Since January 2020 Elsevier has created a COVID-19 resource centre with free information in English and Mandarin on the novel coronavirus COVID-19. The COVID-19 resource centre is hosted on Elsevier Connect, the company's public news and information website.

Elsevier hereby grants permission to make all its COVID-19-related research that is available on the COVID-19 resource centre - including this research content - immediately available in PubMed Central and other publicly funded repositories, such as the WHO COVID database with rights for unrestricted research re-use and analyses in any form or by any means with acknowledgement of the original source. These permissions are granted for free by Elsevier for as long as the COVID-19 resource centre remains active.
Combining L-Arginine with vitamin C improves long-COVID symptoms: The LINCOLN Survey

Raffaele Izzo a,1, Valentina Trimarco b,1, Pasquale Mone c, Teresita Aloè d, Massimo Capra Marzani e, Antonio Diana f, Giovanni Fazio g, Mario Mallardo h, Mauro Maniscalco i, Giuseppe Marazzi j, Nunzia Messina k, Simone Mininni l, Chiara Mussi m, Girolamo Pelaia n, Alfio Pennisi o, Pierachille Santus p, Francesco Scarpelli q, Francesco Tursi r, Alessandro Zanforlin s, Gaetano Santulli a, c, *, Bruno Trimarco a

Department of Advanced Biomedical Sciences, Federico II University, Naples, Italy
Department of Neuroscience, Reproductive Sciences and Dentistry, Federico II University, Naples, Italy
Department of Medicine, Albert Einstein College of Medicine, New York, NY, USA
San Martino Hospital, Genoa, Italy
Alessandra Hospital, Alessandria, Italy
Aversa Hospital, Aversa, Caserta, Italy
Palermo Hospital, Palermo, Italy
"San Gennaro" Hospital, Naples, Italy
Istituti Clinici Scientifici Maugeri IRCCS, Telese Terme, Italy
San Raffaele Hospital, Rome, Italy
ASL Napoli 1, Naples, Italy
Associazione Scientifica Interdisciplinare Aggiornamento Medico (ASIAM), Florence, Italy
Department of Biomedical and Metabolic Sciences and Neuroscience, University of Modena and Reggio Emilia, Modena, Italy
Department of Health Science, Magna Graecia University, Catanzaro, Italy
Catania Hospital, Catania, Italy
"Luigi Sacco" University Hospital, Milan, Italy
Trani Hospital, Trani, BT, Italy
Candogia Hospital, ASST Lodi, Lodi, Italy
Health Authority of South Tyrol, Bolzano, Italy

ARTICLE INFO

Keywords:
Ascorbic acid
COVID-19
Endothelial dysfunction
L-Arginine
Long-COVID
Oxidative stress
Survey

ABSTRACT

Introduction: Recent evidence suggests that oxidative stress and endothelial dysfunction play critical roles in the pathophysiology of COVID-19 and Long-COVID. We hypothesized that a supplementation combining L-Arginine (to improve endothelial function) and Vitamin C (to reduce oxidation) could have favorable effects on Long-COVID symptoms.

Methods: We designed a survey (LINCOLN: L-Arginine and Vitamin C improves Long-COVID), assessing several symptoms that have been associated with Long-COVID to be administered nationwide to COVID-19 survivors; the survey also included effort perception, measured using the Borg scale. Patients receiving the survey were divided in two groups, with a 2:1 ratio: the first group included patients that received L-Arginine + Vitamin C, whereas the second group received a multivitamin combination (alternative treatment).

Results: 1390 patients successfully completed the survey. Following a 30-day treatment in both groups, the survey revealed that patients in the L-Arginine + Vitamin C treatment arm had significantly lower scores compared to patients who had received the multivitamin combination. There were no other significant differences between the two groups. When examining effort perception, we observed a significantly lower value (p < 0.0001) in patients receiving L-Arginine + Vitamin C compared to the alternative-treatment arm.

Conclusions: Our survey indicates that the supplementation with L-Arginine + Vitamin C has beneficial effects in Long-COVID, in terms of attenuating its typical symptoms and improving effort perception.

* Corresponding author at: Department of Medicine, Albert Einstein College of Medicine, New York, NY, USA and Department of Advanced Biomedical Sciences, Federico II University, Naples, Italy.
E-mail address: gsantulli001@gmail.com (G. Santulli).
1 Share first authorship.

https://doi.org/10.1016/j.phrs.2022.106360
Received 10 June 2022; Received in revised form 17 July 2022; Accepted 18 July 2022
Available online 19 July 2022
1043-6618/© 2022 Elsevier Ltd. All rights reserved.
1. Introduction

The coronavirus disease 2019 (COVID-19) pandemic caused by the severe acute respiratory syndrome coronavirus-2 (SARS-CoV-2) has led to a global healthcare crisis [1]. The global pandemic has resulted also in job losses and economic hardships [2]. Although efficacious vaccines have been recently developed [3–6], it is important to better understand SARS-CoV-2-mediated pathology, especially because many survivors are experiencing chronic symptoms of the disease, including metabolic disturbances, even months after the initial infection occurred (“Long-COVID”) [7–18].

While initial studies on COVID-19 primarily focused on the pulmonary manifestations of the disease [19–24], other organs including heart, brain, kidney, and the pancreas, were shown to be affected by COVID-19 [25–27]. In the Spring of 2020, we were among the first groups to indicate a link between COVID-19 and endothelial dysfunction [28,29], and our view has been later confirmed by other investigators, associating the manifestations of the disease to a direct or indirect involvement of the endothelium [30–39]. Indeed, endothelial cells express all co-factors necessary for the internalization of SARS-CoV-2 in host cells, including angiotensin converting enzyme 2 (ACE2), transmembrane protease serine 2 (TMPRSS2), cathepsin B and D, neuropilin-1, transferrin receptor, and others, thereby representing a natural target of SARS-CoV-2 [28,39–45].

Thus, COVID-19 affects not only the epithelial cells of the lung [46,47], but also endothelial cells across the whole body, leading to a generalized endothelial damage. Endothelial cells are known to play instrumental roles in the maintenance of vascular homeostasis and in the regulation of vascular tone [48–50]. Endothelial dysfunction, caused directly by SARS-CoV-2 infection and/or by the ensuing inflammation, can shift the vascular equilibrium towards an altered vascular tone, an increased permeabilization, with subsequent tissue edema, and a pro-coagulant state, which can lead to thromboembolism; most of these findings have been substantiated by autopsies of COVID-19 patients since the outbreak of the pandemic [51–53]. Further supporting our theory of a central role of the endothelium in COVID-19, patients with endothelial dysfunction (e.g. hypertension, smoking, pre-existing diabetes, obesity, and presence of cardiovascular disease) are particularly vulnerable and have adverse outcomes in COVID-19 [54]. Equally important, clinical trials testing whether interventions that ameliorate endothelial dysfunction can have beneficial effects in COVID-19 patients are ongoing [55–63] and we have obtained very encouraging interim results in a randomized study testing L-Arginine oral supplementation in COVID-19 hospitalized patients [55].

L-Arginine is an amino acid that acts as a substrate for endothelial nitric oxide (NO) synthase (eNOS), a key enzyme in endothelial cells [64–72]; it has been previously shown to significantly improve endothelial function, providing a strong rationale for its use in COVID-19 patients [58]. Additionally, beneficial effects of L-Arginine on the regulation of immune responses have been reported [73,74]. Similarly, recent clinical trials have demonstrated that Vitamin C improves the oxidative imbalance and vascular remodeling induced by different stressors and attenuates endothelial barrier permeability, an aspect that has major implications in infectious disorders (including COVID-19), which are also known to cause a systemic surge in oxidative stress [75]. So, the antioxidant roles of Vitamin C and its protective effects on endothelial permeability could come into effect especially in post-infection recovery [76–82].

Therefore, we inferred that associating Vitamin C with other nutraceuticals playing similar actions, like L-Arginine, could be useful. On these grounds, considering that both L-Arginine and Vitamin C are known to improve endothelial function and reduce vascular permeability during infectious disorders [75,83–86], and based on the emerging role of vascular permeability in Long-COVID [87,88], we designed a survey to assess the effects of a combination of L-Arginine and Vitamin C in Long-COVID.

2. Methods

2.1. Survey design and participants

This survey (LINCOLN: L-Arginine and Vitamin C improves Long-COVID) starts from the medical need to understand Long-COVID disease. For this reason, in January 2022, we designed a questionnaire based on a seminal Nature Medicine paper published in April 2021 [89] and on the national recommendations on the management of Long-COVID. Our research group includes 16 key opinion leaders belonging to different Italian Regions; a group of physicians (please see appendix) participated in actually administering the LINCOLN survey to patients. The questionnaire was revised by the key opinion leaders and shared with the general practitioners of the nation with the aim to enroll the highest number of patients.

The LINCOLN survey included the following evaluations: patient age, sex, days from RT-qPCR negativization, and hospitalization for moderate/severe COVID-19. Furthermore, the physicians needed to assess on a scale from 1 to 3 (where 1 is absence of the symptom, 2 is the presence of a mild symptom, and 3 is the presence of a severe symptom) - the following symptoms: fatigue, shortness of breath, chest tightness, dizziness, gastrointestinal disorders, headache, anosmia, difficulties in concentrating, sleep disturbances.

Finally, effort perception was evaluated in the survey using the Borg modified 0–10 Rating of perceived exertion scale, with a score from 0 to 10 (where 10 indicate no tolerance), as we previously described [90]. The survey was administered to two groups of COVID-19 survivors who had been COVID-19 negative (confirmation to be SARS-CoV-2 negative via RT-qPCR test [91]) for at least 4 weeks. The first group included patients who had received 2 vials/day of L-Arginine 1.66 g in association with 500 mg of liposomal Vitamin C. The second group (alternative treatment) had been treated with a multivitamin combination (Vitamin B1: 388 mg; Vitamin B2: 443 mg; Nicotinamide:18 mg; Folic Acid: 200 µg; Pantothenic acid: 2493 g; Vitamin B6: 831 mg; Vitamin B12: 416 µg). Physicians were asked to maintain a ~2:1 ratio (L-Arginine + Vitamin C vs alternative treatment) in administering the survey; therefore, they were not blinded; actually, when the questionnaire was administered, patients of both groups had already completed 30 days of treatment.

2.2. Inclusion and exclusion criteria

Eligible patients were men and non-pregnant women who were at least 18-year-old, screened by applying the following eligibility criteria:

**Inclusion criteria**
- Presence of the following conditions:
  - Previous diagnosis of COVID-19 confirmed by RT-qPCR;
  - COVID-19 negativization confirmed by RT-qPCR from at least 4 weeks;
  - Presence of COVID-19 sequelae that extend beyond four weeks after initial infection.

**Exclusion criteria**
- Refusal to participate;
- History of intolerance to L-Arginine, Vitamin C, or components of the multivitamin combination;
- Pregnancy or breastfeeding;
- Cancer;
- Diagnosis of chronic pulmonary disease - currently under treatment;
- Use of immunosuppressive drugs, or cytotoxic chemotherapy within the previous three weeks;
- Enrollment into an investigational treatment study for COVID-19 in the 30 days before screening.

**Survey design**

This survey included the following evaluations: patient age, sex, days from RT-qPCR negativization, and hospitalization for moderate/severe COVID-19. Furthermore, the physicians needed to assess on a scale from 1 to 3 (where 1 is absence of the symptom, 2 is the presence of a mild symptom, and 3 is the presence of a severe symptom) - the following symptoms: fatigue, shortness of breath, chest tightness, dizziness, gastrointestinal disorders, headache, anosmia, difficulties in concentrating, sleep disturbances.

Finally, effort perception was evaluated in the survey using the Borg modified 0–10 Rating of perceived exertion scale, with a score from 0 to 10 (where 10 indicate no tolerance), as we previously described [90]. The survey was administered to two groups of COVID-19 survivors who had been COVID-19 negative (confirmation to be SARS-CoV-2 negative via RT-qPCR test [91]) for at least 4 weeks. The first group included patients who had received 2 vials/day of L-Arginine 1.66 g in association with 500 mg of liposomal Vitamin C. The second group (alternative treatment) had been treated with a multivitamin combination (Vitamin B1: 388 mg; Vitamin B2: 443 mg; Nicotinamide:18 mg; Folic Acid: 200 µg; Pantothentic acid: 2493 g; Vitamin B6: 831 mg; Vitamin B12: 416 µg). Physicians were asked to maintain a ~2:1 ratio (L-Arginine + Vitamin C vs alternative treatment) in administering the survey; therefore, they were not blinded; actually, when the questionnaire was administered, patients of both groups had already completed 30 days of treatment.
3. Results

1590 patients were initially enrolled, of which 1390 fulfilled the eligibility criteria and successfully completed the survey, as shown in the flow chart in Fig. 1.

There were no significant differences in terms of age, male sex, or hospitalization due to COVID-19 between the group treated with L-Arginine + Vitamin C and the arm treated with the alternative treatment (Table 1). Notably, no difference in the time from SARS-CoV-2 negativity was noted between the two groups of patients (33.8 ± 24.9 vs 31.5 ± 24.0 days; ns).

Following a 30-day treatment with L-Arginine + Vitamin C, the survey revealed that patients in this treatment group had significantly lower scores (which means less severe long-COVID symptoms) compared to the other group (L-Arginine + Vitamin C: 8.15 ± 1.3 vs Alternative treatment: 13.9 ± 2.3, p < 0.001); remarkably, the treatment with L-Arginine + Vitamin C had favorable effects on all the symptoms explored by the survey (Table 2).

When examining the effort perception (modified Borg scale), we observed a significantly lower value (p < 0.0001; Fig. 2) in patients receiving L-Arginine + Vitamin C, compared to the alternative-treatment arm, indicating that L-Arginine + Vitamin C led to a better effort perception.

4. Discussion

This survey is the first to show the beneficial effects of the combination of L-Arginine and Vitamin C in Long-COVID. Our investigation was based on a robust rationale, i.e., targeting endothelial dysfunction in Long-COVID. Indeed, endothelial cell infection with consecutive inflammatory cell recruitment and endothelial dysfunction could explain the impaired microcirculation observed across vascular beds in COVID-19, triggering vasoconstriction, ischemia, and a pro-coagulant state [92-95]. Consistent with our view, several investigators had proposed that endotheliitis could be a critical mechanism underlying systemic impaired microcirculatory function observed in different vascular beds in patients experiencing Long-COVID symptoms [92,96].

Of note, a recent clinical study has demonstrated that COVID-19 patients develop endothelial dysfunction, which remains significantly impaired compared to healthy controls subjects at a 6-month follow-up [97], implicating that endothelial dysfunction is a main player in both acute COVID-19 and Long-COVID. Interestingly, increased numbers of circulating endothelial cells, a biomarker of vascular injury [98], most likely detached from the vessel wall due to pathological insults, were found to significantly correlate with the severity of COVID-19 outcome [99]. Strikingly, elevated levels of circulating endothelial cells persisted in recovered COVID-19 convalescent patients [100], denoting long-term detrimental effects of SARS-CoV-2 infection on endothelial function. Actual signs of endothelial dysfunction (denoted by glycocalyx damage) have been reported in convalescent COVID-19 patients after mild disease progression without hospitalization [101]. Other studies evidenced that sustained endotheliopathy is common in convalescent COVID-19 subjects [102], and that Long-COVID symptoms, specifically non-respiratory symptoms, are mainly due to persistent endothelial dysfunction [103].

Our data are fully in agreement with previous reports implying that after the acute phase of COVID-19, the disease is dominated by immunopathological pro-inflammatory elements [104-110]. In fact, previous investigations have actually demonstrated that reduced levels of L-Arginine increase the generation of reactive oxygen species (ROS), aggravating inflammation [111]. Besides, in vitro assays revealed that the T cell proliferative capacity is significantly reduced among COVID-19 patients [92-95]. Consistent with our view, several investigators had proposed that endotheliitis could be a critical mechanism underlying systemic impaired microcirculatory function observed in different vascular beds in patients experiencing Long-COVID symptoms [92,96].
Fatigue is a prevailing symptom in Long-COVID patients [121], and throughout Italy. Nevertheless, we do reckon that our research is not exempt from limitations. For instance, we did not assess blood levels of nitrates and nitrite, Vitamin C, or citrulline in our patients; moreover, we did not administer the survey at baseline but only after 30 days. Hence, one may argue that the patients who received L-Arginine + Vitamin C could have been somehow healthier than the other group before starting the treatment. However, albeit we do not have full clinical data for all patients, this hypothesis seems unlikely since patients were enrolled by their family physicians who had prescribed different treatments to patients with similar clinical characteristics only on account of little guidance on managing Long-COVID. Further dedicated interventional studies are warranted to endorse our findings.

Table 2
Survey results in the two groups of patients.

| Condition                  | Alternative Treatment (N=521) | L-Arginine + Vitamin C (N=869) | p    |
|----------------------------|-------------------------------|--------------------------------|------|
|                            | Absent (%)                    | Mild (%)                       | Severe (%) | Absent (%) |
| Asthenia                   | 0.4                           | 5.2                            | 94.9      | 4.0        | <0.0001 |
| Dyspnea                    | 55.1                          | 19.5                           | 1.1       | 0.4        | <0.0001 |
| Chest tightness            | 50.9                          | 22.8                           | 13.4      | 0.5        | <0.0001 |
| Dizziness                  | 25.9                          | 7.5                            | 11.6      | 1.1        | <0.0001 |
| Gastrointestinal disorders | 26.7                          | 10.0                           | 11.7      | 0.6        | <0.0001 |
| Headache                   | 44.1                          | 16.8                           | 16.8      | 1.4        | <0.0001 |
| Anosmia                    | 34.0                          | 14.0                           | 11.0      | 1.8        | <0.0001 |
| Concentration difficulty   | 46.8                          | 20.4                           | 19.4      | 1.5        | <0.0001 |
| Sleeplessness              | 42.6                          | 39.2                           | 17.5      | 1.8        | <0.0001 |

As a critical driver of inflammation and oxidative stress [117], endothelial dysfunction has also been involved in the pathophysiology of the neurological symptoms of COVID-19 and Long-COVID [118–120]. Fatigue is a prevailing symptom in Long-COVID patients [121], and previous trials have evidenced a significant decrease in fatigue in subjects treated with Vitamin C [122,123], fully consistent with our present results. Supporting our strategy to combine L-Arginine to Vitamin C, previous investigations have shown that ascorbic acid can synergistically improve the effects of other agents: for example, if added to glucagon-like peptide 1 (GLP-1) agonists, it reduces ROS generation in diabetic patients [124], in combination with metformin, it reduces macro- and microvascular diabetic complications [125].

Strengths of this survey include the large population enrolled and the fact that the questionnaire was administered in multiple centers throughout Italy. Nevertheless, we do reckon that our research is not exempt from limitations. For instance, we did not assess blood levels of nitrates and nitrite, Vitamin C, or citrulline in our patients; moreover, we did not administer the survey at baseline but only after 30 days. Hence, one may argue that the patients who received L-Arginine + Vitamin C could have been somehow healthier than the other group before starting the treatment. However, albeit we do not have full clinical data for all patients, this hypothesis seems unlikely since patients were enrolled by their family physicians who had prescribed different treatments to patients with similar clinical characteristics only on account of little guidance on managing Long-COVID. Further dedicated interventional studies are warranted to endorse our findings.

CRediT authorship contribution statement

Raffaele Izzo: Writing - original draft, Writing - review & editing. Valentina Trimarco: Conceptualization, Data curation, Formal analysis, Software. Pasquale Mone: Visualization, Writing – review & editing. Teresita Aloe: Investigation, Visualization. Massimo Capra Marzani: Investigation, Visualization. Antonio Diana: Investigation, Visualization. Giovanni Fazio: Investigation, Visualization. Mario Mallardo: Investigation, Visualization. Mauro Maniscalco: Investigation, Visualization. Giuseppe Marazzi: Investigation, Visualization. Nunzia Messina: Investigation, Visualization. Simone Mininni: Investigation, Visualization. Chiara Musi: Investigation, Visualization. Girolamo Pelaia: Investigation, Visualization. Alfio Pennisi: Investigation, Visualization. Pierachille Santus: Investigation, Visualization. Francesco ScarPELLi: Investigation, Visualization. Francesco Tursi: Investigation, Visualization. Alessandro Zanforlin: Investigation, Visualization. Gaetano Santulli: Writing - original draft, Writing - review & editing. Bruno Trimarco: Writing - original draft.

Declaration of Competing Interest

None.

Data availability

Data will be made available on request.
A. Sharma, et al., A randomized open-label trial to evaluate the efficacy and safety of triple therapy with aspirin, atorvastatin, and nicorandil in hospitalised patients with SARS-CoV-2 infection: a structured summary of a study protocol for a randomized controlled trial, Trials 22 (2021) 451, https://doi.org/10.1186/s13063-021-00536-y.

S. Srivastava, I. Garg, A.A. Hembrom, B. Kumar, Assessment of nitric oxide (NO) potential to mitigate COVID-19 severity, Viruses 21 (2021) 589–594, https://doi.org/10.3390/v13040492.

A.M. Isidori, et al., Targeting the NO-cGMP-PDE5 pathway in COVID-19 infection. The DEDALO project, Andrology 9 (2021) 33, https://doi.org/10.1111/andr.12837.

P. Mone, et al., L-Arginine enhances the effects of cardiac rehabilitation on restoration of myocardial function in patients with COVID-19: results of a placebo-controlled, randomized trial, JACC Basic Transl. Sci. 6 (2021) 935–944, https://doi.org/10.1016/j.jbts.2021.09.013.

S. Charfeddine, et al., Long COVID 19 syndrome: is it related to microcirculation syndrome, J. Thromb. Haemost. 19 (2021) 2546–2554, https://doi.org/10.1111/jth.15490.

C. Wang, et al., Long COVID: the nature of thrombotic sequelae determines the long-term consequences, Front Cell Dev. Biol. 10 (2022), 824851, https://doi.org/10.3389/febcl.2022.824851.

C.H. Sudre, et al., Attributes and predictors of long COVID, Nat. Med. 27 (2021) 626–631, https://doi.org/10.1038/s41591-021-01292-y.

P. Mone, et al., L-Arginine enhances the effects of cardiac rehabilitation on physical performance: new insights for managing cardiovascular patients during the COVID-19 pandemic, J. Pharm. Exp. Ther. 381 (2022) 197–203, https://doi.org/10.1124/jpet.122.020114.

S.A. Bustin, T. Nolan, RT-qPCR testing of SARS-CoV-2: a primer, Int. J. Mol. Sci. 21 (2020), https://doi.org/10.3390/ijms21030382.

D. Zhao, M. Fu, Y. Qian, Vascular endothelial glycodelin and potential targeted therapy in COVID-19, Cells 11 (2022) 1972, https://doi.org/10.3390/cells11111972.

J. Jamieson, et al., Role of endothelial miR-24 in COVID-19 cerebrovascular events, Crit. Care 25 (2021) 306, https://doi.org/10.1186/s13054-021-03731-2.

P. Andrusiak, et al., Endothelial dysfunction in COVID-19: a unifying mechanism and a potential therapeutic target, Biomolecules 10 (2022), https://doi.org/10.3390/biom10040812.

J. Jamieson, et al., Long COVID 19 syndrome: is it related to microcirculation syndrome, J. Thromb. Haemost. 19 (2021) 2546–2554, https://doi.org/10.1111/jth.15490.

A. Seitz, P. Ong, Endothelial dysfunction in COVID-19: a potential predictor of long-COVID, Int. J. Cardiol. 349 (2022) 155–156, https://doi.org/10.1016/j.ijcard.2021.11.051.

E. Oikonomou, et al., Endothelial dysfunction in acute and long standing COVID-19: a prospective cohort study, Vasc. Pharmac. 144 (2022), 106975, https://doi.org/10.1016/j.vph.2022.106975.

A.D. Blann, et al., Circulating endothelial cells, Biomark. Vasc. Dis. Thromb. Haemost. 93 (2005) 228–235, https://doi.org/10.1016/S2665-9913(20)30275-7.

S. Guerville, et al., Circulating endothelial cells as a marker of endothelial injury in severe COVID-19, J. Infect. Dis. 222 (2020) 1789–1793, https://doi.org/10.1093/infdis/jiaa528.

F.W. Chioh, et al., Convalescent COVID-19 patients are susceptible to endothelial dysfunction due to persistent immune activation, Elife 10 (2021), https://doi.org/10.7554/elife.64909.

V. Bronte, P. Zanovello, Regulation of immune responses by L-arginine metabolism, Nat. Rev. Immunol. 5 (2005) 641–654, https://doi.org/10.1038/nri1667.

J. Parnell, et al., Persistent endotheliopathy in the pathogenesis of long COVID syndrome, J. Thromb. Haemost. 19 (2021) 2546–2553, https://doi.org/10.1111/jth.15490.

S. Charfeddine, et al., Long COVID syndrome: is it related to microcirculation and endothelial dysfunction? Insights from TUN-EndoCV study, Front Cardiovasc. Med. 8 (2021), 745758, https://doi.org/10.3390/fcmv2021.745758.

S.D. Datta, A. Talwar, J.T. Lee, A proposed framework and timeline of the spectrum of disease due to SARS-CoV-2 infection: illness beyond acute infection and public health implications, JAMA 324 (2020) 2251–2252, https://doi.org/10.1001/jama.2020.22717.

M. Taboada, V. Carazo, A. Naveira, P.G. Atanassoff, Corticosteroids and the hyper-inflammatory phase of the COVID-19 disease, J. Clin. Anesth. 66 (2020), https://doi.org/10.1016/j.jclinane.2020.09.026.

L.F. Garcia, Immune response, inflammation, and the clinical spectrum of COVID-19, Front Immunol. 11 (2020) 1441, https://doi.org/10.3389/fimmu.2020.01441.

J.J. Manon, et al., COVID-19-associated hyperinflammation and exacerbation of patient care: a retrospective longitudinal cohort study, Lancet Rheuma 2 (2020) e594–e602, https://doi.org/10.1016/S2665-9913(20)30275-7.

M.Z. Tay, C.M. Poh, L. Renia, P.A. MacAry, L.F.P. Ng, The trinity of COVID-19: immunity, inflammation and intervention, Nat. Rev. Immunol. 20 (2020) 363–374, https://doi.org/10.1038/s41577-020-0311-8.

G.R.S. Buding, A.V. Marthin, K.M. Ridge, B.D. Singer, R.G. Wunderink, Distinctive features of COVID-19 pneumonia, J. Mol. Cell Med. 21 (2021) 741–745, https://doi.org/10.1111/jmcm.149412.
[111] R. Lucas, et al., Arginase 1: an unexpected mediator of pulmonary capillary barrier dysfunction in models of acute lung injury, Front Immunol. 4 (2013) 228, https://doi.org/10.3389/fimmu.2013.00228.

[112] F. Reizine, et al., SARS-CoV-2-induced ARDS associates with MDSC expansion, lymphocyte dysfunction, and arginine shortage, J. Clin. Immunol. 41 (2021) S15–S25, https://doi.org/10.1007/s10875-020-00928-5.

[113] M.J. Dean, et al., Transcriptome and functions of granulocytic myeloid-derived suppressor cells determine their association with disease severity of COVID-19, medRxiv (2021), https://doi.org/10.1101/2021.03.26.21254441.

[114] A. Sacchì, et al., Expansion of myeloid derived suppressor cells contributes to platelet activation by L-Arginine deprivation during SARS-CoV-2 infection, Cells 10 (2021), https://doi.org/10.3390/cells10082111.

[115] M. Gelzo, et al., Inducible Nitric Oxide Synthase (iNOS): why a different production in COVID-19 patients of the two waves, Viruses 14 (2022), https://doi.org/10.3390/v14030393.

[116] R. Cinar, M.R. Iyer, G. Kunos, Dual inhibition of CB1 receptors and iNOS, as a potential novel approach to the pharmacological management of acute and long COVID-19, Br. J. Pharm. 179 (2022) 2121–2127, https://doi.org/10.1111/bph.15461.

[117] H. Crook, S. Raza, J. Nowell, M. Young, P. Edison, Long covid-mechanisms, risk factors, and management, BMJ 374 (2021) n1648, https://doi.org/10.1136/bmj.n1648.

[118] S.A. Azizi, S.A. Azizi, Neurological injuries in COVID-19 patients: direct viral invasion or a bystander injury after infection of epithelial/endothelial cells, J. Neurovirol. 26 (2020) 631–641, https://doi.org/10.1007/s13365-020-00903-7.

[119] D. Castañares-Zapatero, et al., Pathophysiology and mechanism of long COVID: a comprehensive review, Ann. Med. 54 (2022) 1473–1487, https://doi.org/10.1080/07853890.2022.2076601.

[120] I. Ruktai, et al., Neuropathology and virus in brain of SARS-CoV-2 infected non-human primates, Nat. Commun. 13 (2022) 1745, https://doi.org/10.1038/s41467-022-29440-z.

[121] N. Prasanan, et al., Impaired exercise capacity in post-COVID syndrome: the role of VWF-ADAMTS13 axis, Blood Adv. (2022), https://doi.org/10.1182/bloodadvances.2021006944.

[122] S.Y. Suh, et al., Intravenous vitamin C administration reduces fatigue in office workers: a double-blind randomized controlled trial, Nutr. J. 11 (2012) 7, https://doi.org/10.1186/1475-2891-11-7.

[123] C.M. Stephenson, R.D. Levin, T. Spector, C.G. Liu, Phase I clinical trial to evaluate the safety, tolerability, and pharmacokinetics of high-dose intravenous ascorbic acid in patients with advanced cancer, Cancer Chemother. Pharm. 72 (2013) 139–146, https://doi.org/10.1007/s00280-013-2179-9.

[124] A. Ceriello, et al., Vitamin C further improves the protective effect of glucagon-like peptide-1 on acute hypoglycemia-induced oxidative stress, inflammation, and endothelial dysfunction in type 1 diabetes, Diabetes Care 36 (2013) 4104–4108, https://doi.org/10.2337/dc13-0750.

[125] S.W. Gillani, S.A.S. Sulaiman, M.I.M. Abdul, M.R. Baig, Combined effect of metformin with ascorbic acid versus acetyl salicylic acid on diabetes-related cardiovascular complication; a 12-month single blind multicenter randomized control trial, Cardiovasc. Diabetol. 16 (2017) 103, https://doi.org/10.1186/s12933-017-0584-9.