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A secondary approach with conventional medicines and supplements to recuperate current COVID-19 status

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ABSTRACT

Novel coronavirus 2019 (COVID-19) is a zoonosis that revised the global economic and societal progress since early 2020. The SARS-CoV-2 has been recognized as the responsible pathogen for COVID-19 with high infection and mortality rate potential. It has spread in 192 countries and infected about 1.5% of the world population, and still, a proper therapeutic approach is not unveiled. COVID-19 indication starts with fever to shortness of breathing, leading to ICU admission with the ventilation support in severe conditions. Besides the symptomatic mainstay clinical therapeutic approach, only Remdesivir has been approved by the FDA. Several pharmaceutical companies claimed different vaccines with exceptionally high efficacy (90–95%) against COVID-19; how long these vaccines can protect and long-term safety with the new variants are unpredictable. After the worldwide spread of the COVID-19 pandemic, numerous clinical trials with different phases are being performed to find the most appropriate solution to this condition. Some of these trials with old FDA-approved drugs showed promising results. In this review, we have precisely compiled the efforts to curb the disease and discussed the clinical findings of Ivermectin, Doxycycline, Vitamin-D, Vitamin-C, Zinc, and cannabidiol and their combinations. Additionally, the correlation of these molecules on the prophylactic and diseased administration against COVID-19 has been explored.

1. Introduction

Coronaviruses (CoVs) are single-stranded RNA viruses that consist of spike glycoproteins on the envelope, which look like a crown (*coronam* as Latin term of the crown) [1]. CoVs are subdivided into four groups: Alpha-coronavirus (α -CoV), Beta-coronavirus (β -CoV), Delta-coronavirus (Δ -CoV), and Gamma-coronavirus (γ -CoV). Severe-Acute-Respiratory-Syndrome CoV-2 (SARS-CoV-2) is belonging to the β -CoV [2]. SARS-CoV-2 genome has 96% similarity of β -CoV RaTG13 of bats though the origin is still unknown [3]. S2 of (one of the spike proteins) SARS-CoV-2 has an amino acid site that allows it to bind with the Angiotensin-Converting Enzyme 2 (ACE2) receptor and infuse its RNA to the host cells. Host cells replicate this RNA rapidly and produce an excessive immune reaction along with redox imbalance in the host which, finally causes the novel coronavirus disease at the end of 2019, renowned as COVID-19 [4,5]. Fever, dry cough, fatigue, loss of taste or smell, nasal congestion, conjunctivitis (also known as red eyes),

sore throat, headache, muscle or joint pain, different types of skin rash, nausea or vomiting, diarrhea, and chills or dizziness are the symptoms developed in COVID-19 [6].

The most susceptible to the COVID-19 are the persons with low immunity system, elderly aged people with different health concerns like cancer, chronic kidney diseases, chronic obstructive pulmonary diseases, a heart condition, pregnancy, and type 2 diabetics mellitus [7]. Interestingly, there might be a role of an individual's blood group to be at risk of SARS-CoV-2 infection [8].

The current therapeutic approach is mostly symptomatic, but many clinical trials are being performed to figure out preventive management. Lately, Remdesivir got approval from the FDA (Food and Drug Administration) to treat hospitalized COVID-19 patients [9]. Different pharmaceutical companies are trying to develop vaccines employing versatile methodologies. Companies like Pfizer/BioNtech, Moderna claimed their mRNA (messenger RNA) vaccines are 94–95% effective against SARS-CoV-2 [10]. Viral vector vaccines (ChAdOx1 nCoV-19) of

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AstraZeneca and the University of Oxford were studied in 23,848 participants in UK, Brazil, and South Africa. Here, a low dose followed by a standard dose was 90% effective against COVID-19 [11]. Even though, all the vaccines are promising, they are yet to be approved by the WHO for mass usage. Side by side, how long these vaccines can provide protection and their long-term consequences are still unpredictable.

Due to these panic circumstances, peoples from all groups are eager to get information about the viral cause and cure irrespective of their area. Therefore, in this study, we would like to discuss some FDA-approved drugs and supplements studied against COVID-19 separately or as adjuvant. These drugs were used mostly to treat COVID-19, but some of them were used as prophylaxis. Hence, this document will provide comprehensive information about pharmaceutical efforts against COVID-19 in a precise way.

2. Current COVID-19 epidemic status

Since the arrival of novel coronavirus-2019, it has spread around 192 countries and infected over 162 million people (2.1% of the total world population) around the world [12]. At the time of writing (May 2021) according to Johns Hopkins University, the total recovery from COVID-19 is 91.25 million with more than 3.4 million fatalities [13]. According to the WHO as of 18th May 2021, 40% cumulative cases are accumulated in the north and south Americas, 33% in Europe, 17% in South-East Asia, 6% in Eastern Mediterranean, 2% in Africa, and 2% in Western Pacific region [12].

From the 31st January 2020 to 18th May 2021 (17 months) the cases and deaths per month were extracted from the data provided by WHO (Supplementary data) [14]. Those data were plotted in Figs. 1 and 2 where the whole world was divided into six regions (Americas, Europe, South-East Asia, Eastern Mediterranean, Africa, Western Pacific). In Fig. 1A and B, the number of cases and deaths were arranged for all the six regions, and as cases per month were changing in these regions with the months and created a wave shape. The first highest peak was considered as the first wave and the second highest peak as the second wave. The number of cases in the 2nd wave was much greater than the 1st wave (Fig. 1C) which was predicted earlier by Atangana et al. [15]. Cases per month were highest in the Americas but the highest deaths were observed in Europe region although, infection rate and fatality are coming down gradually. But in South-East Asia especially India now reached a critical situation and about to face the most devastating hit by COVID-19.

3. Evolution of COVID-19 around the world for the last 17 months

Though the number of cases and deaths are related to each other the ratio between cases and death is behaving differently in certain regions (Fig. 2). In the Americas region the total case has come down but the death rate is still going up especially in Brazil, U.S.A, Argentina, and Colombia (Fig. 2A). In Europe, death is steady with a fall down in the number of cases per month (Fig. 2B). South-East Asia has now become the burning ground of COVID-19 with the maximum number of cases and death especially in India (Fig. 2C). In Eastern Mediterranean the number of cases has become worst of all time particularly in Iran (Fig. 2D). Interestingly, in Africa, the number of cases has fallen but the death rate is spiking up due to the new range of cases in South Africa (Fig. 2E). The Western Pacific which was meant to be the most controlled place also had its highest peak of cases per month lately especially for Japan, the Philippines, and Malaysia (Fig. 2F).

The most probable reasons for these fluctuations of cases and death rates might be related to the application of lockdown protocol, maintaining social distancing, vaccination, and adequate healthcare facility. However, these data are collected based on regions where different countries followed different approaches. The most of the regions where the condition is worsening, the most common reasons are the saturation

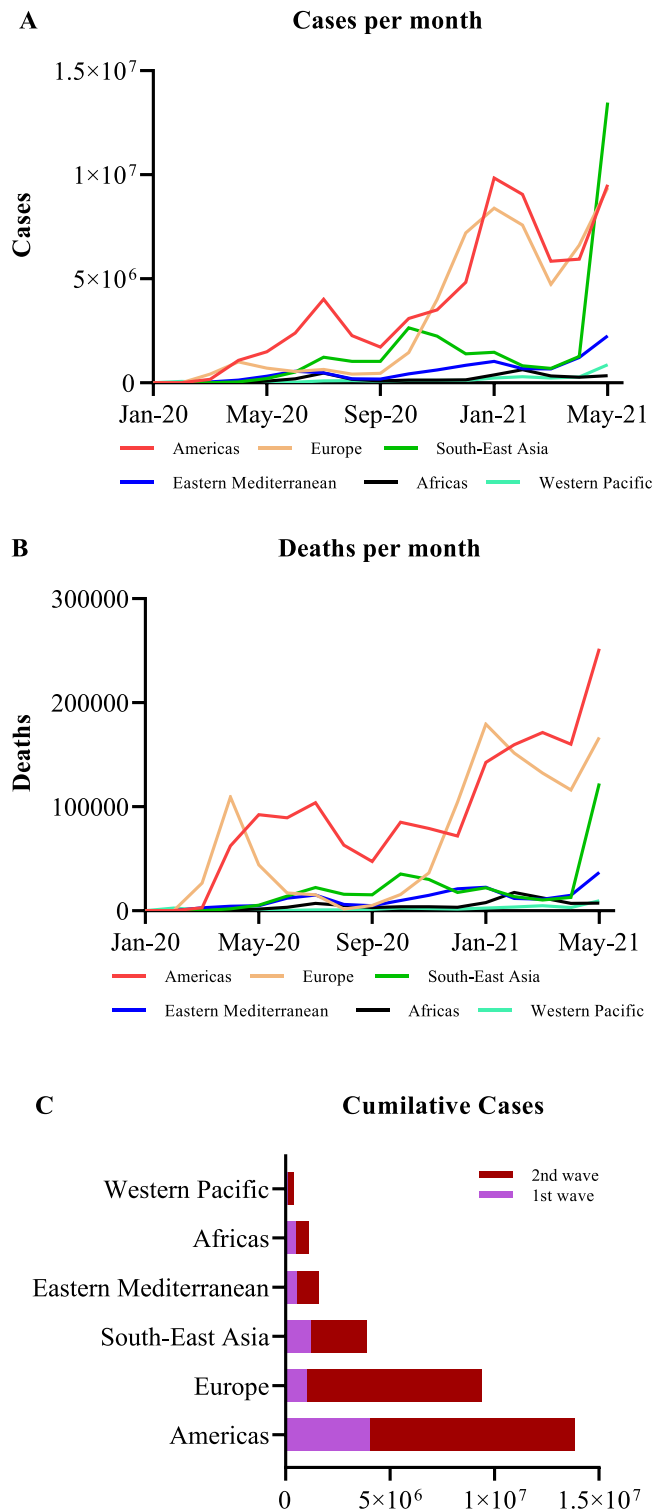


Fig. 1. COVID-19 epidemiological status till 2nd May 2021; these graphs explain the condition of the COVID-19 situation in the world. A. Illustrate the total cases that have been reported in different regions (1 million = 1×10^7). B. States the cumulative deaths in different regions. C. Cumulative cases comparison between the 1st wave and 2nd wave of COVID-19.

of health care scope and vaccination. At present, the death percentage of south-east Asia is raising sharply against the number of cases which might be the influence of new variants and poor pandemic management in this region.

Nevertheless, the mutation in a virus is inevitable and can change its

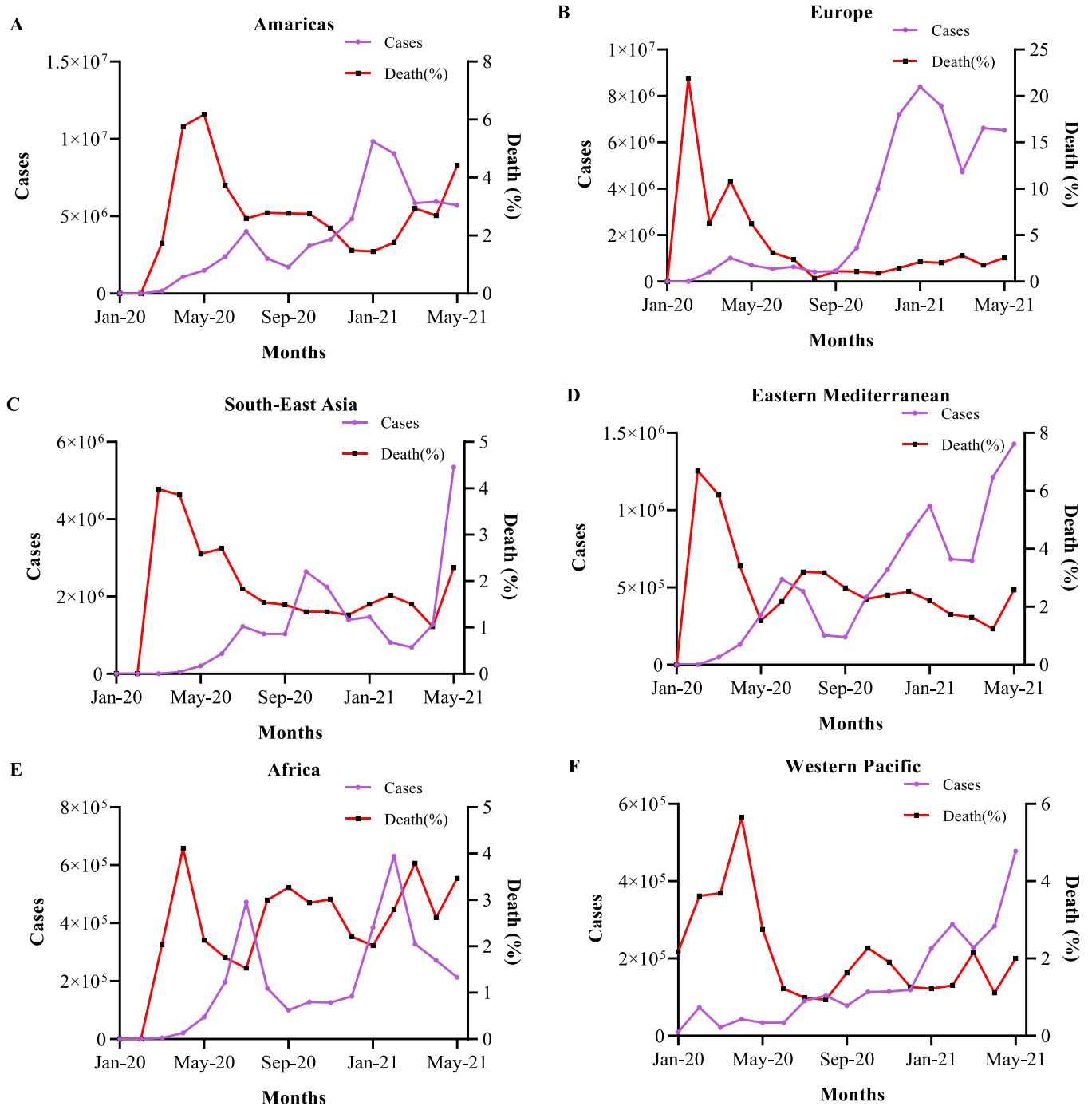


Fig. 2. Comparison of the number of cases and percentage of death per month; all the data are plotted in these graphs from January 2020 to 18th May 2021 (17 months). A, B, C, D, E, and F are cumulative cases vs percentage of death per month in Americas, Europe, South-East Asia, Eastern Mediterranean, Africa, and Western Pacific regions respectively.

contagious profile. SARS-CoV-2 has also mutated as it jumped through lots of individuals. Particular four variants (B.1.1.7, B.1.351, B.1.1.28.1, and B.1.617) of SARS-CoV-2 have raised certain concerns due to their greater transmissibility than the native virus [16]. Around 152 countries have reported with B.1.1.7 variant patients followed by 106 countries with B.1.351, 61 countries with B.1.1.28.1, and 62 countries with new variant B.1.617. Among these four variants B.1.1.7 is the most alarming not only because of the greater risk of mortality and hospitalization but the recent outbreak in India with B.1.617 and its mutated variants (B.1.617.1, B.1.617.2, B.1.617.3, B.1.617.x) has become more serious and also reported in failure with conventional detection. Experts suspect

that this new variant B.1.617 has more capacity for a human to human transmission than the others [17]. Though present vaccine provider companies (Pfizer-BioNTech, Moderna, and Oxford-AstraZeneca) states that their vaccines still active in B.1.1.7 variant but show reduced efficacy on B.1.351 phenotype [18].

4. Recent treatment protocol

SARS-CoV-2 spike protein configuration is different from its ancestor SARS-CoV and MERS-CoV (Middle East respiratory syndrome coronavirus) which makes COVID-19 management complicated. Due to the fast

infection rate, mutation, and complex mechanism of action, conventional anti-viral therapy leads to poor management [19]. Many clinical trials are going on for the search on suitable COVID-19 treatment plans.

On February 9, 2021, FDA had issued Emergency use authorization for monoclonal antibody (bamlanivimab plus etesevimab) treatment for the outpatient with mild to moderate symptoms but not for the hospitalized patients or those who required oxygen therapy [20]. Remdesivir or Remdesivir plus Dexamethasone or Dexamethasone treatment is recommended for those who are hospitalized and require supplemental oxygen. Only Dexamethasone for the hospitalized patient with invasive mechanical ventilation [21].

5. An alternative approach to COVID-19 management

On September 24th of 2020, my cousin and her whole family (5 members) got infected with COVID-19 and they followed the prescription which follows Ivermectin (12 mg) for 5 days, Doxycycline (200 mg) for 7 days, Vitamin-D (2000IU) 15 days, Vitamin C (chewable table) 1 month. They were quarantined and had these medications according to the prescription. In the following 21 days, they all were recovered. This real event intrigued me to investigate more about these drugs against the COVID-19. Another most important reason which interests me that these drugs are available in every country and their safety profile is outstanding. Side by side, countries with lower economic growth have little reachability to the vaccine which made them even more vulnerable to COVID-19. Scientists and clinical practitioners are more driven in the search for alternative management because of poor outcomes from the recent treatment protocol. We have discussed below six potential candidates (Fig. 3) for better management based on several clinical trials (Table 2). These drugs might not only approachable for COVID-19 treatment but also might protect if taken in a prophylactic manner.

6. Ivermectin

Ivermectin (IMT) is a macrocyclic lactone derived from the Nobel

prize-winning compound avermectin [22]. IMT is a broad-spectrum anti-parasitic drug (FDA approved) [23] and also proven as an anti-viral drug in a wide range [24–26], anti-cancer [27], topical anti-inflammatory and wound healing [28,29], antibacterial [30], anti-mitotic activity [31].

Nonstructural protein 5 (NS5) is a crucial component to replicate RNA viruses like ZIKA, Dengue, West Nile, and HIV-1 [32,33]. NS5 complies with the importin β 1 nuclear localization signal (NLS) and α/β NLS [34]. Various investigations proved that IMT could inhibit NLS by disassociating importin α/β 1 heterodimer and thus blind NS5 to recognize importin mediated NLS [32]. Coronavirus is also a positive-sense single-stranded RNA virus, making it a suitable target for IMT treatment [19]. Clay et al. was the first to investigate the antiviral effect of IMT on Covid-19 in vitro and reported results elucidated an efficient remedy with \sim 5000-fold reduction of viral RNA in 48 h [35]. IMT also can downregulate the inflammation through nuclear factor kappa-light-chain-enhancer of activated B cells (NF- κ B) and MAPK pathway [36]. Recently five days randomized clinical trial was conducted on 72 patients infected with mild Covid-19. Patients were divided into three groups (IMT, IMT + doxycycline, and placebo) and received 12 mg of IMT or IMT 12 mg and doxycycline (DC) 200 mg once a day for five days except for the control group. Interestingly the viral load was found negative in 9.7 days and fully recovered within 14 days. No adverse effect or drug-drug interaction was noticed during this trial [37]. Additionally, another phase-3 clinical trial (NCT04523831) was conducted on 400 patients with mild symptoms by combining IMT and DC. In this study, patients have received IMT 6 mg twice on the first day of the treatment and discontinued for the rest of the course. DC 100 mg twice a day (1 + 0 + 1) was continued for five days. Within seven days, Body temperature and oxygen saturation become normal (37.5°C and $\text{SpO}_2 > 93\%$) for 60.7% of the treatment group, 12 days for 23%, 14 days for 7.7%, and one month for 8.7%. Interestingly the mortality rate was 0.0%, which was very encouraging. Another small trial (NCT04591600) on IMT also suggested the same indication, but they also included that early administration of IMT can lead to better

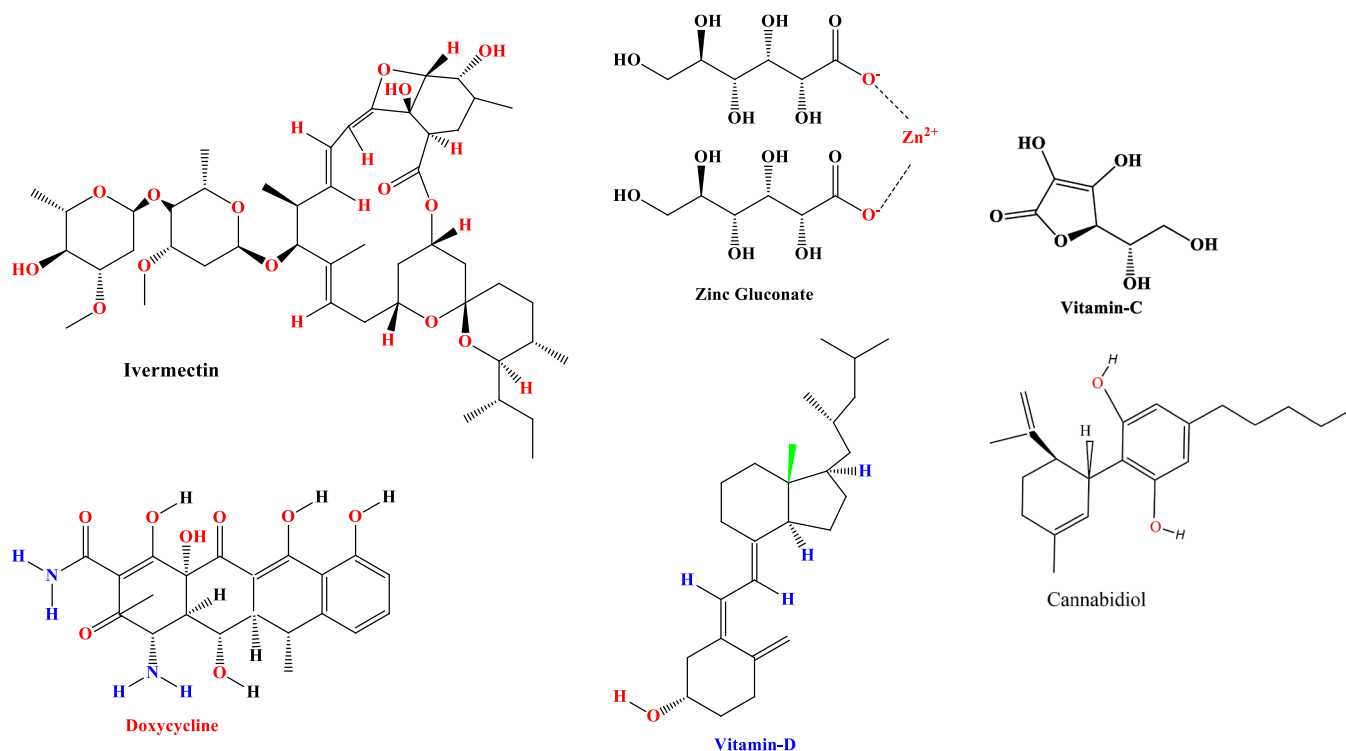


Fig. 3. Potential candidate for COVID-19 management; this diagram displays six promising conventional molecules (2D structure) that might present a life savior option to treat COVID-19.

Table 1

Alternative option of COVID-19 management; the list of drugs and supplements with their dose and regime to use either as preventive measure or treatment approach against COVID-19.

Generic name	Dose	Dose regimen (prophylaxis)	Dose regimen (treatment)	Availability	Role on COVID-19
Ivermectin	12 mg/day	Once	Physician's decision	On prescription	Inhibit viral replication, Immunomodulation.
Doxycycline	40–200 mg/day	5 days	Physician's decision	On prescription	Inhibit viral replication, anti-inflammation.
Zinc	15–50 mg/day	20–30 days	Physician's decision	OTC	Inhibit viral activity.
Vitamin-D	1000–4000 IU/day	20–30 days	Physician's decision	OTC	Increase immunity.
Vitamin-C	250–1000 mg/day	20–30 days	Physician's decision	OTC	Reduce viral infection chance, redox balance.
Cannabidiol	Physician's decision	Physician's decision	Physician's decision	On prescription	Incorporate viral invasion, reduce ARDS

management [38].

7. Doxycycline

Doxycycline (DC) is an FDA-approved antibiotic that comes from the tetracycline family. DC possesses good intra-cellular penetration and broad-spectrum antibacterial effect [39]. DC has been reported for various infectious disease control including pasteurellosis, brucellosis, borreliosis, trepanomatosis, rickettsioses, leptospirosis, cholera, Q fever, urinary and pulmonary infections by Chlamydia and Mycoplasma, anthrax, and gonococci. It is also useful for malaria prophylaxis because of its anti-Plasmodium capabilities [40]. In dermatology, DC is indicated for rosacea and acne [41]. DC can bind with bacterial 30S ribosomal subunit and inhibit bacterial protein synthesis [42]. Besides its core effect, DC is also studied for its antiviral activity against the chikungunya virus, Japanese encephalitis virus, Respiratory Syncytial Virus, and Dengue virus [43–46]. Rothan et al. showed that DC is very potent to inhibit RNA viruses, e.g., chikungunya [44]. DC can inhibit matrix metalloproteinases and bind with E2 envelope glycoprotein, which is important for viral replication and survival [47]. Investigation results elucidated that DC can have a potential effect on SARS-CoV-2 entry and its post replication. Side by side, it can act as an anti-inflammatory agent by reducing various pro-inflammatory cytokines expressions [48]. In May 2020, a small study was conducted on 89 COVID-19 positive patients with DC (100 mg) for seven days. Initially, they were treated with hydroxychloroquine and azithromycin, but afterward, medication was replaced with DC because of both of their cardiotoxic behavior. This study reported a recovery rate of 85%, and only 3% of patients were rebounded for hospitalization due to their clinical deuteriation after ten days [49]. As previously described, the IMT + DC combination was remarkable on COVID-19 recovery (NCT04523831, NCT04591600).

8. Zinc

Zinc (Zn) is an essential macronutrient that involves growth development, neurosensory functions, DNA synthesis, gene expression, and immune system regulation. Nowadays, 17.3% of the global population is Zn deficient, and Zn deficiency can lead to growth retardation, diarrhea, hair loss (alopecia), inflammation of the tongue (glossitis), nail dystrophy, and premature birth [50]. As Zn's source, Oysters comprise the most, but the major Zn comes from red meat and poultry. Refined rice only contains 20% Zn of daily requirement although, the countries where rice is the staple food has more susceptible to Zn deficiency [51, 52]. Insufficient Zn can downregulate antibody production and, consequently, reduce cell-mediated immunity, leading to more infection vulnerability [53]. Admission of Zn can reduce the duration of common cold symptoms [54] and upper respiratory tract infection morbidity [55]. It also can inhibit the hepatitis C virus through IFN- λ 3 signaling [56] and ease people living with HIV infections [57]. *In vitro* study showed that Zn cations could inhibit SARS-CoV-2 RNA polymerase activity by decreasing replication [58]. Zn with or without chloroquine can increase Zn abundance inside the cells, which can intensify cell-mediated immunity [59]. SARS-CoV-2 penetrates the cells through angiotensin-converting enzyme 2 (ACE2), and Zn was studied on the modulation of ACE2. Exposure of Zn reduced recombinant human ACE-2

activity in rat lungs, which might be a potential approach against COVID-19 [60]. Five hundred twenty patients with COVID-19 were recruited for a clinical trial (NCT04342728) with ascorbic acid (8000 mg/day) and Zn (50 mg/day) for 28 days. This group was divided into four subgroups (Ascorbic Acid or Zinc Gluconate or Ascorbic Acid and Zinc Gluconate Dietary Supplement and Standard of Care). This study was expected to reduce symptoms in 28 days, though the results are yet to develop. A case study of four patients who received a high dose of Zn improved significantly after one day of treatment, and the researcher suggested that Zn might play a crucial part in the recovery phase from COVID-19 [61].

9. Vitamin D

Vitamin D (Vit-D) is one of the essential vitamins that can be involved in calcium homeostasis and involves the immune system modulation of our body. Lack of Vit-D can lead to autoimmune diseases and increase the vulnerability to infections [62]. Vit-D deficiency (serum 25-hydroxyvitamin D < 20 ng/ml or 50 nmol/L) is very common worldwide. Currently, 40% in Europe, 37% in Canada, 24% in the U.S., > 20% population in India, Tunisia, Pakistan, and Afghanistan are reported as Vit-D deficient [63]. Generally, adults [64], darker skin [65], and obese [66] people are more susceptible to Vit-D lacking. Elizabeth et al. recorded 10,926 death from 17,278,392 COVID-19 patients and found that more elderly, obese, and darker skin people died [67], which might have a co-relation with Vit-D deficiency. Another study showed that COVID-19 mortality and infection are inversely proportional to the population's Vit-D level [68]. A blood clot in the lung tissue and ventilation is more prevalent to lower Vit-D leveled people and has a greater mortality rate [69,70].

Vit-D can be obtained by sun exposure, diet, and dietary supplement; however, direct sunlight to the skin is the primary source of Vit-D [71]. A small portion can be obtained from fish, egg yolk, animal liver, and milk [72]. The daily necessary intake of Vit-D for an adult is 15 μ g (600 IU) to 100 μ g (4000 IU) [73], but due to lifestyle and lack of sunlight, 50% of the world population is at risk of Vit-D deficiency [74]. A clinical trial (NCT04449718) of 240 patients displayed that one single bolus dose of Vit-D did not improve the outcome. Another small study showed that a high dose (60,000 IU/day) for seven days might help against COVID-19 and a significant decrease in fibrinogen [75].

10. Vitamin C

Vitamin C (Vit-C) or ascorbic acid is a powerful antioxidant that helps decrease inflammation and improve immune function. Vit-C deficiency or low dietary intake can lead to low immunity and greater susceptibility to viral infection [76]. Vit-C has well documented against the common cold including various studies against herpes simplex virus-1, influenza type A virus, AIDS, poliovirus type 1, viral upper respiratory tract infections, acute respiratory infections, and rhinovirus have proved this [77]. In a viral infection, redox homeostasis is imbalanced and causes tissue damage, leading to inflammation [78]. Studies showed that Vit-C could shorten the length of ICU (intensive care unit) with respiratory diseases [79] and I.V. (intravenous) dose helps patients to recover from septic shock [80]. In ICU, Vit-C can improve blood

Table 2

List of clinical trials and case studies. The clinical studies related to the IMT, DC, Zn, Vit-C, Vit-D, and CBD are listed in this table with their clinical trial identification number and actual or estimated participants amount. The studies which don't have identification numbers are referred to a case studies.

Title	Number of patients enrolled	Identification number
A five-day course of ivermectin for the treatment of COVID-19 may reduce the duration of illness.	72	none
Clinical Trial of Ivermectin Plus Doxycycline for the Treatment of Confirmed Covid-19 Infection.	400	NCT04523831
Effectiveness of Ivermectin and Doxycycline on COVID-19 Patients.	140	NCT04591600
Clinical Outcomes of Early Treatment with Doxycycline for 89 High-Risk COVID-19 Patients in Long-Term Care Facilities in New York.	89	none
Coronavirus 2019 (COVID-19)- Using Ascorbic Acid and Zinc Supplementation (COVIDAtoZ).	520	NCT04342728
Vitamin D Supplementation in Patients With COVID-19.	240	NCT04449718
Short Term, High Dose Vitamin D Supplementation for COVID-19 (SHADE).	30	NCT04459247
Prophylactic Ivermectin in COVID-19 Contacts.	340	NCT04422561
Role of ivermectin in the prevention of SARS-CoV-2 infection among healthcare workers in India: A matched case-control study.	117	none
Zinc Versus Multivitamin Micronutrient Supplementation in the Setting of COVID-19 (ZnCOVID-19).	4500 (estimated)	NCT04551339
A Preventive Treatment for Migrant Workers at High-risk of COVID-19.	4257	NCT04446104
A Study of Hydroxychloroquine, Vitamin C, Vitamin D, and Zinc for the Prevention of COVID-19 Infection (HELPCOVID-19).	600 (estimated)	NCT04335084
Safety and Efficacy of Hydroxychloroquine for the Treatment & Prevention of Coronavirus Disease 2019 (COVID-19) Caused by Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2)	5000 (estimated)	NCT04590274
Povidone-Iodine Vs Essential Oil Vs Tap Water Gargling For COVID-19 Patients (GARGLES).	20	NCT04410159
Study to Evaluate the Safety and Antiviral Activity of Remdesivir (GS-5734™) in Participants With Moderate Coronavirus Disease (COVID-19) Compared to Standard of Care Treatment.	596	NCT04292730
Study to Evaluate the Safety and Antiviral Activity of Remdesivir (GS-5734™) in Participants With Severe Coronavirus Disease (COVID-19)	397	NCT04292899
Outcomes Mandate National Integration with Cannabis as Medicine for Prevention and Treatment of COVID-19 (OMNI-Can)	200,000 (estimated)	NCT03944447
Cannabidiol for Covid-19 patients With Mild to Moderate Symptoms (CANDIDATE)	104 (estimated)	NCT04467918

pressure, bronchoconstriction, infections, atrial fibrillation, and acute renal injury [81]. So, a mega-dose of Vit-C might rescue severely ill patients from COVID-19 [82]. During the pandemic in China, 10 g and 20 g Vit-C I.V. was prescribed from moderate to severe COVID-19 patients for 5–7 days and that improved the disease resolution period [77]. Though regular 250 mg (twice daily) Vit-C supplement might present a preventive measure to the general population [83].

Besides Vit-C, quercetin (QCT) is a natural plant-derived flavonoid, can act as a viral protease inhibitor thus block RNA production and viral replication of different RNA viruses like HIV, Hepatitis-C, and SARS-CoV-1. Concurrent administration of Vit-C, QCT, and Vit-B3 can

reduce the vulnerability and mortality from influenza H1N1. QCT is also believed to bind with the 3CL-like-protease of SARS-CoV-2 and inhibit its proteolytic activity. Ascorbate (conjugate base form of Vit-C) able to recycle QCT and increase its bioavailability [84,85].

11. Cannabidiol

Cannabidiol (CBD) is a non-psychoactive cannabinoid derived from the cannabis plant. CBD exerts a broad range of therapeutic effects like anti-inflammatory, antioxidant, sedative, anticonvulsive, neuro-protective, hypnotic, antipsychotic, and anti-cancer activities. Recently CBD has been hypothesized against COVID-19 [86]. SARS-CoV-2 binds with ACE2 receptor followed by the consequent activation of NF-κB and stormy release of interleukin-6 (IL-6), tumor necrosis factor-α, IL-1α/β, chemokine ligand-2, and C-X-C motif chemokine ligand-10. This event leads to activation of T-cells and reduction of total T-cells amount leads to disrupt immunity system [87]. Wang et al. illustrated that CBD might able to manifest a possible hindrance at the infiltration gateway by reducing the expression of ACE2 and transmembrane serine protease-2 (TMPRSS2) which helps the spike protein to bind with the ACE2 receptor [88]. Another study suggested that CBD can modulate apelin (a secondary catalytic substrate for ACE2) and might improve COVID-19 related acute respiratory distress syndrome (ARDS) [89]. A phase 2 clinical study (NCT03944447) is undergoing to evaluate the application of CBD as a preventive and diseased management tactic. Another small randomized, double-blinded phase 2 study (NCT04467918) is going on with CBD (300 mg/day) for 14 days. This study aims to observe changes in proinflammatory cytokine concentration, parenchymal lung damage, and adverse event with CBD treatment.

12. Prophylactic use of the proposed drugs

A randomized clinical trial (NCT04422561) was done with 340 participants in Egypt to evaluate the prophylactic use of IMT. In this study, they reported only 15 participants out of 203 developed symptoms who receive prophylactic IMT after 14 days. Similarly, a group of health professionals in India received IMT at a dose of 300 µg/kg for 72 h. These healthcare workers were directly in contact with COVID-19 patients, and 73% of these people were uninfected for the following month [90].

DC showed promising results in preventing viral infection, replication, and associated inflammation. This drug has also been used in several clinical trials against COVID-19, suggesting that a low dose of DC can be given (40 mg/day) to take preventive measures [91].

At the time of writing this article, a clinical study (NCT04551339) of 4500 patients is being conducted where the researcher proposed 11 mg of Zn tablet per day for the next three months. This study will be conducted to observe the prophylaxis of Zn. Similar kinds of clinical trials (NCT04446104, NCT04335084, NCT04590274) are also ongoing, but they also see how Zn is standing out hydroxychloroquine, DC, IMT, Vit-C, Vit-D, Azithromycin, and Quercetin.

A study based on the Israeli population was done on 7807 individuals with serum Vit-D test reports and found that only 10.1% were COVID-19 positive, and the rest were not infected. The Vit-D level in the infected group was lower than the non-infected group, which suggests that the Vit-D can boost the immune system to fight against COVID-19 and prevent from getting infected [92].

Different studies against different viral respiratory tract infections showed that Vit-C could decrease the severity of illness and mortality rate [93,94]. Several randomized clinical trials suggested that early intake of Vit-C can reduce the 50% chance of getting caught with the common cold [95].

ACE2 and TMPRSS2 inhibitors can restrict SARS-CoV-2 invasion [96] so, high-cannabidiol containing mouth wash can be a preventive strategy due to the ability of ACE2 downregulation in EpiIntestinal and EpiAirway tissues [86,88].

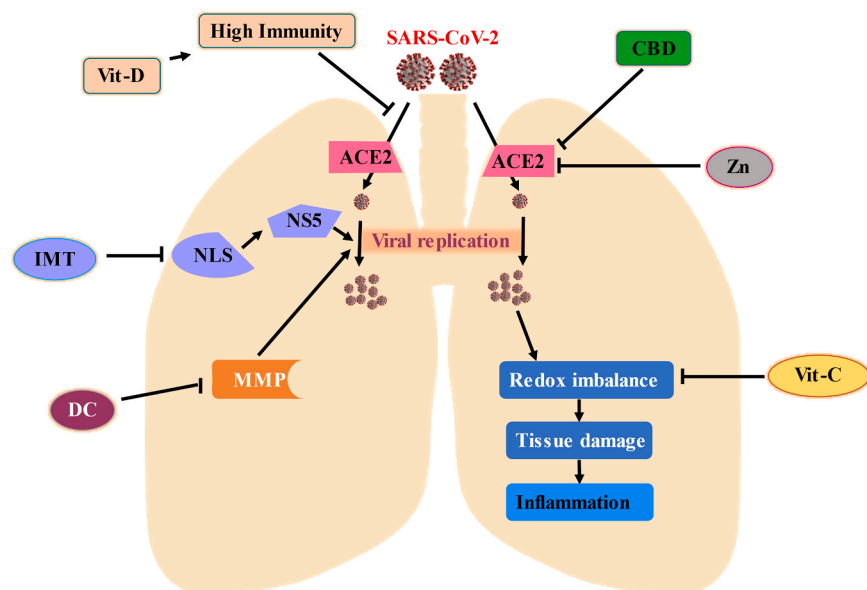


Fig. 4. Proposed mechanism of action against SARS-CoV-2 infection; this illustration reflects a probable mechanism of action with the combination of the Ivermectin (IMT), Doxycycline (DC), Zinc (Zn), Vitamin-D (Vit-D), Vitamin-C (Vit-C), and Cannabidiol (CBD) might present preventive measure against COVID-19. SARS-CoV-2 invades the host body through Angiotensin-Converting Enzyme 2 (ACE2) and consumes host machinery to replicate itself which leads to extensive tissue damage, immune reaction, redox imbalance, and inflammation. IMT induces conformational changes in nuclear localization signal (NLS) and inhibits the contribution of nonstructural protein-5 (NS5) to replicate SARS-CoV-2 along with the immunomodulation to the host. Matrix metalloproteinase (MMP) is also a vital component for viral replication which can be repressed by DC. Vit-C is a well-known antioxidant that can help to counterbalance redox stress. Zn and CBD might modulate ACE2 which encumber SARS-CoV-2 to penetrate the host body. Vit-D can boost the immunity system and might give an extra layer of protection.

13. Safety profile

Though IMT is a very potent anti-parasites, its antiviral capability becomes more prominent in COVID-19 treatment. IMT has a good safety profile and has a minimum adverse effect when taken orally. The safe therapeutic doses of IMT range from 20 to 80 ng/ml in the blood [97].

Like other tetracyclines, DC is also contraindicated in pregnant women, especially after the second trimester, breastfeeding mothers and children. The overuse of DC can also lead to antimicrobial resistance, leading to higher medical costs, prolonged hospital stays, and increased mortality [98]. However, DC is a well-tolerated drug with some side effects like stomach upset and some cutaneous abnormalities [39].

Daily recommended Zn intake for men and women is 7.5–12.7 mg/day and 9.4–16.3 mg/day, respectively. The upper limit is set at 40 mg/day, but higher and prolonged-term intake might cause Zn toxicity, leading to nausea, vomiting, fatigue, epigastric pain, lethargy, and permanent loss of smell [99].

Vit-D has been considered a very safe medication, even in a very high dose. If the serum 25-hydroxyvitamin D has to exceed 150 ng/ml (375 nM/L) with higher calcium intake, Vit-D toxicity might occur, which is extremely rare [100]. According to the European food safety authority (EFSA), 250 µg/day was established as no observed adverse effect level (NOAEL) [101].

Though the daily recommended dose of Vit-C for adults is 90 mg/day during acute infection, this can be raised to 2 g. Does higher than 2 g per day might cause diarrhea, nausea, and abdominal pain [102]. A larger dose (10 mg/day) might cause oxalate kidney stones, but nothing was observed in sepsis in a clinical trial [103].

CBD is non-psychoactive but the smoking of cannabis is not recommended. CBD might lead to hepatic abnormalities, vomiting, diarrhea, fatigue, and CBD-induced drug-drug interactions [104]. So, medical CBD has to be applied under strict monitoring.

14. Comparison between Remdesivir and proposed drug combination

At present, Remdesivir (RDV) is the only drug that has been approved by the FDA for COVID-19 management. The antiviral activity of this drug acts by inhibiting viral replication. The active metabolite of RDV (Remdesivir triphosphate or, GS-443902) is a synthetic analog and competes with the natural nucleosides which inhibit viral RNA replication [105]. RDV nucleoside analog inhibits or delays transportation of

RNA-dependent RNA polymerase of SARS-CoV-2 and suppresses its viral replication [106]. Two clinical trials (NCT04292730 and NCT04292899) were conducted with RDV for moderate to severe COVID-19 patients where they reported that RDV is more suitable for moderate patients when followed five days regime [107,108]. Similar results were also shown by IMT and DC clinical studies. Eweas et al. exhibited that both IMT and RVD were very potent candidates against SARS-CoV-2 [109].

Our proposed drugs act on SARS-CoV-2 in multiple sites (Table 1) and side by side the most advantageous point is the ability of prophylactic uses which might not be attained with RDV. Other advantages of these drugs are not only cost-effective but also accessible for most of these drugs.

15. Expert opinion

The most fundamental way to stay away from COVID-19 infection is to wear a mask, wash hands frequently, avoid touching the nose and eyes with unwashed hands, maintain social distance, and stay healthy as much as possible [110]. Apart from these, taking essential vitamins and minerals can give a fighting chance to stay uninfected.

As previously discussed, the IMT and DC have been used for the preventive approach on COVID-19. However, the main concern is the dose regimen because frequent intake as a preventive measure can make this drug-resistant, especially DC IMT and DC are not available as OTC (over the counter) drugs so, before starting to take these drugs, one has to discuss with the physician or licensed pharmacist to get a prescription. During these meditations, each individual should be under daily monitoring regarding any side effects.

Zn level in the body is inversely proportional to the interleukin-6, and in the case of COVID-19, Zn was found low. For the patients who died in COVID-19, their Zn levels were much lower than those who survived [111]. Another study also suggested that insufficient Zn level results in poor outcomes in COVID-19 [112].

Vit-D is a very safe supplement, but the dose must be adjusted according to the patient body mass index (BMI) to obtain the optimum level. Vit-D is fat-soluble, so overweight individuals (BMI = 26–30) are suggested to take a 1.5 times greater dose than healthy weighed (BMI ≤ 25) individuals. For obese people (BMI > 30), a high dose (even up to 3-fold) is recommended to obtain the average Vit-D level [113]. EFSA suggested 100 µg/day of Vit-D as an upper limit for adults, including pregnant and lactating women. For 1–10 years old, the upper limit was

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