Milano), Rho, Italia
7Ambulatorio di Allergologia, Clinica San Carlo, Paderno Dugnano, Italia
Severe asthma in adults does not significantly affect the outcome of COVID-19 disease: Results from the Italian Severe Asthma Registry

To the Editor,

Severe asthma is a chronic disease affecting around 3%-8% of adult asthma population in Europe, with the refractory form estimated to occur in 0.1% of the general population.1,2 Severe asthma is characterized by increased use of healthcare resources (ie, emergency room/hospital admissions, access to intensive care units (ICU), and use of biologics) due to exacerbations compared with the less severe form. In the current SARS-CoV-2 pandemic, there is an ongoing debate on the role of asthma and use of immunomodulating drugs, like corticosteroids and biologics, on COVID-19 outcomes. According to available data on COVID-19 hospitalizations, asthma seems to play little role on the clinical severity or access to health resources, unlike other chronic conditions such as hypertension, obesity, and chronic obstructive pulmonary disease.3 However, to date, no information is available on the burden of severe asthma on COVID-19 severity and hospitalization rates.

A questionnaire was submitted to clinicians of the Italian Registry of Severe Asthma (IRSA) network,4 assessing the prevalence and clinical characteristics of patients with severe asthma who contracted COVID-19 during the outbreak in Italy (February 24, 2020 to May 18, 2020), and 41 out of 78 centers distributed evenly among different Italian regions participated to the survey, covering 65.6% of the total subjects of the IRSA registry (Figure 1).

Among the 558 subjects surveyed, 7 subjects contracted COVID-19 (1.25% of the national sample), 2 showing suggestive symptoms, 4 confirmed by positive reverse transcriptase-polymerase chain reaction and 1 with positive serology testing, with an average age of 54.5 years: 5 isolated at home/received home care (71.5%), while 2 subjects were admitted to the hospital (28.5%), none required access to ICU and no deaths were reported. All COVID-19 subjects with severe asthma came from 2 regions of Northern Italy (6 Lombardy, 1 Emilia-Romagna, 3.7% of the regional population), all showing one or more comorbidities and were treated with medium/high-dose inhaled corticosteroids plus long-acting beta-2 agonists (ICS-LABA) and biologics (Table 1).

We then compared our results with data provided by the Italian Department for Civil Protection in the same time period from the affected geographic areas,5 and we observed that the frequency of COVID-19 among subjects referred to IRSA centers strongly correlated with the prevalence of SARS-CoV-2 infection in the corresponding province (r = .798, P = .025, Spearman rho test).