Bibliometric Analysis of International Scientific Production on Pharmacologic Treatments for SARS-CoV-2/COVID-19 During 2020

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Background: COVID-19 is causing a grave global health and economic crisis and the fight against the pandemic has led to unprecedented scientific activity. Bibliometrics could be a useful tool for guiding future research lines and promoting international collaboration for an effective treatment. For this purpose, we have conducted a bibliometric analysis of scientific publications on drugs and therapies used to treat COVID-19 during 2020.

Methods: Data source: Web of Science. We gathered data on scientific production relating to drugs used to treat COVID-19. We calculated impact factors and analyzed production by institution, country, and journal, visualizing our results in bibliometric networks.

Results: In 1 year, production relating to COVID-19 exceeded 100,000 publications, with over 6,500 on Drugs and COVID-19. Research into hydroxychloroquine and chloroquine, remdesivir, lopinavir and ritonavir, tocilizumab and convalescent plasma is particularly noteworthy. Mean citations/study range from 11.9 to 15.4. Producer institutions fall into three groups: one in the US and centered on Harvard Medical School; another in Europe led by INSERS; and another in China led by Huazhong University of Science and Technology. Production by journal is widespread but the Journal of Medical Virology, International Journal of Antimicrobial Agents, and American Journal of Transplantation are noteworthy.

Conclusions: The volume of research that is currently under way is comparable to the magnitude of the pandemic itself. Such a high volume of studies is infrequent and the impact they have achieved has no known precedent. The producing countries are those with highest incidence of the pandemic and greatest scientific potential; moreover, inter-agency and international collaboration has reached extraordinarily high levels.

Keywords: COVID-19, SARS-CoV-2, scientific production, bibliometric analysis, pharmacologic treatments, bibliometric network, visualization
INTRODUCTION

Coronaviruses are single-stranded RNA genome viruses capable of infecting both humans and animals causing respiratory, gastrointestinal, and neurological illnesses (1, 2). They are characterized by being surrounded by a shell of transmembrane glycoproteins (S proteins), giving them a characteristic morphology in the shape of a crown (3). The virus uses these proteins which anchor themselves to receptors on the host cells they infect.

In the past 20 years, coronaviruses have caused three significant health emergencies, including the current one. In 2002, SARS-CoV (4) was discovered in Guandong, China; it was named thus as it caused a Severe Acute Respiratory Syndrome in patients. It was transmitted from nasal and oral fluids and by physical contact with contaminated surfaces (5). It infected some 8,000 individuals with a mortality rate of 9.5%. In 2012, a new coronavirus with SARS-like symptoms appeared in the Middle East and was named MERS-CoV (Middle East respiratory syndrome). This virus is still in circulation and according to the World Health Organization (WHO) has caused 862 deaths with 35% mortality (5). Its transmission rate is lower than that of SARS-CoV with an R0 close to 1. Suddenly, in December 2019, a group of patients in Wuhan, China, was diagnosed with pneumonia of unknown origin and a new species of coronavirus, called SARS-CoV-2 due to its similarity to SARS-CoV, was identified. The disease that causes SARS-CoV-2 was named “Coronavirus Disease of 2019” (COVID-19) by the WHO. This new coronavirus has infected an infinitely larger number than its predecessors, with an R0 of between 2 and 3.5 (5, 6). Mortality is around 2% (7), and at the time of writing (October 2021) deaths are approaching 5 million worldwide, with more than 240 million diagnosed cases (8). The WHO declared COVID-19 a global pandemic on 11 March 2020.

This pandemic is causing a grave health crisis with serious consequences for the world economy due to the rigorous confinement measures imposed. The current situation has led to substantial investment in research funding that has, in turn, triggered the hitherto unprecedented volume of production of studies on SARS-CoV-2/COVID-19. Major publishers and biomedical journals are sharing their content in open access to facilitate the visibility of research as a basis for the generation of new knowledge and the rapid search for solutions (9). A good example is the permanently up-to-date collection of the International Journal of Epidemiology > COVID-19, which hosts possibly the best series of papers on the incidence and epidemiological characteristics of COVID-19 in different regions and countries around the world. The massive amount of scientific information circulating has led to a rapid response in studies on scientific production related to COVID-19 from both the general (10–18) and socio-economic perspectives (19, 20). Some of these analyse COVID-19 as an unprecedented informational phenomenon and use a wide range of media as their source of data; bibliometric studies, however, specifically use the Web of Science (WoS), Scopus and Medline databases, which are considered more suitable for this type of analysis. Bibliometrics may provide exhaustive evaluations of the most active journals, countries, institutions, or authors in a given research field. For this reason, bibliometric analyses are very important for guiding future research trends and promoting international collaboration between institutions and countries.

Alongside the scientific race against time to develop and administer the first vaccines during 2020, the search for therapeutic and pharmacological treatments for COVID-19 that aim to find palliative remedies for the disease became a priority. Typically, this research is based on repurposing pre-existing drugs, currently prescribed for other pathologies, and now undergoing clinical trials to determine their capacity to prevent the virus from binding to human cells by modifying S proteins or receptor proteins present in cells (3, 21). Other are intended to try to stop replication of the virus, alleviate its inflammatory effects or regulate the disproportionate immune process (cytokine storms) that it triggers. In this line of work, numerous reviews of the research generated have already been published, indicating the speed with which the research has been produced and the need to synthesize the scientific information available (22–27). Furthermore, repositories such as The COVID-19 Real-Time Learning Network, sponsored by the Infectious Diseases Society of America (28), are especially useful. For its part, the WHO and agencies such as the Food and Drug Administration (FDA) also update possible applications available for use in COVID-19 (29, 30). However, there is still no controlled scientific evidence for an completely effective treatment. Science has recently published a trial of plitidepsin (aplidin) that reports it has an antiviral potency 100 times greater than that of remdesivir (31). At 11 February 2021, another clinical trial has shown that tocilizumab is effective against COVID-19, reducing mortality by 4% in severe infections and, potentially, by 50% when administered with dexamethasone (32). The numerous earlier studies of tocilizumab administration in COVID-19 have been the subject of a meta-analysis (33). Right after COVID-19 was declared a global pandemic and during 2020, the most investigated drugs against COVID-19 included hydroxychloroquine, remdesivir, lopinavir, ritonavir, tocilizumab, azithromycin, among others. After that, some of the drugs listed above along with some new ones including favipiravir, thalidomide, ivermectin, and umifenovir have been considered as the most promising ones during the recent year 2021 (34). Although they are not the goal of the present bibliometrics analysis, it is important to note that vaccines are the most efficient preventive treatment against COVID-19. As a consequence of the rapid development and massive research funding, m-RNA vaccines have been successfully employed for the first time. These m-RNA-based vaccines, such as Pfizer BioNTech and Moderna, consist of messenger RNA (ribonucleic acid) molecules, which contain the genes necessary for viral proteins production, that trigger in an immune response by the host.

The aim of the present study is to analyse international scientific publications through bibliometric indicators on drugs and pharmacological treatments for use in treating COVID-19 during 2020. This work provides novel quantitative data that will be useful for promoting international collaboration between institutions and countries to search for possible clinical responses to the pandemic, as well as for the development of new research.
MATERIALS AND METHODS

Data Source
We have gathered our data from scientific production indexed in the WOS databases (35). This multi-disciplinary international source references the most prestigious scientific publications in the world and is an essential starting point for bibliometric studies providing indicators of production and scientific impact. WOS has been seen to match the current pace of publishing, quickly indexing the special COVID-19 sections that journals have created (Online articles, Articles in press, Early Access, Latest issue, etc.) thus enhancing their diffusion and visibility. We found no differences in coverage between WOS and the LitCovid-PUBMED repository. We searched WOS “All Databases,” which includes the biomedical coverage of MEDLINE, BIOSIS and SCIELO. We launched our searches between late December 2020 and January 2021. Given the level of ongoing scientific production, the final data provided in this study may vary, although any differences do not significantly affect our analysis of results.

Search Strategies and Data Treatment and Analysis
The search strategies used to recover the scientific production indexed in WOS on the subject of study, as well as the treatment and analysis of the data obtained, are described in Supplementary Material.

RESULTS AND DISCUSSION

Evolution of Scientific Production on Coronavirus
Prior to the arrival of SARS-CoV-2/COVID-19, SARS-CoV, and MERS-CoV caused significant worldwide health emergencies that had been studied by the scientific community. Data show that the corresponding scientific production was significant with 1,625 documents dated in 2003 or later (Figure 1)—months after SARS-COV had been detected. Peak production on this coronavirus amounted to 4,018 studies published between 2004 and 2006. Subsequently, production fell to 2,000 documents in 2007–2009; a process that continued until 2017, with 400–500 documents per year. MERS-CoV, identified in 2012, followed a similar evolution but with a smaller volume of studies, corresponding to its lower epidemic potential. Peak production occurred in 2016–2018, with almost 1,000 documents. However, the upsurge in COVID-19–related production between December 2019 and 2020 has confounded all conceivable predictions of statistical values. In 2020, production rose from almost 0 to over 100,000 documents recorded by WOS, and the number of documents continues to grow. In a single year, all-time production on all coronaviruses has grown seven-fold, reaching around 15,000–20,000 documents. This enormous differences in scientific production between SARS-CoV-2 and other coronaviruses (SARS-CoV and MERS-CoV) can be observed as well analyzing the data obtained in previous works (36, 37).

Drugs and SARS-CoV-2/COVID-19 Production and Impact
At the end of 2020, scientific production on Drugs and SARS-CoV-2/COVID-19 amounted to 6,533 documents (7% of all SARS-CoV-2/COVID-19 production). Logically, the co-occurrence of these two themes only appears in 2020, when their contents coincided as an object of investigation in the same studies (Table 1). The presence of earlier documents must be attributed to errors in record indexing. The typology of the studies is dominated by articles (60%), followed by the reviews (15%), case reports and clinical trials (10%), early access (10%) and letters (5%). Note that prior to 2020, research on drugs and their repurposing for other pathologies already existed, with substantial annual production rates (Table 1).

It is estimated that hundreds of drugs and therapeutic applications are currently being investigated in relation to COVID-19. The present study focuses on those that have received most attention in scientific publications. Table 2 lists the drugs that have led to the production of ≥150 studies, and shows indicators of production and impact measured in terms of citation. In terms of production, hydroxychloroquine— with about 2,000 studies—and chloroquine—with more than 1,000 studies in just 1 year, stand out. These are followed by the antivirals group formed by remdesivir, lopinavir, and ritonavir, with around 1,000 studies produced since February
Finally, it is important to note that several bibliometric analyses about COVID-19 in general terms and vaccines have been already published. However, the present manuscript offers, for the first time, important bibliometric indicators on drugs treatments against COVID-19.

Producer Institutions, Countries, and Journals

We will now focus on agents of production. Table 3 lists the institutions producing ≥35 studies distributed by drugs. The French INSERS (Institut National de la Santé et de la Recherche Médicale) center leads the list with 213 studies. This multi-center in Health Sciences is considered among the best in the world and ranks alongside the US National Institutes of Health. It stands out for having produced 66 studies on hydroxychloroquine, 41 on chloroquine, and others on remdesivir, lopinavir, and ritonavir. It is followed by the University of Milan, Italy, and Harvard Medical School, which also stand out for studies on hydroxychloroquine and antiviral drugs. With production levels of >100 studies we find Aix-Marseille University, Harvard University, Huazhong University of Science and Technology (Wuhan), the Chinese Academy of Medical Sciences (Beijing) and the Assistance Publique Hôpitaux, Paris, indicating that France and the US are increasingly important producers, as is the People’s Republic of China (PRC). The remaining institutions, with more disperse production, underscore the pre-eminence of
these countries although many Italian centers appear, as do some in Canada and Iran.

Some differences can be appreciated when analyzing bibliometrics of global scientific research on COVID-19 (38). In COVID-19 general research, Huazhong University of Science and Technology and Wuhan University, where the disease was discovered, are the most producers in terms of publications by far, with 300 and 170 (launched October, 2020) (38). As mentioned above, our results highlights INSERS, the University of Milan, and Harvard Medical School as the top 3 on research on pharmacological treatments against COVID-19 and lead Huazhong University of Science and Technology, and Wuhan University to the 6th and 16th position. This points out the different research specialties among different institutions.

Our visualization of the bibliometric network of institutions (co-occurrence ≥ 25) includes the first 68 institutions and depicts three clusters (Figure 2); one located in the US and Canada and centered around Harvard Medical School; another in Europe led by French and Italian universities; and a third in the PRC, with fewer but tightly grouped nodes, centered on Huazhong University of Science and Technology. The US has strong connections indicating tight inter-institutional collaboration; it is closer to the European group and interacts less with the PRC. The PRC cluster shows a marked closeness between actors, although connections are less intense than those observed in the US; three subgroups are evident: one centered on Huazhong University of Science and Technology and connected with Fudan University and Wuhan University; another around the Chinese Academy of Science; and a third around Capital Medical University (Beijing). The Chinese cluster’s links to the outside show some interaction with the European cluster through the Chinese Academy of Science and the National University of Singapore. The European cluster shows marked, tight interconnections, especially among the many Italian institutions that interact well with their French counterparts, University College London in the UK, and the National and Kapodistrian University of Athens, Greece. In Europe, we see two peculiarities: on the one hand, some English institutions are absent (University of Oxford and Imperial College London); on the other, INSERM does not appear (ranked 1st in Table 3). This is because the data in Figure 2 give preference to the affiliation of the first signatory, thus reflecting its level of prominence. In contrast, Table 3 shows any institution present among the signatories, and INSERM, although it finances and is involved in a very important part of the work, never appears as the principal institution.

If we look in detail at the collaboration patterns of other centers—such as MacMaster University (Ontario, Canada), Imperial College London, University of Oxford, University of Barcelona, and the University of São Paulo (Brazil)—we see that their positions within the network may be due to the fact that the strong internal interaction of large groups moves them to less well-defined positions. Finally, we would also wish to highlight the peripheral position of Iranian and Indian institutions and the absence of Russian and German institutions, which fail to reach the co-occurrence of collaboration threshold.

Our visualization of the bibliometric network by country (Figure 3) allows us to indicate other elements: among the poorly-represented countries we can see those with institutions
TABLE 3 | Number of studies produced by institution and drug (≥35).

| Institutions | HY | CH | RE | LO | RI | TO | CP | AZ | MA | AC | INF | Total |
|--------------|----|----|----|----|----|----|----|----|----|----|-----|-------|
| Inst Nat San et la Recher Medi. Inserm | 66 | 41 | 16 | 18 | 15 | 12 | 24 | 13 | 8  | 213 |
| University of Milan | 42 | 20 | 17 | 19 | 20 | 21 | 17 | 14 | 170 |
| Johns Hopkins Univ | 53 | 30 | 27 | 12 | 12 | 29 | 15 | 147 |
| China Acad Sci | 44 | 25 | 24 | 11 | 13 | 17 | 16 | 149 |
| Assistance Publish Hôpitaux Paris APHP | 30 | 21 | 18 | 20 | 16 | 20 | 8  | 133 |
| Columbia Univ | 42 | 19 | 13 | 12 | 19 | 10 | 11 | 96  |
| Univ Paris | 38 | 27 | 14 | 19 | 12 | 17 | 16 | 98  |
| Univ of Washington | 35 | 38 | 12 | 16 | 12 | 17 | 15 | 94  |
| Chinese Acad Sci | 26 | 19 | 17 | 12 | 17 | 91  |
| Mayo Clin | 31 | 16 | 17 | 11 | 15 | 90  |
| University of California System | 28 | 22 | 14 | 14 | 9  | 87  |
| Johns Hopkins Univ | 41 | 17 | 17 | 17 | 7  | 75  |
| Wuhan Univ | 16 | 21 | 12 | 12 | 12 | 74  |
| Mcmaster Univ (Ontario) | 24 | 16 | 13 | 13 | 10 | 69  |
| Fudan Univ | 26 | 14 | 13 | 13 | 18 | 58  |
| Stanford Univ | 27 | 15 | 16 | 15 | 58  |
| Brigham Womens’ Hospital | 33 | 22 | 14 | 12 | 55  |
| Cent South Univ Changsha, Hunan | 25 | 16 | 12 | 12 | 53  |
| New York University | 39 | 13 | 13 | 13 | 52  |
| Tehran Univ Med Sci | 17 | 12 | 13 | 10 | 52  |
| Univ Naples Federico II | 13 | 13 | 13 | 13 | 52  |
| Shahid Beheshti Univ Med Sci | 19 | 16 | 14 | 49  |
| Univ Hong Kong | 13 | 16 | 17 | 46  |
| Univ Campania Luigi Vanvitelli | 14 | 14 | 15 | 43  |
| Zhejiang Univ | 15 | 15 | 13 | 43  |
| Vanderbilt Univ Nashville, Tenn | 23 | 17 | 17 | 40  |
| Ins de Recherche Pour le Developpement IRD | 22 | 17 | 17 | 39  |
| Capital Med Univ Beijing | 22 | 12 | 12 | 35  |
| Sapienza Univ Rome | 21 | 14 | 35  |
| Total | 887 | 428 | 246 | 251 | 248 | 136 | 131 | 120 | 90 | 41 | 2,826 |

HY, Hydroxychloroquine; CH, Chloroquine; RE, Remdesivir; LO, Lopinavir; RI, Ritonavir; TO, Tocilizumab; CP, Convalescent Plasma; AZ, Azithromycin; MA, Monoclonal Antibodies; AC, Angiotensin-Converting Enzyme 2 Receptor; INF, Interferon.

that have produced > 100 works, which have research potential and have experienced considerable incidence of the pandemic; the positions of the UK, Germany, Canada, Australia and Japan are strengthened or appear for the first time; the positions of Iran and India are strengthened, with significant collaboration in Asia and the Arab world; Saudi Arabia appears; and Spain, Brazil, Belgium, and Turkey are more clearly visible. Bibliometric analysis of global research on COVID-19 highlighted United States, China, Italy United Kingdom, and India as the top countries in number of publications about general research on COVID-19 (38). The research on pharmacological treatments against COVID-19 follows a similar trend, with the same countries in the main positions (Figure 3).

In terms of journal production, the distribution is more disperse. Table 4 shows journals that have published ≥ 40 studies. The Journal of Medical Virology [IF (Impact Factor) 2019 = 2.021, Q4, Virology] with 174 papers stands out and is the only journal that has published on all the drugs under study. With more than 100 papers, we find the International Journal of Antimicrobial Agents (IF 2019 = 4.62, Q1, Microbiology); and the International Journal of Infectious Diseases (IF 2019 = 3.20, Q2, Infectious Diseases). The production of the American Journal of Transplantation (IF 2019 = 7.33, Q1, Transplantation) is striking, with 134 studies. The journal, which specializes in organ and tissue transplantation, has created a special section entitled COVID-19 in Transplantation Infectious Disease. Their interest in the issue may be due to the fact that transplanted
patients are permanently immunocompromised and, therefore, one of the main groups at risk from COVID-19. A similar explanation may justify the production of two journals that have published by far the highest number of papers on a single drug: *Annals of the Rheumatic Diseases* (Rheumatology, IF 2019 = 16.102; Position 2/32) with 63 hydroxychloroquine studies—the drug most frequently researched in the context of COVID-19 and traditionally administered in diseases such as rheumatoid arthritis. *Transfusion* (IF 2019 = 2.80; 40/76, Q3, Hematology) with 61 papers on convalescent plasma; the journal is sponsored by the American Association of Blood Banks and has a panel of experts that issues recommendations for the use of convalescent plasma in COVID-19.

Finally, given the clinical condition presented by patients with COVID-19, the presence of Medicine, General, and Internal journals at the top of the rankings is logical. Here we highlight the presence of the *New England Journal of Medicine* (IF 2019 = 74.69, Q1, first in its speciality, 1/65) with 100 papers; and the *Lancet* (IF 2019 = 60.39, Q1, 2/65), with 50 studies, but ranking in an intermediate position. The remaining journals appear in a long list included publications in the field of Pharmacology and Pharmacy, and dominated by those in Microbiology, Infectious Diseases, and Immunology.

**CONCLUSIONS**

The present study clearly demonstrates that global research on drugs and pharmacological treatments against COVID-19 has been massive and has reached unprecedented levels in terms of number of publications, citations, impact factor, and cooperation between countries and institutions. Our results showed that the therapy against COVID-19 has been focused on the efficiency of existing medical treatments for their application to this disease. The studies and citations they have received during a single year (2020) have reached unprecedented levels (14–20 MCS for azithromycin and ACE2, for example) in comparison with the most prestigious journals, i.e., *Nature* or *Science* years before (8.4 and 9.6 MCS during in 2018–2019, respectively). Similarly, h-index scores have reached equally unprecedented values higher than 50 in 2020, while the highest scores of articles in *Nature* or *Science* only reached 40. However, it should be noted that to a large extent, the aforementioned citation activity feeds back into itself with 14–32% of self-citation between studies, a figure higher than that typical in periods of normal scientific activity. Producer countries and institutions correlate with three variables: the origin of SARS-CoV-2; the countries most affected by the pandemic; and world leaders in...
TABLE 4 | Production in terms of number of studies by journal and drug (≥40).

| Journals                      | HY  | CH  | RE  | LO  | RI  | TO  | CP  | AZ  | MA  | AC  | INF | Total |
|-------------------------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-------|
| J Med Virol                   | 21  | 14  | 13  | 28  | 29  | 11  | 5   | 10  | 8   | 8   | 8   | 174   |
| Am J Transplantation          | 39  | 13  | 15  | 15  | 26  | 16  | 10  |     |     |     |     | 134   |
| Int J Antimicrob AG           | 37  | 29  | 10  | 8   | 8   | 5   | 21  | 5   | 5   |     |     | 128   |
| Int J Infect Dis              | 30  | 6   | 14  | 12  | 16  | 7   | 19  | 10  |     |     |     | 114   |
| New Engl J Med               | 29  | 32  | 21  | 19  |     |     |     |     |     |     |     | 101   |
| Cureus                        | 27  | 10  | 14  | 8   | 7   | 10  | 5   | 18  |     |     |     | 99    |
| J Biosol Struct Dynamics      | 20  | 14  | 20  | 14  |     |     |     |     |     |     |     | 98    |
| Ann Rheum Dis                | 63  | 7   |     | 18  | 7   |     |     |     |     |     |     | 95    |
| Trials                        | 30  | 12  | 6   | 9   | 5   | 6   | 9   | 7   |     |     |     | 92    |
| Am J Trop Med Hyg             | 29  | 13  | 6   | 13  | 15  |     |     |     |     |     | 13   | 89    |
| BMJ (Br Med J)               | 29  | 22  |     |     |     |     |     |     |     |     |     | 68    |
| Biorxiv                       | 6   | 17  |     |     |     |     |     |     |     |     |     | 72    |
| Medical Hypotheses            | 17  | 12  | 5   | 6   | 5   |     |     |     |     |     |     | 67    |
| Transfusion                   |     |     |     |     |     |     |     |     |     |     |     | 61    |
| Viruses                       |     |     |     |     |     |     |     |     |     |     |     | 60    |
| JAMA                          |     |     |     |     |     |     |     |     |     |     |     | 59    |
| Lancet                        |     |     |     |     |     |     |     |     |     |     |     | 50    |
| Front Pharmacol               |     |     |     |     |     |     |     |     |     |     |     | 49    |
| Clin Infect Dis               |     |     |     |     |     |     |     |     |     |     |     | 40    |
| Eur Rev Med Pharmacol Sci    |     |     |     |     |     |     |     |     |     |     |     | 44    |
| Front Immunol                 |     |     |     |     |     |     |     |     |     |     |     | 43    |
| J Antimicrob Chemother        |     |     |     |     |     |     |     |     |     |     |     | 43    |
| Eur J Pharmacol               |     |     |     |     |     |     |     |     |     |     |     | 41    |
| Nature                        |     |     |     |     |     |     |     |     |     |     |     | 41    |
| Pharmacol Res                |     |     |     |     |     |     |     |     |     |     |     | 41    |
| Ann Intern Med               |     |     |     |     |     |     |     |     |     |     |     | 40    |

HY, Hydroxychloroquine; CH, Chloroquine; RE, Remdesivir; LO, Lopinavir; RI, Ritonavir; TO, Tocilizumab; CP, Convalescent Plasma; AZ, Azithromycin; MA, Monoclonal antibodies; AC, Angiotensin-converting enzyme 2 Receptor; INF, Interferon.

research. The institutions are also those that conduct research into the most promising drugs, including the antimalarials hydroxychloroquine and chloroquine; antiviral drugs remdesivir, lopinavir and ritonavir; the antibiotic azithromycin; tocilizumab, and convalescent plasma. Collaboration between institutions and between countries is significant, and transparency and the exchange of research results has certainly led to rapid progress in the fight against the disease and clinical treatments have been approved. Substantial inter-institutional research has taken place between centers in the same country and between countries in the same geographical area.

The results here provided can be very useful for the development of new and existing research lines on pharmacological treatments against COVID-19, as well as to promote international collaboration, which in turn would help to achieve an effective cure against this horrible disease.

AUTHOR CONTRIBUTIONS

MAR-F and RR-P: conceptualization, formal analysis, and investigation. EJ-C, CR-F, and RR-P: methodology. MAR-F, EJ-C, and RR-P: software and writing—original draft preparation. MAR-F, CR-F, and RR-P: writing—review and editing. MAR-F: visualization. RR-P: supervision and project administration. RR-P and EJ-C: funding acquisition. All authors contributed to the article and approved the submitted version.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fpubh.2021.778203/full#supplementary-material
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