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Review of the Emerging Evidence Demonstrating the Efficacy of Ivermectin in the Prophylaxis and Treatment of COVID-19

Pierre Kory, MD,^{1*} Gianfranco Umberto Meduri, MD,² Joseph Varon, MD,³
Jose Iglesias, DO,⁴ and Paul E. Marik, MD⁵

Background: After COVID-19 emerged on U.S shores, providers began reviewing the emerging basic science, translational, and clinical data to identify potentially effective treatment options. In addition, a multitude of both novel and repurposed therapeutic agents were used empirically and studied within clinical trials.

Areas of Uncertainty: The majority of trialed agents have failed to provide reproducible, definitive proof of efficacy in reducing the mortality of COVID-19 with the exception of corticosteroids in moderate to severe disease. Recently, evidence has emerged that the oral antiparasitic agent ivermectin exhibits numerous antiviral and anti-inflammatory mechanisms with trial results reporting significant outcome benefits. Given some have not passed peer review, several expert groups including Unitaid/World Health Organization have undertaken a systematic global effort to contact all active trial investigators to rapidly gather the data needed to grade and perform meta-analyses.

Data Sources: Data were sourced from published peer-reviewed studies, manuscripts posted to preprint servers, expert meta-analyses, and numerous epidemiological analyses of regions with ivermectin distribution campaigns.

Therapeutic Advances: A large majority of randomized and observational controlled trials of ivermectin are reporting repeated, large magnitude improvements in clinical outcomes. Numerous prophylaxis trials demonstrate that regular ivermectin use leads to large reductions in transmission. Multiple, large “natural experiments” occurred in regions that initiated “ivermectin distribution” campaigns followed by tight, reproducible, temporally associated decreases in case counts and case fatality rates compared with nearby regions without such campaigns.

Conclusions: Meta-analyses based on 18 randomized controlled treatment trials of ivermectin in COVID-19 have found large, statistically significant reductions in mortality, time to clinical recovery, and time to viral clearance. Furthermore, results from numerous controlled prophylaxis trials report significantly

¹Front-Line Covid-19 Critical Care Alliance, Madison, WI; ²Memphis VA Medical Center—University of Tennessee Health Science Center, Pulmonary, Critical Care, and Research Services, Memphis, TN; ³University of Texas Health Science Center, Critical Care Service, Houston, TX; ⁴Department of Medicine, Hackensack School of Medicine, Seton Hall, NJ; and ⁵Eastern Virginia Medical School, Division of Pulmonary and Critical Care, Norfolk, VA.

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P. Kory and G. U. Meduri have contributed equally to this work.

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Off-Label Use: This manuscript includes discussion of off-label use in COVID-19 of the FDA-approved medication ivermectin.

*Address for correspondence: Pierre Kory, MD, MPA, Front-Line Covid-19 Critical Care, 2002 L St NW, Suite 500, Washington, D.C. 20036. E-mail: pkory@flccc.net

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reduced risks of contracting COVID-19 with the regular use of ivermectin. Finally, the many examples of ivermectin distribution campaigns leading to rapid population-wide decreases in morbidity and mortality indicate that an oral agent effective in all phases of COVID-19 has been identified.

Keywords: ivermectin, COVID-19, infectious disease, pulmonary infection, respiratory failure

INTRODUCTION

In early 2020, on the onset of the spreading pandemic, many providers and institutions began to continuously review the rapidly emerging basic science, translational, and clinical data to identify potentially effective treatment options for COVID-19. Although there is now a small and increasing number of therapeutics showing some efficacy in important clinical outcomes, chief of which are corticosteroids in moderate to severe illness, the world continues to suffer from a worsening crisis with the potential of again overwhelming hospitals and intensive care units (ICU). As of February 21, 2020, the number of deaths attributed to COVID-19 in the United States reached 510,248 with more than 9.3 million active cases, the highest number to date. In addition, multiple European countries have imposed new rounds of restrictions and lockdowns.

Further compounding these alarming developments was a wave of recently published results from therapeutic randomized controlled trials conducted on medicines believed effective for COVID-19 that found a lack of impact on mortality in hospitalized patients with the use of remdesivir, hydroxychloroquine, lopinavir/ritonavir, interferon, convalescent plasma, and monoclonal antibody therapy.¹⁻⁴ One year into the pandemic, the only therapy considered “proven” as a life-saving treatment in COVID-19 is the use of corticosteroids in patients with moderate to severe illness.^{5,6} Similarly, most concerning is the fact that no agent has yet proven effective in outpatients to prevent disease progression to prevent hospitalization.

More recently, trial results of ivermectin, a widely used antiparasitic medicine with known antiviral and anti-inflammatory properties, have been showing benefits in multiple important clinical and virologic outcomes, including mortality. Although growing numbers of the studies supporting this conclusion have passed through peer review, approximately half of the remaining trials data are from manuscripts uploaded to medical preprint servers, a now standard practice for both rapid dissemination and adoption of new therapeutics throughout the pandemic. Following is a comprehensive review of the available efficacy data as of December 12, 2020, taken from in vitro, animal, clinical, and real-world studies all showing the above impacts of ivermectin in COVID-19.

History of ivermectin

In 1975, Professor Satoshi Omura at the Kitasato institute in Japan isolated an unusual *Streptomyces* bacterium from the soil near a golf course along the southeast coast of Honshu, Japan. Omura, along with William Campbell, found that the bacterial culture could cure mice infected with the roundworm *Heligmosomoides polygyrus*. Campbell isolated the active compounds from the bacterial culture, naming them “avermectins” and the bacterium *S. avermitilis* for the compounds’ ability to clear mice of worms.⁷ Despite decades of searching around the world, the Japanese microorganism remains the only source of avermectin ever found. Ivermectin, a derivative of avermectin, then proved revolutionary. Originally introduced as a veterinary drug, it soon made historic impacts in human health, improving the nutrition, general health, and well-being of billions of people worldwide ever since it was first used to treat onchocerciasis (river blindness) in humans in 1988. It proved ideal in many ways, given that it was highly effective, broad-spectrum, safe, well tolerated, and could be easily administered.⁷ Although it was used to treat a variety of internal nematode infections, it was most known as the essential mainstay of 2 global disease elimination campaigns that has nearly eliminated the world of two of its most disfiguring and devastating diseases. The unprecedented partnership between Merck & Co. Inc. and the Kitasato Institute combined with the aid of international health care organizations has been recognized by many experts as one of the greatest medical accomplishments of the 20th century. One example was the decision by Merck & Co to donate ivermectin doses to support the Mectizan Donation Program that then provided more than 570 million treatments in its first 20 years alone.⁸ Ivermectin’s impacts in controlling onchocerciasis and lymphatic filariasis, diseases which blighted the lives of billions of the poor and disadvantaged throughout the tropics, is why its discoverers were awarded the Nobel Prize in Medicine in 2015 and the reason for its inclusion on the World Health Organization’s (WHO) “List of Essential Medicines.” Furthermore, it has also been used to successfully overcome several other human diseases and new uses for it are continually being found.⁷

Preclinical studies of Ivermectin's activity against SARS-CoV-2

Since 2012, a growing number of cellular studies have demonstrated that ivermectin has antiviral properties against an increasing number of RNA viruses, including influenza, *Zika*, HIV, *Dengue*, and most importantly, SARS-CoV-2.⁹⁻¹⁷ Insights into the mechanisms of action by which ivermectin both interferes with the entrance and replication of SARS-CoV-2 within human cells are mounting. Caly et al¹⁸ first reported that ivermectin significantly inhibits SARS-CoV-2 replication in a cell culture model, observing the near absence of all viral material 48 hours after exposure to ivermectin. However, some questioned whether this observation is generalizable clinically given the inability to achieve similar tissue concentrations used in their experimental model using standard or even massive doses of ivermectin.^{19,20} It should be noted that the concentrations required for an effect in cell culture models bear little resemblance to human physiology given the absence of an active immune system working synergistically with a therapeutic agent, such as ivermectin. Furthermore, prolonged durations of exposure to a drug likely would require a fraction of the dosing in short-term cell model exposure. Furthermore, multiple coexisting or alternate mechanisms of action likely explain the clinical effects observed, such as the competitive binding of ivermectin with the host receptor-binding region of SARS-CoV-2 spike protein, as proposed in 6 molecular modeling studies.²¹⁻²⁶ In 4 of the studies, ivermectin was identified as having the highest or among the highest of binding affinities to spike protein S1 binding domains of SARS-CoV-2 among hundreds of molecules collectively examined, with ivermectin not being the particular focus of study in 4 of these studies.²⁷ This is the same mechanism by which viral antibodies, in particular, those generated by the Pfizer and Moderna vaccines contain the SARS-CoV-2 virus. The high binding activity of ivermectin to the SARS-CoV-2 spike protein could limit binding to either the ACE-2 receptor or sialic acid receptors, respectively, either preventing cellular entry of the virus or preventing hemagglutination, a recently proposed pathologic mechanism in COVID-19.^{21,22,26-28} Ivermectin has also been shown to bind to or interfere with multiple essential structural and nonstructural proteins required by the virus to replicate.^{26,29} Finally, ivermectin also binds to the SARS-CoV-2 RNA-dependent RNA polymerase (RdRp), thereby inhibiting viral replication.³⁰

Arevalo et al investigated in a murine model infected with a type 2 family RNA coronavirus similar to SARS-CoV-2, (mouse hepatitis virus), the response to 500 µg/kg of ivermectin versus placebo.³¹ The study included 40 infected mice, with 20 treated with ivermectin, 20 with phosphate-buffered saline, and then 16 uninfected control

mice that were also given phosphate-buffered saline. At day 5, all the mice were killed to obtain tissues for examination and viral load assessment. The 20 nonivermectin-treated infected mice all showed severe hepatocellular necrosis surrounded by a severe lymphoplasmacytic inflammatory infiltration associated with a high hepatic viral load (52,158), whereas in the ivermectin-treated mice a much lower viral load was measured (23,192; $P < 0.05$), with only few livers in the ivermectin-treated mice showing histopathological damage such that the differences between the livers from the uninfected control mice were not statistically significant.

Dias De Melo et al³² recently posted the results of a study they did with golden hamsters that were intranasally inoculated with SARS-CoV-2 virus, and at the time of the infection, the animals also received a single subcutaneous injection of ivermectin at a dose of 0.4 mg/kg on day 1. Control animals received only the physiologic solution. They found the following among the ivermectin-treated hamsters: a dramatic reduction in anosmia (33.3% vs. 83.3%, $P = 0.03$), which was also sex dependent in that the male hamsters exhibited a reduction in clinical score while the treated female hamsters failed to show any sign of anosmia. They also found significant reductions in cytokine concentrations in the nasal turbinates and lungs of the treated animals, despite the lack of apparent differences in viral titers.

Despite these mounting insights into the existing and potential mechanisms of action of ivermectin both as a prophylactic and treatment agent, it must be emphasized that significant research gaps remain and that many further in vitro and animal studies should be undertaken to better define not only these mechanisms but also to further support ivermectin's role as a prophylactic agent, especially in the optimal dose and frequency required.

Preclinical studies of ivermectin's anti-inflammatory properties

Given that little viral replication occurs in the later phases of COVID-19, nor can virus be cultured, and only in a minority of autopsies can viral cytopathic changes be found,³³⁻³⁵ the most likely pathophysiologic mechanism is that identified by Li et al³⁶ where they showed that the nonviable RNA fragments of SARS-CoV-2 lead to a high mortality and morbidity in COVID-19 through the provocation of an overwhelming and injurious inflammatory response. Based on these insights and the clinical benefits of ivermectin in the late phase of disease to be reviewed below, it seems that the increasingly well-described in vitro properties of ivermectin as an inhibitor of inflammation are far more clinically potent than previously recognized. The growing list of studies demonstrating the anti-inflammatory properties of ivermectin include its

ability to inhibit cytokine production after lipopolysaccharide exposure, downregulate transcription of NF- κ B, and limit the production of both nitric oxide and prostaglandin E₂.^{37–39}

Exposure prophylaxis studies of ivermectin's ability to prevent transmission of COVID-19

Data are also now available showing large and statistically significant decreases in the transmission of COVID-19 among human subjects based on data from 3 randomized controlled trials (RCTs) and 5 observational controlled trials (OCTs) with 4 of the 8 (2 of them RCTs) published in peer-reviewed journals.^{40–46}

Elgazzar and colleagues⁴⁵ at Benha University in Egypt randomized 200 health care and household contacts of patients with COVID-19 where the intervention group consisted of 100 patients given a high dose of 0.4 mg/kg on day 1 and a second dose on day 7 in addition to wearing personal protective equipment, whereas the control group of 100 contacts wore personal protective equipment alone. They reported a large and statistically significant reduction in contacts testing positive by Reverse Transcriptase Polymerase Chain Reaction (PCR) when treated with ivermectin versus controls, 2% versus 10%, $P < 0.05$.

Shouman conducted an RCT at Zagazig University in Egypt, including 340 (228 treated and 112 control) family members of patients positive for SARS-CoV-2 through PCR.⁴⁴ Ivermectin (approximately 0.25 mg/kg) was administered twice, on the day of the positive test and 72 hours later. After a two-week follow-up, a large and statistically significant decrease in COVID-19 symptoms among household members treated with ivermectin was found, 7.4% versus 58.4%, $P < 0.001$.

Recently, Alam et al from Bangladesh performed a prospective observational study of 118 patients who were evenly split into those who volunteered for either the treatment or control arms, described as a persuasive approach. Although this method, along with the study being unblinded, likely led to confounders, the difference between the 2 groups was so large (6.7% vs. 73.3%, $P < 0.001$) and similar to the other prophylaxis trial results that confounders alone are unlikely to explain such a result.⁴⁷ Carvallo et al also performed a prospective observational trial where they gave healthy volunteers ivermectin and carrageenan daily for 28 days and matched them to similarly healthy controls who did not take the medicines.⁴⁰ Of the 229 study subjects, 131 were treated with 0.2 mg of ivermectin drops taken by mouth 5 times per day. After 28 days, none of those receiving ivermectin in the prophylaxis group had tested positive for SARS-COV-2 versus 11.2% of patients in the control arm ($P < 0.001$). In a much larger follow-up prospective, observational controlled trial by the same

group that included 1195 health care workers, they found that over a 3-month period there were no infections recorded among the 788 workers who took weekly ivermectin prophylaxis, whereas 58% of the 407 controls had become ill with COVID-19. This study demonstrates that remarkable protection against transmission can be achieved among high-risk health care workers by taking 12 mg once weekly.⁴⁰ The Carvallo IVERCAR protocol was also separately tested in a prospective RCT by the Health Ministry of Tucuman, Argentina, where they found that among 234 health care workers, the intervention group that took 12 mg once weekly, only 3.4% contracted COVID-19 versus 21.4% of controls, $P < .0001$.⁴⁶

The need for weekly dosing in the Carvallo study over a 4-month period may not have been necessary given that, in a recent RCT from Dhaka, Bangladesh, the intervention group ($n = 58$) took 12 mg once monthly for a similar 4-month period and also reported a large and statistically significant decrease in infections compared with controls, 6.9% versus 73.3%, $P < 0.05$.⁴⁷ Then, in a large retrospective observational case-control study from India, Behera et al⁴¹ reported that among 186 case-control pairs ($n = 372$) of health care workers, they identified 169 participants who had taken some form of prophylaxis, with 115 participants that had taken ivermectin. After matched pair analysis, they reported that in the workers who had taken 2 dose ivermectin prophylaxis, the odds ratio for contracting COVID-19 was markedly decreased (0.27, 95% confidence interval (CI) 0.15–0.51). Notably, one dose prophylaxis was not found to be protective in this study. Based on both their study finding and the Egyptian prophylaxis study, the All India Institute of Medical Sciences instituted a prophylaxis protocol for their health care workers where they now take two 0.3 mg/kg doses of ivermectin 72 hours apart and repeat the dose monthly.

Data that further illuminates the potential protective role of ivermectin against COVID-19 come from a study of nursing home residents in France which reported that in a facility that suffered a scabies outbreak where all 69 residents and 52 staff were treated with ivermectin,⁴¹ they found that during the period surrounding this event, 7 of the 69 residents fell ill with COVID-19 (10.1%). In this group with an average age of 90 years, only one resident required oxygen support and no resident died. In a matched control group of residents from surrounding facilities, they found 22.6% of residents fell ill and 4.9% died.

Further evidence supporting the efficacy of ivermectin as a prophylaxis agent was published recently in the *International Journal of Antimicrobial Agents* where a group of researchers analyzed data using the prophylactic chemotherapy databank administered by the WHO along with case counts obtained by Worldometers, a public data

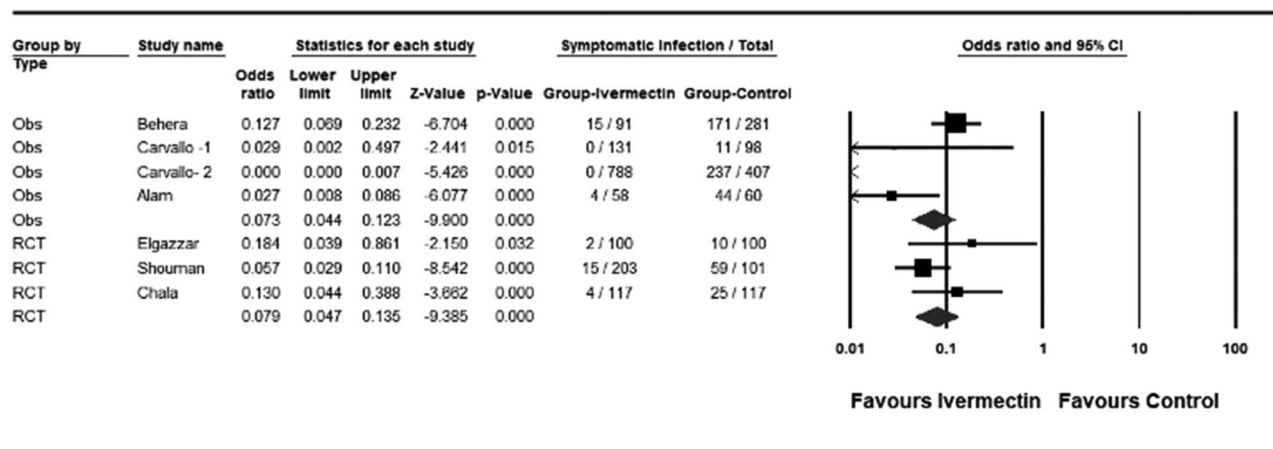


FIGURE 1. Meta-analysis of ivermectin prophylaxis trials in COVID-19. OBS, observational study; RCT, randomized controlled trial. Symbols: Squares: Indicate treatment effect of an individual study. Large diamond: Reflect summary of study design immediately above. Size of each symbol correlates with the size of the confidence interval around the point estimate of treatment effect with larger sizes indicating a more precise confidence interval.

aggregation site used by among others, the Johns Hopkins University.⁴² When they compared the data from countries with active ivermectin mass drug administration programs for the prevention of parasite infections, they discovered that the COVID-19 case counts were significantly lower in the countries with recently active programs, to a high degree of statistical significance, $P < 0.001$.

Figure 1 presents a meta-analysis performed by the study authors of the controlled ivermectin prophylaxis trials in COVID-19.

Further data supporting a role of ivermectin in decreasing transmission rates can be found from South American countries where, in retrospect, large “natural experiments” seem to have occurred. For instance, beginning as early as May, various regional health ministries and governmental authorities within Peru, Brazil, and Paraguay initiated “ivermectin distribution” campaigns to their citizen populations.⁴⁸ In one such example from Brazil, the cities of Itajai, Macapa, and Natal distributed massive amounts of ivermectin doses to their city’s

population, where in the case of Natal, 1 million doses were distributed. The distribution campaign of Itajai began in mid-July, in Natal they began on June 30th, and in Macapa, the capital city of Amapa and others nearby, they incorporated ivermectin into their treatment protocols in late May after they were particularly hard hit in April. The data in Table 1 were obtained from the official Brazilian government site and the national press consortium and show large decreases in case counts in the 3 cities soon after distribution began compared with their neighboring cities without such campaigns.

The decreases in case counts among the 3 Brazilian cities given in Table 1 were also associated with reduced mortality rates as summarized in Table 2.

Clinical studies on the efficacy of ivermectin in treating mildly ill outpatients

Currently, 7 trials that include a total of more than 3000 patients with mild outpatient illness have been completed, a set composed of 7 RCTs and 4 case series.⁴⁹⁻⁶⁰

Table 1. Comparison of case count decreases among Brazilian cities with and without ivermectin distribution campaigns.

| Region | New cases | June | July | August | Population 2020 (1000) | % Decline in new cases between June and August 2020 |
|------------|---------------|-------------|-------------|-------------|------------------------|---|
| South | Itajai | 2123 | 2854 | 998 | 223 | -53% |
| | Chapecó | 1760 | 1754 | 1405 | 224 | -20% |
| North | Macapá | 7966 | 2481 | 2370 | 503 | -70% |
| | Ananindeua | 1520 | 1521 | 1014 | 535 | -30% |
| North East | Natal | 9009 | 7554 | 1590 | 890 | -82% |
| | João Pessoa | 9437 | 7963 | 5384 | 817 | -43% |

Bolded cities distributed ivermectin, neighboring regional city below did not.

Table 2. Change in death rates among neighboring regions in Brazil.

| Region | State | % Change in average deaths/week compared with 2 weeks before |
|------------|----------------------------|--|
| South | Santa Catarina | -36% |
| | PARANÁ | -3% |
| | Rio Grande do Sul | -5% |
| North | Amapá | -75% |
| | AMAZONAS | -42% |
| | Pará | +13% |
| North East | Rio Grande do Norte | -65% |
| | CEARÁ | +62% |
| | Paraíba | -30% |

Bolded regions contained a major city that distributed ivermectin to its citizens, the other regions did not.

The largest, a double-blinded RCT by Mahmud⁴⁹ was conducted in Dhaka, Bangladesh, and targeted 400 patients with 363 patients completing the study. In this study, as in many other of the clinical studies to be reviewed, either a tetracycline (doxycycline) or macrolide antibiotic (azithromycin) was included as part of the treatment. The importance of including antibiotics such as doxycycline or azithromycin is unclear; however, both tetracycline and macrolide antibiotics have recognized anti-inflammatory, immunomodulatory, and even antiviral effects (58–61). Although the posted data from this study does not specify the amount of mildly ill outpatients versus hospitalized patients treated, important clinical outcomes were profoundly affected, with increased rates of early improvement (60.7% vs. 44.4% $P < 0.03$) and decreased rates of clinical deterioration (8.7% vs. 17.8%, $P < 0.02$). Given that mildly ill outpatients mainly comprised the study cohort, only 2 deaths were observed (both in the control group).

Ravikirti performed a double-blinded RCT of 115 patients, and although the primary outcome of PCR positivity on day 6 was no different, the secondary outcome of mortality was 0% versus 6.9%, $P = .019$.⁶⁰ Babalola in Nigeria also performed a double-blinded RCT of 62 patients, and in contrast to Ravikirti, they found a significant difference in viral clearance between both the low-dose and high-dose treatment groups and controls in a dose dependent fashion, $P = .006$.⁵⁹

Another RCT by Hashim et al⁵³ in Baghdad, Iraq, included 140 patients equally divided; the control group received standard care, and the treated group included a combination of both outpatient and hospitalized patients. In the 96 patients with mild-to-moderate outpatient illness, they treated 48 patients with a combination of ivermectin/doxycycline and standard of care and compared outcomes with the 48 patients treated with standard of care alone. The standard of care in this trial

included medicines such as dexamethasone 6 mg/d or methylprednisolone 40 mg twice per day if needed, vitamin C 1000 mg twice/day, zinc 75–125 mg/d, vitamin D3 5000 IU/day, azithromycin 250 mg/d for 5 days, and acetaminophen 500 mg as needed. Although no patients in either group progressed or died, the time to recovery was significantly shorter in the ivermectin-treated group (6.3 days vs. 13.7 days, $P < 0.0001$).

Chaccour et al conducted a small, double-blinded RCT in Spain where they randomized 24 patients to ivermectin versus placebo, and although they found no difference in PCR positivity at day 7, they did find statistically significant decreases in viral loads, patient days of anosmia (76 vs. 158, $P < 0.05$), and patient days with cough (68 vs. 98, $P < 0.05$).⁵⁷

Another RCT of ivermectin treatment in 116 outpatients was performed by Chowdhury et al in Bangladesh where they compared a group of 60 patients treated with the combination of ivermectin/doxycycline to a group of 60 patients treated with hydroxychloroquine/doxycycline with a primary outcome of time to negative PCR.⁵⁴ Although they found no difference in this outcome, in the treatment group, the time to symptomatic recovery approached statistical significance (5.9 days vs. 7.0 days, $P = 0.07$). In another smaller RCT of 62 patients by Podder et al, they also found a shorter time to symptomatic recovery that approached statistical significance (10.1 days vs. 11.5 days, $P > 0.05$, 95% CI, 0.86–3.67).⁵⁵

A medical group in the Dominican Republic reported a case series of 2688 consecutive symptomatic outpatients seeking treatment in the emergency department, most whom were diagnosed using a clinical algorithm. The patients were treated with a high-dose ivermectin of 0.4 mg/kg for one dose along with 5 days of azithromycin. Remarkably, only 16 of the 2688 patients (0.59%) required subsequent hospitalization with only a single death recorded.⁶¹

In another case series of 100 patients in Bangladesh, all treated with a combination of 0.2 mg/kg ivermectin and doxycycline, they found that no patient required hospitalization nor died, and all patients' symptoms improved within 72 hours.⁶²

A case series from Argentina reported on a combination protocol that used ivermectin, aspirin, dexamethasone, and enoxaparin. In the 135 mild illness patients, all survived.⁵⁰ Similarly, a case series from Mexico of 28 consecutively treated patients with ivermectin, all were reported to have recovered with an average time to full recovery of only 3.6 days.⁵⁸

Clinical studies of the efficacy of ivermectin in hospitalized patients

Studies of ivermectin among more severely ill hospitalized patients include 6 RCTs, 5 OCTs, and a database analysis study.^{45,51–53,63–70}

The largest RCT in hospitalized patients was performed concurrent with the prophylaxis study reviewed above by Elgazzar et al.⁴⁵ Four hundred patients were randomized among 4 treatment groups of 100 patients each. Groups 1 and 2 included mild/moderate illness patients alone, with group 1 treated with one dose 0.4 mg/kg ivermectin plus standard of care (SOC) and group 2 received hydroxychloroquine 400 mg twice on day 1 then 200 mg twice daily for 5 days plus standard of care. There was a statistically significant lower rate of progression in the ivermectin-treated group (1% vs. 22%, $P < 0.001$), with no deaths and 4 deaths, respectively. Groups 3 and 4 included only severely ill patients, with group 3 again treated with a single dose of 0.4 mg/kg plus SOC, whereas group 4 received hydroxychloroquine plus SOC. In this severely ill subgroup, the differences in outcomes were even larger, with lower rates of progression 4% versus 30% and mortality 2% versus 20% ($P < 0.001$).

The one largely outpatient RCT conducted by Hashim reviewed above also included 22 hospitalized patients in each group. In the ivermectin/doxycycline-treated group, there were 11 severely ill patients and 11 critically ill patients, whereas in the standard of care group, only severely ill patients ($n = 22$) were included because of their ethical concerns of including critically ill patients in the control group (45). This decision led to a marked imbalance in the severity of illness between these hospitalized patient groups. However, despite the mismatched severity of illness between groups and the small number of patients included, beneficial differences in outcomes were seen, but not all reached statistical significance. For instance, there was a large reduction in the rate of progression of illness (9% vs. 31.8%, $P = 0.15$) and, most importantly, there was a large difference in mortality among the severely ill groups that reached a borderline statistical significance (0% vs. 27.3%, $P = 0.052$). Another

important finding was the relatively low mortality rate of 18% found among the subset of critically ill patients, all of whom were treated with ivermectin.

A recent RCT from Iran found a dramatic reduction in mortality with ivermectin use.⁶⁵ Among multiple ivermectin treatment arms (different ivermectin dosing strategies were used in the intervention arms), the average mortality was reported as 3.3%, whereas the average mortality within the standard care and placebo arms was 18.8%, with an odds ratio (OR) of 0.18 (95% CI 0.06–0.55, $P < 0.05$).

Spoorthi⁶⁴ and Sasanak performed a prospective trial of 100 hospitalized patients whereby they treated 50 with ivermectin and doxycycline, whereas the 50 controls were given a placebo consisting of vitamin B6. Although no deaths were reported in either group, the ivermectin treatment group had a statistically significant shorter hospital length of stay (LOS) 3.7 days versus 4.7 days, $P = 0.03$, and shorter time to complete resolution of symptoms, 6.7 days versus 7.9 days, $P = 0.01$.

The largest OCT ($n = 280$) in hospitalized patients was conducted by Rajter et al at Broward Health Hospitals in Florida and was recently published in the major medical journal *Chest* (43). They performed a retrospective OCT using a propensity-matched design on 280 consecutive treated patients and compared those treated with ivermectin to those without. One hundred seventy-three patients were treated with ivermectin (160 received a single dose and 13 received a second dose at day 7) while 107 were not.⁶³ In both unmatched and propensity-matched cohort comparisons, similar, large, and statistically significant lower mortality was found among ivermectin-treated patients (15.0% vs. 25.2%, $P = 0.03$). Furthermore, in the subgroup of patients with severe pulmonary involvement, mortality was profoundly reduced when treated with ivermectin (38.8% vs. 80.7%, $P = 0.001$).

Another large OCT in Bangladesh compared 115 patients treated with ivermectin to a standard care cohort consisting of 133 patients.⁵¹ Despite a significantly higher proportion of patients in the ivermectin group being men (ie, with well-described, lower survival rates in COVID), the groups were otherwise well matched, yet the mortality decrease was statistically significant (0.9% vs. 6.8%, $P < 0.05$). The largest OCT is a study from Brazil, published as a letter to the editor and included almost 1500 patients.⁶⁶ Although the primary data were not provided, they reported that in 704 hospitalized patients treated with a single dose of 0.15 mg/kg ivermectin, compared with 704 controls, overall mortality was reduced (1.4% vs. 8.5%, HR 0.2, 95% CI 0.12–0.37, $P < 0.0001$). Similarly, in the patients on mechanical ventilation, mortality was also reduced (1.3% vs. 7.3%). A small study from Baghdad, Iraq, compared 16 ivermectin-treated patients with 71

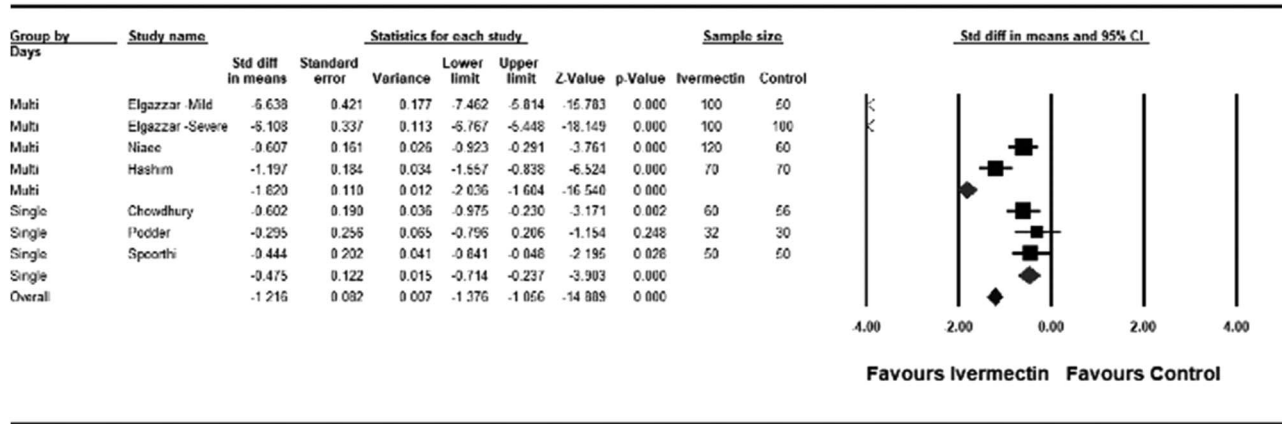


FIGURE 2. Meta-analysis of the outcome of time to clinical recovery from controlled trials of ivermectin treatment in COVID-19. OBS, observational study; RCT, randomized controlled trial. Symbols: Squares: Indicate treatment effect of an individual study. Large diamond: Reflect summary of study design immediately above. Small diamond: Sum effect of all trial designs. Size of each symbol correlates with the size of the confidence interval around the point estimate of treatment effect with larger sizes indicating a more precise confidence interval.

controls.⁵² This study also reported a significant reduction in length of hospital stay (7.6 days vs. 13.2 days, $P < 0.001$) in the ivermectin group. In a study reporting on the first 1000 patients treated in a hospital in India, they found that in the 34 patients treated with ivermectin alone, all recovered and were discharged, whereas in more than 900 patients treated with other agents, there was an overall mortality of 11.1%.⁷⁰

Meta-analyses of the above controlled treatment trials were performed by the study authors focused on

the 2 important clinical outcomes: time to clinical recovery and mortality (Figures 2 and 3). The consistent and reproducible signals leading to large overall statistically significant benefits from within both study designs are remarkable, especially given that in several of the studies treatment was initiated late in the disease course.

Details of the prophylaxis, early, and late treatment trials of ivermectin in COVID-19 can be found in Table 3.

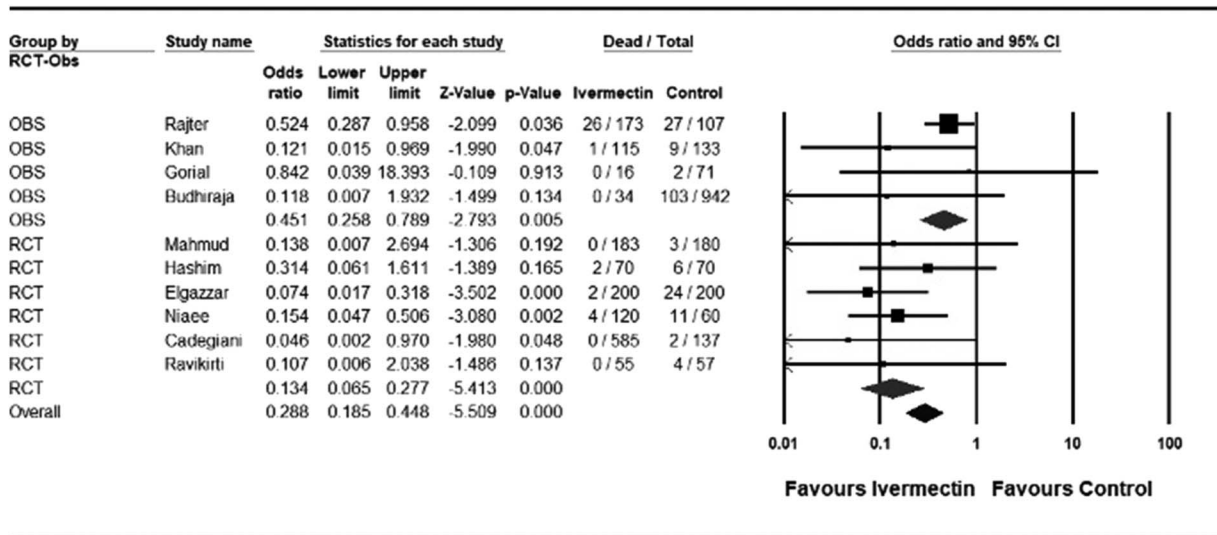


FIGURE 3. Meta-analysis of the outcome of mortality from controlled trials of ivermectin treatment in COVID-19. OBS, observational study; RCT, randomized controlled trial. Symbols: Squares: Indicate treatment effect of an individual study. Large diamond: Reflect summary of study design immediately above. Small diamond: Sum effect of all trial designs. Size of each symbol correlates with the size of the confidence interval around the point estimate of treatment effect with larger sizes indicating a more precise confidence interval.

Table 3. Clinical studies assessing the efficacy of ivermectin in the prophylaxis and treatment of COVID-19.

| Prophylaxis Trials Author, Country, source | Study design, size | Study subjects | Ivermectin dose | Dose frequency | Clinical outcomes reported |
|--|--------------------------------------|---|--|-----------------------------|---|
| Prophylaxis trials | | | | | |
| Shouman W, Egypt <i>www.clinicaltrials.gov</i> NCT04422561 | RCT N = 340 | Household members of pts with +COVID-19 PCR test | 40–60 kg: 15 mg, 60–80 kg: 18 mg, and > 80 kg: 24 mg | Two doses, 72 hours apart | 7.4% versus 58.4% developed COVID-19 symptoms, $P < 0.001$ |
| Elgazzar A, Egypt ResearchSquare doi.org/10.21203/rs.3.rs-100956/v1 | RCT N = 200 | Health care and household contacts of pts with +COVID-19 PCR test | 0.4 mg/kg | Two doses, day 1 and day 7 | 2% versus 10% tested positive for COVID-19 $P < 0.05$ |
| Chala R, Argentina NCT04701710 <i>Clinicaltrials.gov</i> | RCT N = 234 | Health care workers | 12 mg | Every 7 d | 3.4% versus 21.4%, $P = 0.0001$. |
| Carvallo H, Argentina <i>Journal of Biochemical Research and Investigation</i> doi.org/10.31546/2633–8653.1007 | OCT N = 229 | Healthy patients negative for COVID-19 PCR test | 0.2 mg drops | 1 drop 5 times a d x 28 d | 0.0% versus 11.2% contracted COVID-19 $P < 0.001$ |
| Alam MT, Bangladesh <i>European J Med Hlth Sciences</i> 10.24018/ejmed.2020.2.6.599 | OCT N = 118 | Health care workers | 12 mg | Monthly | 6.9% versus 73.3%, $P < 0.05$ |
| Carvallo H, Argentina <i>Journal of Biochemical Research and Investigation</i> doi.org/10.31546/2633–8653.1007 | OCT N = 1195 | Health care workers | 12 mg | Once weekly for up to 10 wk | 0.0% of the 788 workers taking ivermectin versus 58% of the 407 controls contracted COVID-19. |
| Behera P, India <i>medRxiv</i> doi.org/10.1101/2020.10.29.20222661 | OCT N = 186 case control pairs | Health care workers | 0.3 mg/kg | Day 1 and day 4 | 2 doses reduced odds of contracting COVID-19 (OR 0.27 95% CI 0.16–0.53) |
| Bernigaud C, France <i>Annales de Dermatologie et de Venereologi</i> doi.org/10.1016/j.annder.2020.09.231 | OCT N = 69 case control pairs | Nursing home residents | 0.2 mg/kg | Once | 10.1% versus 22.6% residents contracted COVID-19 0.0% versus 4.9% mortality |
| Hellwig M, USA <i>J Antimicrobial Agents</i> doi.org/10.1016/j.ijantimicag.2020.106,248 | OCT N = 52 countries | Countries with and without IVM prophylaxis programs | Unknown | Variable | Significantly lower-case incidence of COVID-19 in African countries with IVM prophylaxis programs $P < 0.001$ |

Clinical trials–Outpatients

% Ivermectin versus % Controls

| Prophylaxis Trials Author, Country, source | Study design, size | Study subjects | Ivermectin dose | Dose frequency | Clinical outcomes reported |
|---|-------------------------|------------------------------|---|---|--|
| Mahmud R, Bangladesh <i>www.clinicaltrials.gov</i> NCT0452383 | DB-RCT N = 363 | Outpatients and hospitalized | 12 mg + doxycycline | Once, within 3 days of PCR+ test | Early improvement 60.7% versus 44.4%, $P < 0.03$, deterioration 8.7% versus 17.8%, $P < 0.02$ |
| Chowdhury A, Bangladesh <i>Research Square</i> doi.org/10.21203/rs.3.rs-38896/v1 | RCT N = 116 | Outpatients | 0.2 mg/kg + doxycycline | Once | Recovery time 5.9 versus 9.3 days ($P = 0.07$) |
| Ravikirti, India <i>medRxiv</i> doi.org/10.1101/2021.01.05.21249310 | DB-RCT N = 115 | Mild–moderate illness | 12 mg | Daily for 2 d | No diff in day 6 PCR + 0% versus 6.9% mortality, $P = 0.019$ |
| Babalola OE, Nigeria <i>medRxiv</i> doi.org/10.1101/2021.01.05.21249131 | DB-RCT N = 62 | Mild–moderate illness | 6 mg and 12 mg | Every 48 hours × 2 wk | Time to viral clearance: 4.6 days high dose versus 6.0 days low dose versus 9.1 days control ($P = 0.006$) |
| Podder CS, Bangladesh <i>IMC J Med Sci</i> 2020;14(2) | RCT N = 62 | Outpatients | 0.2 mg/kg | Once | Recovery time 10.1 versus 11.5 days (NS), average time 5.3 versus 6.3 (NS) |
| Chaccour C. Spain <i>Research Square</i> doi.org/10.21203/rs.3.rs-116547/v1 | DB-RCT N = 24 | Outpatients | 0.4 mg/kg | Once | No diff in PCR+ day 7, lower viral load d 4 and 7, ($P < 0.05$), 76 versus 158 pts. d of anosmia ($P < 0.05$), 68 versus 98 pts. d of cough ($P < 0.05$) |
| Morgenstern J, Dominican Republic <i>medRxiv</i> doi.org/10.1101/2020.10.29.20222505 | Case series N = 3099 | Outpatients and hospitalized | Outpatients: 0.4 mg/kg hospital patients: 0.3 mg/kg | Outpatients: 0.3 mg/kg × 1 dose Inpatients: 0.3 mg/kg, days 1,2,6, and 7 | Mortality = 0.03% in 2688 outpatients, 1% in 300 non-ICU hospital patients, and 30.6% in 111 ICU patients |
| Carvalho H, Argentina <i>medRxiv</i> doi.org/10.1101/2020.09.10.20191619 | Case series N = 167 | Outpatients and hospitalized | 24 mg = mild, 36 mg = moderate, and 48 mg = severe | Days 0 and 7 | All 135 with mild illness survived, 1/32 (3.1% of hospitalized) patients died |
| Alam A, Banglades <i>J of Bangladesh College Phys and Surg</i> , 2020; 38:10-15 doi.org/10.3329/jbcps.v38i0.47512 | Case series N = 100 | Outpatients | 0.2 mg/kg/kg + doxycycline | Once | All improved within 72 h |

(Continued on next page)

Table 3. (Continued) Clinical studies assessing the efficacy of ivermectin in the prophylaxis and treatment of COVID-19.

| Clinical trials–Outpatients | | | | | | % Ivermectin versus % Controls |
|---|-----------------------|--------------------------------------|---|------------------------------------|---|--------------------------------|
| Prophylaxis Trials Author, Country, source | Study design, size | Study subjects | Ivermectin dose | Dose frequency | Clinical outcomes reported | |
| Espatia-Hernandez G, Mexico <i>Biomedical Research</i> www.biomedres.info/biomed...-proof-of-concept-study-14435.html | Case series N = 28 | Outpatients | 6 mg | Days 1,2, 7, and 8 | All pts recovered average recovery time 3.6 d | |
| Clinical trials–Hospitalized patients | | | | | | % Ivermectin versus % Controls |
| Prophylaxis Trials Author, Country, source | Study design, size | Study subjects | Ivermectin dose | Dose frequency | Clinical outcomes reported | |
| Elgazzar A, Egypt ResearchSquare doi.org/10.21203/rs.3.rs-100956/v1 | OL-RCT N = 400 | Hospitalized patients | 0.4 mg/kg | Daily for 4 days | Moderately ill: worsened 1% versus 22%, $P < 0.001$. Severely ill: worsened 4% versus 30% mortality 2% versus 20% both with $P < 0.001$ | |
| Niaee S. M, Research Square doi.org/10.21203/rs.3.rs-109670/v1 | DB-RCT N = 180 | Hospitalized patients | 0.2, 0.3, and 0.4 mg/kg (3 dosing strategies) | Once versus Days 1,3,5 | Mortality 3.3% versus 18.3%. OR 0.18, (0.06–0.55, $P < 0.05$) | |
| Hashim H, Iraq medRxiv doi.org/10.1101/2020.10.26.20219345 | SB-RCT N=140 | 2/3 outpatients and 1/3 hospital pts | 0.2 mg/kg + doxycycline | Daily for 2–3 d | Recovery time 6.3 versus 13.6 days ($P < 0.001$), 0% versus 27.3% mortality in severely ill ($P = 0.052$) | |
| Spoorthi S, India AIAM, 2020; 7(10):177-182 | PCT N = 100 | Hospitalized patients | 0.2 mg/kg+ doxycycline | Once | Shorter hospital LOS, 3.7 versus 4.7 days, $P = 0.03$, faster resolution of symptoms, 6.7 versus 7.9 days, $P = 0.01$ | |
| Ahmed S. Dhaka, Bangladesh International journal of Infectious disease doi.org/10.1016/j.ijid.2020.11.191 | DB-RCT N = 72 | Hospitalized patients | 12 mg | Daily for 5 d | Faster viral clearance 9.7 versus 12.7 days, $P = 0.02$ | |
| Chachar AZK, Pakistan Int J Sciences doi.org/10.18483/ijSci.2378 | DB-RCT N = 50 | Hospitalized patients-mild | 12 mg | Two doses day 1 and one dose day 2 | 64% versus 60% asymptomatic by day 7 | |
| Portman-Baracco A, Brazil | OCT | | 0.15 mg/kg | Once | | |

(Continued on next page)

Table 3. (Continued) Clinical studies assessing the efficacy of ivermectin in the prophylaxis and treatment of COVID-19.

| Clinical trials–Hospitalized patients | | | | | % Ivermectin versus % Controls |
|---|---------------------------|-----------------------|----------------------------------|---------------------------|---|
| Prophylaxis Trials Author, Country, source | Study design, size | Study subjects | Ivermectin dose | Dose frequency | Clinical outcomes reported |
| Arch Bronconeumol. 2020 doi.org/10.1016/j.arbres.2020.06.011 | N = 1408 | Hospitalized patients | | | Overall mortality 1.4% versus 8.5%, HR 0.2, 95% CI 0.12–0.37, $P < 0.0001$ |
| Rajter JC, Florida Chest 2020 doi.org/10.1016/j.chest.2020.10.009 | OCT N=280 | Hospitalized patients | 0.2 mg/kg + azithromycin | Day 1 and day 7 if needed | Overall mortality 15.0% versus 25.2%, $P = 0.03$, severe illness mortality 38.8% versus 80.7%, $P = 0.001$ |
| Khan X, Bangladesh Arch Bronconeumol. 2020 doi.org/10.1016/j.arbres.2020.08.007 | OCT N = 248 | Hospitalized patients | 12 mg | Once on admission | Mortality 0.9% versus 6.8%, $P < 0.05$, LOS 9 versus 15 days, $P < 0.001$ |
| Gorial FI, Iraq medRxiv doi.org/10.1101/2020.07.07.20145979 | OCT N = 87 | Hospitalized patients | 0.2 mg/kg + HCQ and azithromycin | Once on admission | LOS 7.6 versus 13.2 days, $P < 0.001$, 0/15 versus 2/71 died |
| Budiraja S. India medRxiv doi.org/10.1101/2020.11.16.20232223 | OCT N = 1000 IVM=34 | Hospitalized patients | n/a | n/a | 100% IVM pts recovered 11.1% mortality in non-IVM-treated pts |

DB-RCT, double-blinded randomized controlled trial; HCQ, hydroxychloroquine; IVM, ivermectin; LOS, length of stay; NS, nonstatistically significant, $P > .05$; OCT, observational controlled trial; OL, open label; PCR, polymerase chain reaction; RCT, randomized controlled trial; SB-RCT, single blinded randomized controlled trial.

Ivermectin in post-COVID-19 syndrome

Increasing reports of persistent, vexing, and even disabling symptoms after recovery from acute COVID-19 have been reported and that many have termed the condition as “Long COVID” and patients as “long haulers,” estimated to occur in approximately 10%–30% of cases.^{71–73} Generally considered as a postviral syndrome consisting of a chronic and sometimes disabling constellation of symptoms which include, in order, fatigue, shortness of breath, joint pains, and chest pain. Many patients describe their most disabling symptom as impaired memory and concentration, often with extreme fatigue, described as “brain fog,” and is highly suggestive of the condition myalgic encephalomyelitis/chronic fatigue syndrome, a condition well reported to begin after viral infections, in particular with Epstein–Barr virus. Although no specific treatments have been identified for Long COVID, a recent manuscript by Aguirre-Chang et al from the National University of San Marcos in Peru reported on their experience with ivermectin in such patients.⁷⁴ They treated 33 patients who were between 4 and 12 weeks from the onset of symptoms with escalating doses of ivermectin; 0.2 mg/kg for 2 days if mild and 0.4 mg/kg for 2 days if moderate, with doses extended if symptoms persisted. They found that in 87.9% of the patients, resolution of all symptoms was observed after 2 doses with an additional 7% reporting complete resolution after additional doses. Their experience suggests the need for controlled studies to better test efficacy in this vexing syndrome.

Epidemiological data showing impacts of widespread ivermectin use on population case counts and case fatality rates

Similar to the individual cities in Brazil that measured large decreases in case counts soon after distributing ivermectin in comparison to neighboring cities without such campaigns, in Peru, the government approved the use of ivermectin by decree on May 8, 2020, solely based on the *in vitro* study by Caly et al from Australia.⁴⁸ Soon after, multiple state health ministries initiated ivermectin distribution campaigns in an effort to decrease what was at that time some of the highest COVID-19 morbidity and mortality rates in the world. Juan Chamie,⁴⁸ a data analyst and member of the FLCCC Alliance, recently posted an article based on 2 critical sets of data that he compiled and compared; first, he identified the timing and magnitude of each region’s ivermectin interventions through a review of official communications, press releases, and the Peruvian Situation Room database to confirm the dates of effective delivery, and second, he extracted data on the total all-cause deaths from the region along with COVID-19 case counts in selected age groups over time

from the registry of the National Computer System of Deaths (SINADEF) and from the National Institute of Statistics and Informatics.⁴⁸ It should be noted that he restricted his analyses to only those citizens older than 60 years to avoid the confounding of rises in the numbers of infected younger patients. With these data, he was then able to compare the timing of major decreases in this age group of both total COVID-19 cases and total excess deaths per 1000,000 people among 8 states in Peru with the initiation dates of their respective ivermectin distribution campaigns as shown in Figure 4.

Figure 5 from the same study presents data on the case fatality rates in patients older than 60 years, again among the 8 states in Peru. Note the dramatically decreased case fatality rates among older patients diagnosed with COVID-19 after ivermectin became widely distributed in those areas, a result which cannot be explained by changes in mask-wearing or lock-downs.

In an even more telling example, Chamie compared the case counts and fatality rates of the 8 states above with the city of Lima, where ivermectin was not distributed nor widely used in treatment during the same period. Figure 6 compares the lack of significant or sustained reductions in case counts or fatalities in Lima with the dramatic reductions in both outcomes among the 8 states with widespread ivermectin distribution.

Another example can be seen from the data compiled from Paraguay, again by Chamie who noted that the government of the state of Alto Parana had launched an ivermectin distribution campaign in early September. Although the campaign was officially described as a “deworming” program, this was interpreted as a guise by the regions’ governor to avoid reprimand or conflict with the National Ministry of Health that recommended against the use of ivermectin to treat COVID-19 in Paraguay. The program began with a distribution of 30,000 boxes of ivermectin, and by October 15, the governor declared that there were very few cases left in the state as can be seen in Figure 7.

The evidence base for ivermectin against COVID-19

To date, the efficacy of ivermectin in COVID-19 has been supported by the following:

1. Since 2012, multiple *in vitro* studies have demonstrated that Ivermectin inhibits the replication of many viruses, including influenza, Zika, Dengue, and others.^{9–17}
2. Ivermectin inhibits SARS-CoV-2 replication and binding to host tissue through several observed and proposed mechanisms.¹⁸
3. Ivermectin has potent anti-inflammatory properties with *in vitro* data demonstrating profound

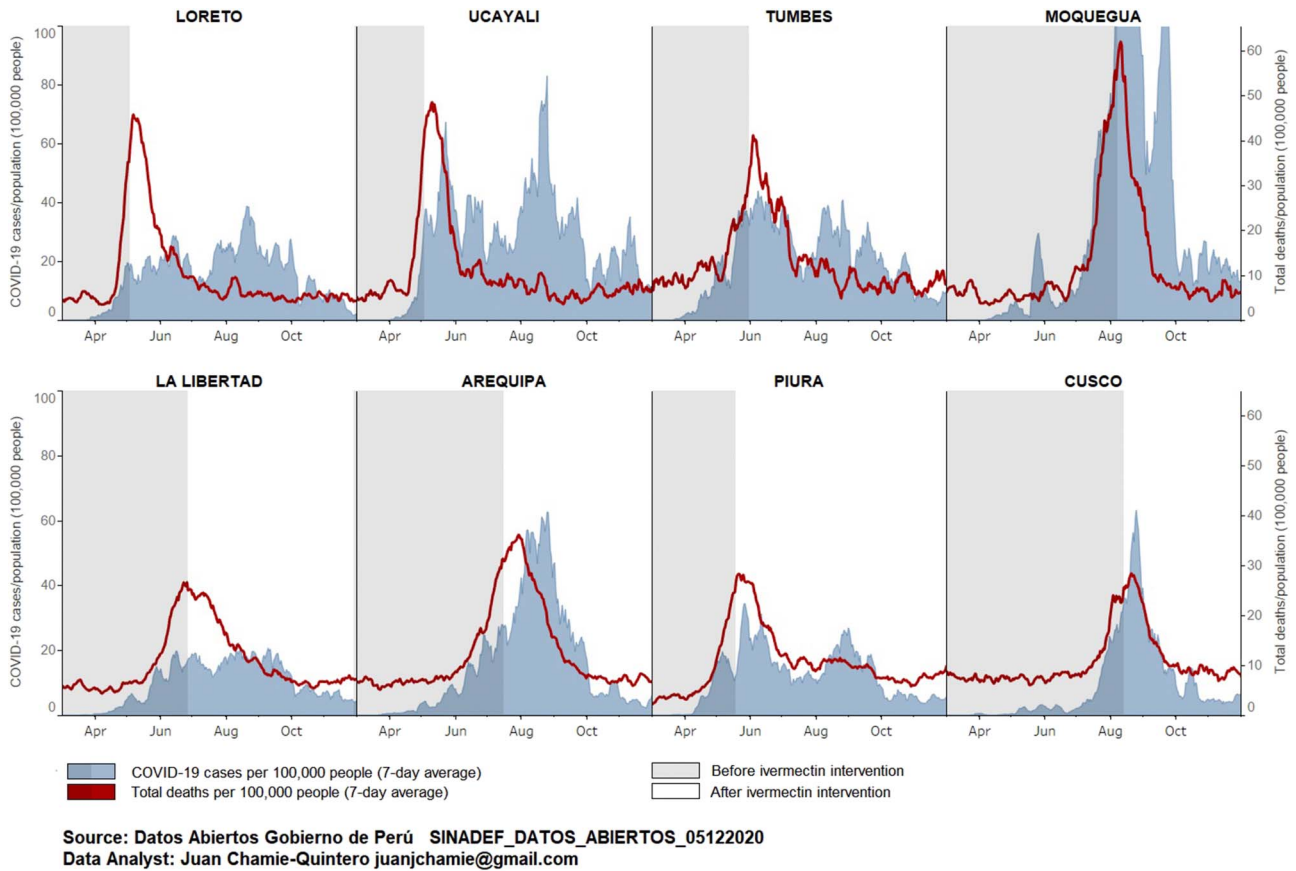


FIGURE 4. Decrease in total case incidences and total deaths/population of COVID-19 in the over 60 population among 8 Peruvian states after deploying mass ivermectin distribution campaigns.

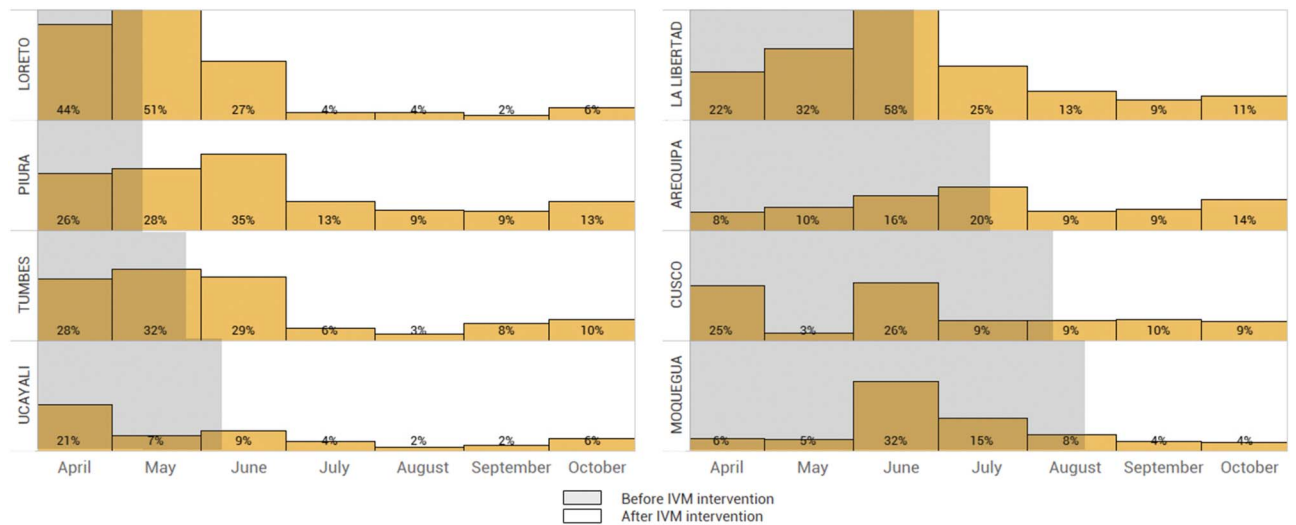


FIGURE 5. Daily total deaths, case fatalities, and case incidence for COVID-19 in populations of patients aged 60 and older for 8 states in Peru deploying early mass ivermectin treatments versus the state of Lima, including the capital city, where ivermectin treatment was applied months later.

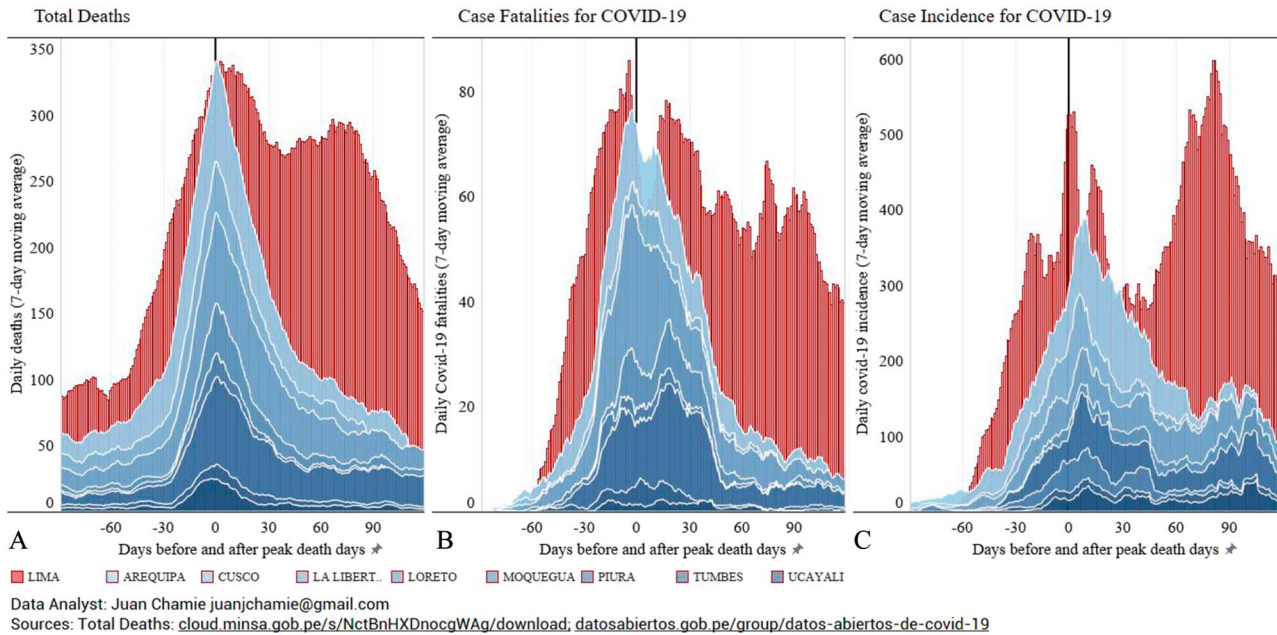


FIGURE 6. Covid-19 case fatalities and total deaths with and without mass ivermectin in different states of Peru.

inhibition of both cytokine production and transcription of nuclear factor- κ B (NF- κ B), the most potent mediator of inflammation.³⁷⁻³⁹

4. Ivermectin significantly diminishes viral load and protects against organ damage in multiple animal

models when infected with SARS-CoV-2 or similar coronaviruses.^{31,32}

5. Ivermectin prevents transmission and development of COVID-19 disease in those exposed to infected patients.⁴⁰⁻⁴⁵

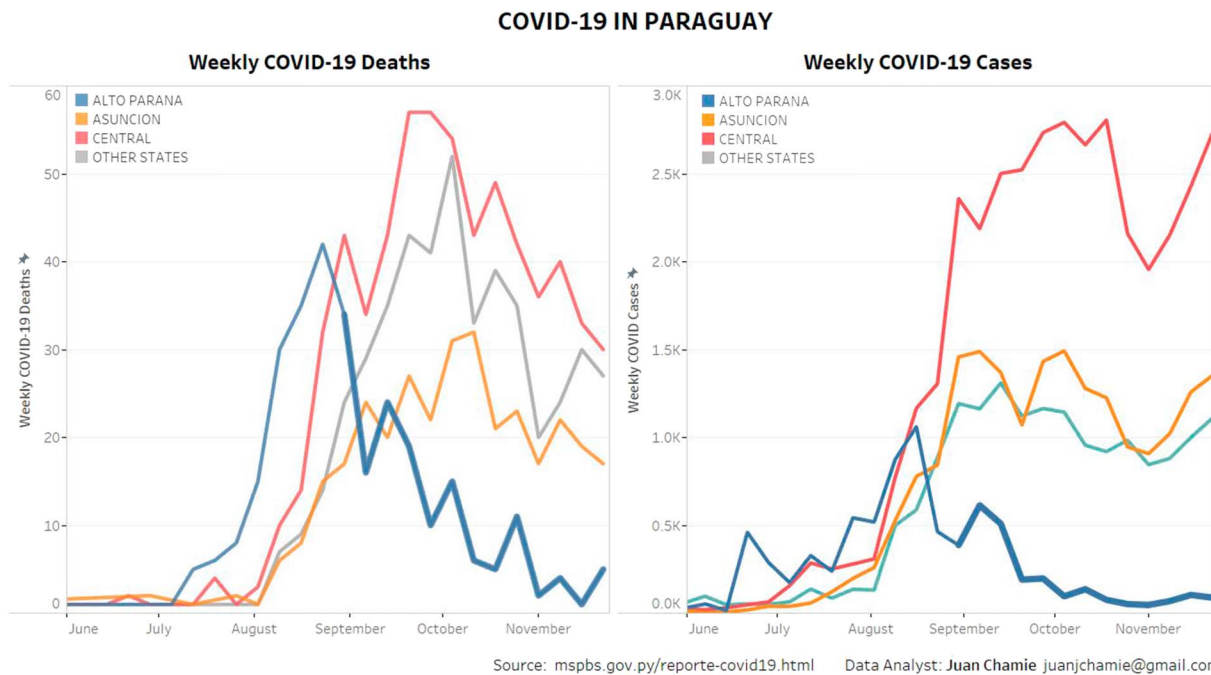


FIGURE 7. Paraguay – COVID-19 case counts and deaths in Alto Parana (bolded blue line) after ivermectin distribution began compared to other regions.

6. Ivermectin hastens recovery and prevents deterioration in patients with mild to moderate disease treated early after symptoms.^{45,49–52,61,62}
7. Ivermectin hastens recovery and avoidance of ICU admission and death in hospitalized patients.^{45,51,53,63–66}
8. Ivermectin reduces mortality in critically ill patients with COVID-19.^{45,53,63}
9. Ivermectin leads to temporally associated reductions in case fatality rates in regions after ivermectin distribution campaigns.⁴⁸
10. The safety, availability, and cost of ivermectin are nearly unparalleled given its low incidence of important drug interactions along with only mild and rare side effects observed in almost 40 years of use and billions of doses administered.⁷⁵
11. The World Health Organization has long included ivermectin on its “List of Essential Medicines.”

A summary of the statistically significant results from the above controlled trials are as follows:

Controlled trials in the prophylaxis of COVID-19 (8 studies)

1. All 8 available controlled trial results show statistically significant reductions in transmission.
2. Three RCTs with large statistically significant reductions in transmission rates, N = 774 patients.^{44–46}
3. Five OCTs with large statistically significant reductions in transmission rates, N = 2052 patients.^{40–43,47}

Controlled trials in the treatment of COVID-19 (19 studies)

1. Five RCTs with statistically significant impacts in time to recovery or hospital length of stay.^{45,49,53,64,65}
2. One RCT with a near statistically significant decrease in time to recovery, $P = 0.07$, N = 130.⁵⁴
3. One RCT with a large, statistically significant reduction in the rate of deterioration or hospitalization, N = 363.⁴⁹
4. Two RCTs with a statistically significant decrease in viral load, days of anosmia, and cough, N = 85.^{57,60}
5. Three RCTs with large, statistically significant reductions in mortality (N = 695).^{45,60,65}
6. One RCT with a near statistically significant reduction in mortality, $P = 0.052$ (N = 140).⁵³
7. Three OCTs with large, statistically significant reductions in mortality (N = 1688).^{51,63,66}

Safety of ivermectin

Numerous studies report low rates of adverse events, with the majority mild, transient, and largely attributed to the body's inflammatory response to the death of the parasites

and include itching, rash, swollen lymph nodes, joint pains, fever, and headache.⁷⁵ In a study that combined results from trials including more than 50,000 patients, serious events occurred in less than 1% and largely associated with administration in Loa loa.⁷⁶ Furthermore, according to the pharmaceutical reference standard *Lexicomp*, the only medications contraindicated for use with ivermectin are the concurrent administration of antituberculosis and cholera vaccines while the anticoagulant warfarin would require dose monitoring. Another special caution is that immunosuppressed or organ transplant patients who are on calcineurin inhibitors, such as tacrolimus or cyclosporine, or the immunosuppressant sirolimus should have close monitoring of drug levels when on ivermectin given that interactions exist that can affect these levels. A longer list of drug interactions can be found on the *drugs.com* database, with nearly all interactions leading to a possibility of either increased or decreased blood levels of ivermectin. Given studies showing tolerance and lack of adverse effects in human subjects given escalating high doses of ivermectin, toxicity is unlikely, although a reduced efficacy because of decreased levels may be a concern.⁷⁷

Concerns of safety in the setting of liver disease are unfounded given that, to the best of our knowledge, only 2 cases of liver injury have ever been reported in association with ivermectin, with both cases rapidly resolved without need for treatment.^{78,79} Furthermore, no dose adjustments are required in patients with liver disease. Some have described ivermectin as potentially neurotoxic, yet one study performed a search of a global pharmaceutical database and found only 28 cases among almost 4 billion doses with serious neurological adverse events, such as ataxia, altered consciousness, seizure, or tremor.⁸⁰ Potential explanations included the effects of concomitantly administered drugs that increase absorption past the blood-brain barrier or polymorphisms in the *mdr-1* gene. However, the total number of reported cases suggests that such events are exceedingly rare. Finally, ivermectin has been used safely in pregnant women, children, and infants.

DISCUSSION

Currently, as of December 14, 2020, there is accumulating evidence that demonstrates both the safety and efficacy of ivermectin in the prevention and treatment of COVID-19. Large-scale epidemiologic analyses validate the findings of in vitro, animal, prophylaxis, and clinical studies. Epidemiologic data from regions of the world with widespread ivermectin use have demonstrated a temporally associated reduction in case counts, hospitalizations, and fatality rates.

In the context of ivermectin's long-standing safety record, low cost, and wide availability along with the

consistent, reproducible, large magnitude of findings on transmission rates, need for hospitalization, and mortality, widespread deployment in both prevention and treatment has been proposed. Although a subset of trials are of an observational design, it must be recognized that in the case of ivermectin (1) half of the trials used a randomized controlled trial design (12 of the 24 reviewed above) and (2) observational and randomized trial designs reach equivalent conclusions on average as reported in a large Cochrane review of the topic from 2014.⁸¹ In particular, OCTs that use propensity-matching techniques (as in the Rajter study from Florida) find near identical conclusions to later-conducted RCTs in many different disease states, including coronary syndromes, critical illness, and surgery.^{82–84} Similarly, as evidenced in the prophylaxis (Figure 1) and treatment trial (Figures 2 and 3) meta-analyses as well as the summary trials table (Table 3), the entirety of the benefits found in both OCT and RCT trial designs aligns in both direction and magnitude of benefit. Such a consistency of benefit among numerous trials of varying sizes designs from multiple different countries and centers around the world is unique and provides strong, additional support.

The continued challenges faced by health care providers in deciding on appropriate therapeutic interventions in patients with COVID-19 would be greatly eased if more updated and commensurate evidence-based guidance came from the leading governmental health care agencies. Currently, in the United States, the treatment guidelines for COVID-19 are issued by the National Institutes of Health. Their most recent recommendation on the use of ivermectin in patients with COVID-19 was last updated on February 11, 2021, where they found that “there was insufficient evidence to recommend for or against ivermectin in COVID-19.” For a more definitive recommendation to be issued by major leading public health agencies (PHA), it is apparent that even more data on both the quality and quantity of trials are needed, even during a global health care emergency, and in consideration of a safe, oral, low-cost, widely available and deployable intervention such as ivermectin.

Fortunately, large teams sponsored by 2 different organizations have embarked on this effort. One team, sponsored by the Unitaid/WHO's ACT Accelerator Program and led by the University of Liverpool Senior Research Fellow Dr. Andrew Hill, is performing a systematic review and meta-analysis focused solely on ivermectin treatment RCTs in COVID-19. Although a preliminary meta-analysis of 17 RCTs was posted to a preprint server in February, it is expected that by March 19, 2021, results from approximately 27–29 RCTs including almost 4500 patients will be presented to the WHO Guidelines Committee and that the epidemiologic studies reviewed above

by Chamie et al were already presented to the committee in early March (personal communication with Dr. Andrew Hill). It is important to note that on February 5, the WHO Guidelines Committee announced that they had begun a review of the accumulating ivermectin data and expected to arrive at their own formal treatment recommendation within 4–6 weeks. If the above benefits in clinical outcomes continue to be reported in the remaining trials, it is hoped that this almost doubling of the current supportive evidence base would merit a recommendation for use by the WHO, NIH, and other PHA's would be forthcoming.

Because of the urgency of the pandemic, and in response to the surprising persistent inaction by the leading PHA's, the British Ivermectin Recommendation Development Panel was recently coordinated by the Evidence-Based Medicine Consultancy Ltd to more rapidly formulate an ivermectin treatment guideline using the standard guideline development process followed by the WHO. Made up of long-time research consultants to numerous national and international public health organizations including the WHO, they convened both a steering committee and a technical working group that then performed a systematic review and meta-analysis. On February 12, 2021, a meeting was held that included an international consortium of 75 practitioners, researchers, specialists, and patient representatives representing 16 countries and most regions of the world. This Recommendation Development Panel was presented the results of the meta-analysis of 18 treatment RCTs and 3 prophylaxis RCTs including more than 2500 patients along with a summary of the observational trials and the epidemiologic analyses related to regional ivermectin use. After a discussion period, a vote was held on multiple aspects of the data on ivermectin, according to standard WHO guideline development processes. The Panel *found the certainty of evidence for ivermectin's effects on survival to be strong and they recommended unconditional adoption for use in the prophylaxis and treatment of COVID-19.*

In summary, based on the totality of the trials and epidemiologic evidence presented in this review along with the preliminary findings of the Unitaid/WHO meta-analysis of treatment RCTs and the guideline recommendation from the international BIRD conference, ivermectin should be globally and systematically deployed in the prevention and treatment of COVID-19.

REFERENCES

1. Hermine O, Mariette X, Tharaux PL, et al. Effect of tocilizumab vs usual care in adults hospitalized with COVID-19 and moderate or severe pneumonia: a randomized clinical trial. *JAMA Intern Med.* 2021;181:32–40.

2. Salvarani C, Dolci G, Massari M, et al. Effect of tocilizumab vs standard care on clinical worsening in patients hospitalized with COVID-19 pneumonia: a randomized clinical trial. *JAMA Intern Med.* 2021;181:24–31.
3. Agarwal A, Mukherjee A, Kumar G, et al. Convalescent plasma in the management of moderate covid-19 in adults in India: open label phase II multicentre randomised controlled trial (PLACID Trial). *BMJ.* 2020;371:m3939.
4. Consortium WST. Repurposed antiviral drugs for covid-19— interim WHO solidarity trial results. *New Engl J Med.* 2020;384:497–511.
5. World Health Organization. Corticosteroids for COVID-19: living guidance. *World Health Organization*, September 2, 2020. Available at: <https://apps.who.int/iris/handle/10665/334125>.
6. Horby P, Lim WS, Emberson JR, et al. Dexamethasone in hospitalized patients with Covid-19—preliminary report. *New Engl J Med.* 2020;384:693–704.
7. Crump A, Omura S. Ivermectin, 'wonder drug' from Japan: the human use perspective. *Proc Jpn Acad Ser B.* 2011;87:13–28.
8. Tambo E, Khater EI, Chen JH, et al. Nobel prize for the artemisinin and ivermectin discoveries: a great boost towards elimination of the global infectious diseases of poverty. *Infect Dis Poverty.* 2015;4:58.
9. Atkinson SC, Audsley MD, Lieu KG, et al. Recognition by host nuclear transport proteins drives disorder-to-order transition in Hendra virus V. *Sci Rep.* 2018;8:358.
10. Yang SNY, Atkinson SC, Wang C, et al. The broad spectrum antiviral ivermectin targets the host nuclear transport importin $\alpha/\beta 1$ heterodimer. *Antivir Res.* 2020;177:104760.
11. Götz V, Magar L, Dornfeld D, et al. Influenza A viruses escape from MxA restriction at the expense of efficient nuclear vRNP import. *Sci Rep.* 2016;6:23138.
12. Lv C, Liu W, Wang B, et al. Ivermectin inhibits DNA polymerase UL42 of pseudorabies virus entrance into the nucleus and proliferation of the virus in vitro and vivo. *Antivir Res.* 2018;159:55–62.
13. Mastrangelo E, Pezzullo M, De Burghgraeve T, et al. Ivermectin is a potent inhibitor of flavivirus replication specifically targeting NS3 helicase activity: new prospects for an old drug. *J Antimicrob Chemother.* 2012;67:1884–1894.
14. Tay MYF, Fraser JE, Chan WKK, et al. Nuclear localization of dengue virus (DENV) 1–4 non-structural protein 5; protection against all 4 DENV serotypes by the inhibitor Ivermectin. *Antiviral Res.* 2013;99:301–306.
15. Varghese FS, Kaukinen P, Gläsker S, et al. Discovery of berberine, abamectin and ivermectin as antivirals against chikungunya and other alphaviruses. *Antivir Res.* 2016;126:117–124.
16. Wagstaff Kylie M, Sivakumaran H, Heaton Steven M, et al. Ivermectin is a specific inhibitor of importin α/β -mediated nuclear import able to inhibit replication of HIV-1 and dengue virus. *Biochem J.* 2012;443:851–856.
17. King CR, Tessier TM, Dodge MJ, et al. Inhibition of human adenovirus replication by the importin $\alpha/\beta 1$ nuclear import inhibitor ivermectin. *J Virol.* 2020;94:94.
18. Caly L, Druce JD, Catton MG, et al. The FDA-approved drug ivermectin inhibits the replication of SARS-CoV-2 in vitro. *Antivir Res.* 2020;178:104787.
19. Bray M, Rayner C, Noël F, et al. Ivermectin and COVID-19: a report in Antiviral Research, widespread interest, an FDA warning, two letters to the editor and the authors' responses. *Antiviral Res.* 2020;178:104805.
20. Schmith VD, Zhou J, Lohmer LR. The approved dose of ivermectin alone is not the ideal dose for the treatment of COVID-19. *Clin Pharmacol Ther.* 2020;103:214–216.
21. Dayer M.R. Coronavirus (2019-nCoV) deactivation via spike glycoprotein shielding by old drugs, bioinformatic study. *Preprints.* 2020. doi: 10.20944/preprints202005.0020.v1.
22. Maurya DK. A combination of ivermectin and doxycycline possibly blocks the viral entry and modulate the innate immune response in COVID-19 patients. *ChemRxiv.* 2020. doi: 10.26434/chemrxiv.12630539.v1.
23. Hussien MA, Abdelaziz AEM. Molecular docking suggests repurposing of brincidofovir as a potential drug targeting SARS-CoV-2 ACE2 receptor and main protease. *Netw Model Anal Health Inform Bioinform.* 2020;9:56–18.
24. Suravajhala R, Parashar A, Malik B, et al. Comparative docking studies on curcumin with COVID-19 proteins. *Preprints.* 2020. doi: 10.20944/preprints202005.0439.v1.
25. Nallusamy S, Mannu J, Ravikumar C, et al. Shortlisting phytochemicals exhibiting inhibitory activity against major proteins of SARS-CoV-2 through virtual screening. *Res Square.* 2020. Doi: 10.21203/rs.3.rs-31834/v1.
26. Lehrer S, Rheinstein PH. Ivermectin docks to the SARS-CoV-2 spike receptor-binding domain attached to ACE2. *Vivo.* 2020;34:3023–3026.
27. Scheim D. From cold to killer: how SARS-CoV-2 evolved without hemagglutinin esterase to agglutinate, then clot blood cells in pulmonary and systemic microvasculature. *SSRN.* 2020. Doi: 10.2139/ssrn.3706347.
28. Dasgupta J, Sen U, Bakshi A, et al. Nsp7 and spike glycoprotein of SARS-CoV-2 are envisaged as potential targets of vitamin D and ivermectin. *Preprints.* 2020. doi: 10.20944/preprints202005.0084.v1.
29. Sen Gupta PS, Biswal S, Panda SK, et al. Binding mechanism and structural insights into the identified protein target of COVID-19 and importin-alpha with in-vitro effective drug ivermectin. *J Biomol Struct Dyn.* 2020:1–10.
30. Swargiary A. Ivermectin as a Promising RNA-dependent RNA Polymerase Inhibitor and a Therapeutic Drug against SARS-CoV2: Evidence from in Silico Studies. *Res Square.* 2020. Doi: 10.21203/rs.3.rs-73308/v1.
31. Arevalo AP, Pagotto R, Porfido J, et al. Ivermectin reduces coronavirus infection in vivo: a mouse experimental model. *bioRxiv.* 2020. doi:10.1101/2020.11.

32. de Melo GD, Lazarini F, Larrous F, et al. Anti-COVID-19 efficacy of ivermectin in the golden hamster. *bioRxiv*. 2020. doi: 10.1101/2020.11.21.392639.
33. Perera RA, Tso E, Tsang OT, et al. SARS-CoV-2 virus culture from the upper respiratory tract: correlation with viral load, subgenomic viral RNA and duration of illness. *MedRxiv*. 2020. doi: 10.1101/2020.07.08.20148783.
34. Polak SB, Van Gool IC, Cohen D, et al. A systematic review of pathological findings in COVID-19: a pathophysiological timeline and possible mechanisms of disease progression. *Mod Pathol*. 2020;33:2128–2138.
35. Young BE, Ong SW, Ng LF, et al. Viral dynamics and immune correlates of COVID-19 disease severity. *Clin Infect Dis*. 2020. Available at: <https://bookcafe.yuntsq.com/ueditor/jsp/upload/file/20200921/1600658531176084970.pdf>.
36. Li Y, Chen M, Cao H, et al. Extraordinary GU-rich single-strand RNA identified from SARS coronavirus contributes an excessive innate immune response. *Microbes Infect*. 2013;15:88–95.
37. Zhang X, Song Y, Xiong H, et al. Inhibitory effects of ivermectin on nitric oxide and prostaglandin E2 production in LPS-stimulated RAW 264.7 macrophages. *Int Immunopharmacol*. 2009;9:354–359.
38. Ci X, Li H, Yu Q, et al. Avermectin exerts anti-inflammatory effect by downregulating the nuclear transcription factor kappa-B and mitogen-activated protein kinase activation pathway. *Fundam Clin Pharmacol*. 2009; 23:449–455.
39. Zhang X, Song Y, Ci X, et al. Ivermectin inhibits LPS-induced production of inflammatory cytokines and improves LPS-induced survival in mice. *Inflamm Res*. 2008;57:524–529.
40. Héctor C, Roberto H, Psaltis A, et al. Study of the efficacy and safety of topical ivermectin+ iota-carrageenan in the prophylaxis against COVID-19 in health personnel. *J Biomed Res Clin Investig*. 2020;2.
41. Behera P, Patro BK, Singh AK, et al. Role of ivermectin in the prevention of COVID-19 infection among healthcare workers in India: a matched case-control study. *medRxiv*. 2020.
42. Hellwig MD, Maia A. A COVID-19 Prophylaxis? Lower incidence associated with prophylactic administration of Ivermectin. *Int J Antimicrob Agents*. 2021;57:106248.
43. Bernigaud C, Guillemot D, Ahmed-Belkacem A. *et al. Bénéfice de l'ivermectine: de la gale à la COVID-19, un exemple de sérendipité.*. *Annales de Dermatologie et de Vénérologie*. 2020;147:A194.
44. Shouman WM, Hegazy AA, Nafae RM, et al. Use of Ivermectin as a Potential Chemoprophylaxis for COVID-19 in Egypt: A Randomized Clinical Trial. *J Clin Diagn Res*. 2021;15.
45. Elgazzar A, Hany B, Youssef SA, Hafez M, Moussa H, eltaweel A. Efficacy and safety of ivermectin for treatment and prophylaxis of COVID-19 pandemic. *Res Square*. 2020. Doi: 10.21203/rs.3.rs-100956/v3.
46. Prophylaxis Chala. Covid-19 in healthcare agents by intensive treatment with ivermectin and iota-carrageenan (Ivercar-Tuc). *ClinicalTrials.gov*. 2020: NCT04701710.
47. Alam M, RM, PFG, et al. Ivermectin as pre-exposure prophylaxis for COVID 19 among healthcare providers in a selected tertiary hospital in Dhaka an observational study. *Eur J Med Health Sci*. 2020;2:1–5.
48. Chamie J. Real-world evidence: the case of Peru. In: *Causality between Ivermectin and COVID-19 Infection Fatality Rate*; 2020.
49. Mahmud R. A Randomized, Double-Blind Placebo Controlled Clinical Trial of Ivermectin Plus Doxycycline for the Treatment of Confirmed Covid-19 Infection *ClinicalTrials.gov*. 2020. Available at: <https://clinicaltrials.gov/ct2/show/results/NCT04523831>.
50. Carvallo HE, Hirsch RR, Farinella ME. Safety and Efficacy of the combined use of ivermectin, dexamethasone, enoxaparin and aspirin against COVID-19. *medRxiv*. 2020. Available at: <https://www.medrxiv.org/content/10.1101/2020.09.10.20191619v1>.
51. Khan MSI, Khan MSI, Debnath CR, et al. Ivermectin treatment may improve the prognosis of patients with COVID-19. *Archivos de Bronconeumología*. 2020;56:828–830.
52. Gorial FI, Mashhadani S, Sayaly HM, et al. Effectiveness of ivermectin as add-on therapy in COVID-19 management (pilot trial). *medRxiv*. 2020. doi: 10.1101/2020.07.07.20145979.
53. Hashim HA, Maulood MF, Rasheed AM, et al. Controlled randomized clinical trial on using Ivermectin with Doxycycline for treating COVID-19 patients in Baghdad, Iraq. *medRxiv*. 2020. doi: 10.1101/2020.10.26.20219345.
54. Chowdhury AT, Shahbaz M, Karim MR, et al. A randomized trial of ivermectin-doxycycline and hydroxychloroquine-azithromycin therapy on COVID19 patients. *Res Square*. 2020. doi: 10.21203/rs.3.rs-38896/v1.
55. Podder CS, Chowdhury N, Sina MI, et al. Outcome of ivermectin treated mild to moderate COVID-19 cases: a single-centre, open-label, randomised controlled study. *IMC J Med Sci*. 2020;14:1–8.
56. Cadebiani FA, Goren A, Wambier CG, et al. Early COVID-19 therapy with azithromycin plus nitazoxanide, ivermectin or hydroxychloroquine in outpatient settings significantly reduced symptoms compared to known outcomes in untreated patients. *medRxiv*. 2020. doi: 10.1101/2020.10.31.20223883.
57. Chaccour C, Casellas A, Blanco-Di Matteo A, et al. The effect of early treatment with ivermectin on viral load, symptoms and humoral response in patients with non-severe COVID-19: A pilot, double-blind, placebo-controlled, randomized clinical trial. *EClinicalMedicine*. 2021;32:100720.
58. Espitia-Hernandez G, Munguia L, Diaz-Chiguer D, et al. *Effects of Ivermectin-Azithromycin-Cholecalciferol Combined Therapy on COVID-19 Infected Patients: A Proof of Concept Study*; 2020.

59. Babalola OE, Bode CO, Ajayi AA, et al. Ivermectin shows clinical benefits in mild to moderate Covid19 disease: a randomised controlled double blind dose response study in Lagos. *medRxiv*. 2021;hcab035. doi: 10.1093/qjmed/hcab035.
60. Ravikirti, Roy R, Pattadar C, et al. Ivermectin as a potential treatment for mild to moderate COVID-19 – a double blind randomized placebo-controlled trial. *medRxiv*. 2021. doi: 10.1101/2021.01.05.21249310.
61. Morgenstern J, Redondo JN, De Leon A, et al. The use of compassionate Ivermectin in the management of symptomatic outpatients and hospitalized patients with clinical diagnosis of COVID-19 at the Medical Center Bournigal and the Medical Center Punta Cana, Rescue Group, Dominican Republic, from may 1 to august 10, 2020. *medRxiv*. 2020. Doi: 10.1101/2020.10.29.20222505.
62. Robin RC, Alam RF, Saber S, et al. A case series of 100 COVID-19 positive patients treated with combination of ivermectin and doxycycline. *J Bangladesh Coll Physicians Surgeons*. 2020;38:10–15. doi: 10.3329/jbcps.v38i0.47512. [published Online First: Epub Date].
63. Rajter JC, Sherman MS, Fatteh N, et al. Use of ivermectin is associated with lower mortality in hospitalized patients with COVID-19 (ICON study). *Chest*. 2021;159:85–92.
64. Spoorthi VSS. Utility of ivermectin and doxycycline combination for the treatment of SARS-CoV2. *Int Arch Integrated Med*. 2020;7:177–182.
65. Niaee MS, Gheibi N, Namdar P, et al. Ivermectin as an adjunct treatment for hospitalized adult COVID-19 patients: A randomized multi-center clinical trial. *Res Square*. 2020. doi: 10.21203/rs.3.rs-109670/v1.
66. Portmann-Baracco A, Bryce-Alberti M, Accinelli RA. Antiviral and anti-inflammatory properties of ivermectin and its potential use in covid-19. *Arch Bronconeumol*. 2020;56:831.
67. Ahmed S, Karim MM, Ross AG, et al. A five day course of ivermectin for the treatment of COVID-19 may reduce the duration of illness. *Int J Infect Dis*. 2021;103:214–216.
68. Chachar AZK, Khan KA, Asif M, et al. Effectiveness of ivermectin in SARS-CoV-2/COVID-19 patients. *Int J Sci*. 2020;9:31–35.
69. Soto-Becerra P, Culquichicón C, Hurtado-Roca Y, et al. Real-world effectiveness of hydroxychloroquine, azithromycin, and ivermectin among hospitalized COVID-19 patients: results of a target trial emulation using observational data from a nationwide healthcare system in Peru. *SSRN*. 2020. Available at: <https://ssrn.com/abstract=3710623>.
70. Budhiraja S, Soni A, Jha V, et al. Clinical Profile of First 1000 COVID-19 cases admitted at tertiary care hospitals and the correlates of their mortality: an Indian experience. *medRxiv*. 2020. doi: 10.1101/2020.11.16.20232223.
71. Rubin R. As their numbers grow, COVID-19 “long haulers” stump experts. *JAMA*. 2020;324:1381–1383.
72. Callard F, Perego E. How and why patients made Long Covid. *Soc Sci Med*. 2021;268:113426.
73. Siegelman JN. Reflections of a COVID-19 long hauler. *JAMA*. 2020;324:2031–2032.
74. Aguirre-Chang G. Post-Acute or prolonged COVID-19: treatment with ivermectin for patients with persistent, or post-acute symptoms *ResearchGate*. 2020. Available at: https://www.researchgate.net/publication/344318845_POST-ACUTE_OR_PROLONGED_COVID-19_IVERMECTIN_TREATMENT_FOR_PATIENTS_WITH_PERSISTENT_SYMPTOMS_OR_POST-ACUTE.
75. Kircik LH, Del Rosso JQ, Layton AM, et al. Over 25 Years of clinical experience with ivermectin: an overview of safety for an increasing number of indications. *J Drugs Dermatol*. 2016;15:325–332.
76. Gardon J, Gardon-Wendel N, Demanga N, et al. Serious reactions after mass treatment of onchocerciasis with ivermectin in an area endemic for Loa loa infection. *Lancet*. 1997;350:18–22.
77. Guzzo CA, Furtek CI, Porras AG, et al. Safety, tolerability, and pharmacokinetics of escalating high doses of ivermectin in healthy adult subjects. *J Clin Pharmacol*. 2002;42:1122–1133.
78. Veit O, Beck B, Steuerwald M, et al. First case of ivermectin-induced severe hepatitis. *Trans R Soc Trop Med Hyg*. 2006;100:795–797.
79. Sparsa A, Bonnetblanc JM, Peyrot I, et al. Systemic adverse reactions with ivermectin treatment of scabies. *Annales de Dermatologie et de Venereologie*. 2006;133:784–787.
80. Chandler RE. Serious neurological adverse events after ivermectin—do they occur beyond the indication of onchocerciasis?. *Am J Trop Med Hyg*. 2018;98:382–388.
81. Anglemeyer A, Horvath HT, Bero L. Healthcare outcomes assessed with observational study designs compared with those assessed in randomized trials. *Cochrane Database Syst Rev*. 2014;MR000034.
82. Dahabreh IJ, Sheldrick RC, Paulus JK, et al. Do observational studies using propensity score methods agree with randomized trials? A systematic comparison of studies on acute coronary syndromes. *Eur Heart J*. 2012;33:1893–1901.
83. Kitsios GD, Dahabreh IJ, Callahan S, et al. Can we trust observational studies using propensity scores in the critical care literature? A systematic comparison with randomized clinical trials. *Crit Care Med*. 2015;43:1870–1879.
84. Lonjon G, Boutron I, Trinquart L, et al. Comparison of treatment effect estimates from prospective nonrandomized studies with propensity score analysis and randomized controlled trials of surgical procedures. *Ann Surg*. 2014;259:18–25.