The Functional Medicine Approach to COVID-19: Virus-Specific Nutraceutical and Botanical Agents

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Background and Introduction

Health professionals and the public must be well informed about the COVID-19 virus, the disease it causes, and how it spreads. This information is readily available and not within the scope of this document. At this time, there are no specific vaccines or uniformly successful treatments for COVID-19. In this context of insufficient evidence, the scope of this document will be to assess the scientific plausibility of promising prevention approaches and therapeutic (nutraceutical and botanical) interventions and then to offer clinical recommendations.

With respect to interventions, the practice of Functional Medicine emphasizes the primacy of safety, validity, and effectiveness. In the novel context of COVID-19, validity in the form of
recommendations represent the functional medicine approach to the COVID-19 crisis:

- Adherence to all health recommendations from official sources to decrease viral transmission.
- Optimizing modifiable lifestyle factors in order to improve overall immune function (an introductory document on ‘Boosting Immunity’ is available [here](#)):  
  - Reduces progression from colonization to illness.
- Personalized consideration of therapeutic agents that may:
  - Favorably modulate cellular defense and repair mechanisms.
  - Favorably modulate viral-induced pathological cellular processes.
  - Promote viral eradication or inactivation.
  - Mitigate collateral damage from other therapeutic agents.
  - Promote resolution of collateral damage and restoration of function.
- Treatment of confirmed COVID-19 illness (as per conventional standards and practice):
  - May reduce the severity and duration of acute symptoms and complications.
  - May support recovery and reduce long-term morbidity and sequelae.

Additional references are being collated and will be made available in the future.

Clinical Recommendations and Mechanisms of Action

**BACKGROUND AND MECHANISMS OF ACTION**

We encourage practitioners to learn about the mechanism of invasion, replication, and pathophysiology of the COVID-19 virus. Much of what we know has been extrapolated from basic science research on SARS-CoV-2. Excellent resources are available online, including the free YouTube lectures through Dr. Roger Seheult:

This document discusses the mechanisms of action of a number of different botanical and nutraceutical agents. These agents can be considered as immunoadjuvants, defined as substances that act to accelerate, prolong, or enhance antigen-specific immune responses by potentiating or modulating the immune response.\(^1\)
A 2016 review article[4] entitled "Natural compounds as regulators of NLRP3 inflammasome-mediated IL-beta production" notes that "resveratrol, curcumin, EGCG [epigallocatechin gallate], and quercetin are potent inhibitors of NLRP3 inflammasome-mediated IL-1beta production, typically acting at more than one element of the involved pathways. However, it should be noted that these polyphenols have an even much broader biological effect, as they influence a variety of pathways.” For example, these polyphenols modulate NF-kB upregulation, which is useful to counteract the COVID-19 'hyper-inflammation'.[6]

A preprint released on March 23, 2020, identified the ability of plant bioactive compounds to inhibit the COVID-19 main protease (M^{pro}),[7] which is necessary for viral replication. There is much excitement surrounding the recent identification of M^{pro}, and it is a current potential pharmaceutical drug target. Kaempferol, quercetin, luteolin-7-glucoside, demethoxycurcumin, naringenin, apigenin-7glucoside, oleuropein, curcumin, catechin, and epicatechin-gallate were the natural compounds that appeared to have the best potential to act as COVID-19 M^{pro} inhibitors. Though further research is necessary to prove their efficacy, this study provides the biologic plausibility and mechanistic support (COVID-19 protease inhibition) to justify their use.

For these reasons, we recommend the following compounds, at standard dosages, to prevent activation of the NLRP3 inflammasome, to decrease NF-kB activation, and to potentially inhibit COVID-19 replication. There is no literature to support a regimen of a single vs. multiple agents. Our recommendation is to use higher dosing and/or multiple agents when patient contextual factors (e.g., patient desire, pre-existing inflammation, multiple co-morbidities, higher risk, etc.) and/or therapeutic decision-making warrant such use.

In the recommendations below, the following criteria are used to identify strength of evidence and risk of harm.

Recommended Interventions

QUERCETIN

Quercetin has been shown to have antiviral effects against both RNA (e.g., influenza and coronavirus) and DNA viruses (e.g., herpesvirus). Quercetin has a pleiotropic role as an antioxidant and anti-inflammatory, modulating signaling pathways that are associated with post-transcriptional modulators affecting post-viral healing.[8]
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<table>
<thead>
<tr>
<th>Mechanism(s) of action against non-COVID-19 viruses</th>
<th>Promote viral eradication or inactivation.¹⁹,¹⁰,¹¹,¹²,¹³</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Inhibition of viral replication</td>
</tr>
<tr>
<td></td>
<td>Favorably modulate viral-induced pathological cellular processes:</td>
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<tr>
<td></td>
<td>• Modulation of NLRP3 inflammasome activation⁵,¹⁴,¹⁵</td>
</tr>
<tr>
<td></td>
<td>Mechanistically promote resolution of collateral damage and restoration of function:</td>
</tr>
<tr>
<td></td>
<td>• Modulation of mast cell stabilization (anti-fibrotic)</td>
</tr>
</tbody>
</table>

| Outcomes data supporting their mitigating          | Reduction of symptoms                                  |
| Strength of evidence                               | Moderate                                               |
| Risk of harm:⁶,⁷,¹⁶,¹⁷                                  | Mild                                                  |

**CURCUMIN**

Curcumin has been shown to modulate the NLRP3 inflammasome,⁵ and a preprint suggests that curcumin can target the COVID-19 main protease to reduce viral replication.¹⁸

<table>
<thead>
<tr>
<th>Intervention</th>
<th>Curcumin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Suggested Dose</td>
<td>500-1,000 mg po bid (of absorption-enhanced curcumin)</td>
</tr>
<tr>
<td>Mechanism(s) of action against non-COVID-19 viruses</td>
<td>Favorably modulate viral-induced pathological cellular processes:</td>
</tr>
<tr>
<td></td>
<td>• Modulation of NLRP3 inflammasome activation⁵,¹⁹,²⁰,²¹</td>
</tr>
<tr>
<td>Outcomes data supporting their mitigating</td>
<td>No data available</td>
</tr>
</tbody>
</table>
EPIGALLOCATECHIN GALLATE (EGCG)

Green tea, in addition to modulating the NLRP3 inflammasome and, based on a preprint, potentially targeting the COVID-19 main protease (M^{bro})[^31] to reduce viral replication, has also been shown to prevent influenza in healthcare workers.[^28]

<table>
<thead>
<tr>
<th>Intervention</th>
<th>Epigallocatechin gallate (EGCG)</th>
</tr>
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<tbody>
<tr>
<td>Suggested Dose</td>
<td>4 cups daily or 225 mg po qd</td>
</tr>
<tr>
<td>Mechanism(s) of action against non-COVID-19 viruses</td>
<td>Favorably modulate viral-induced pathological cellular processes: • Modulation of NLRP3 inflammasome activation[^5],[^28],[^29]</td>
</tr>
<tr>
<td>Outcomes data supporting their mitigating effects on illness from other viral strains</td>
<td>No data available</td>
</tr>
<tr>
<td>Strength of evidence</td>
<td>Conditional</td>
</tr>
<tr>
<td>Risk of harm:[^30],[^31],[^32],[^33],[^34],[^35]</td>
<td>Significant (rare) - Hepatotoxicity</td>
</tr>
</tbody>
</table>

N-ACETYLCYSTEINE (NAC)

N-acetylcysteine promotes glutathione production, which has been shown to be protective in rodents infected with influenza. In a little-noticed six-month controlled clinical study enrolling 262 primarily elderly subjects, those receiving 600 mg NAC twice daily, as opposed to those receiving placebo, experienced significantly fewer influenza-like episodes and days of bed confinement.[^36]
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**RESVERATROL**

Resveratrol, a naturally occurring polyphenol, shows many beneficial health effects. It has been shown to modulate the NLRP3 inflammasome.\[^5\] In addition, resveratrol was shown to have in vitro activity against MERS-CoV.\[^43\]

<table>
<thead>
<tr>
<th>Intervention</th>
<th>Resveratrol</th>
</tr>
</thead>
<tbody>
<tr>
<td>Suggested Dose</td>
<td>100-150 mg po qd</td>
</tr>
</tbody>
</table>
| Mechanism(s) of action against non-COVID-19 viruses | Favorably modulate viral-induced pathological cellular processes
- Modulation of NLRP3 inflammasome activation\[^5\] |
| Outcomes data supporting their mitigating effects on illness from other viral strains | MERS-CoV\[^43\]
Influenza\[^44\],\[^45\] |
| Strength of evidence | Conditional |
| Risk of harm:\[^46\],[^47],[^48],[^49],[^50],[^51],[^52],[^53]\] | Mild |
stimulates the expression of potent antimicrobial peptides (AMPs), which exist in neutrophils, monocytes, natural killer cells, and epithelial cells of the respiratory tract.[54] Vitamin D increases anti-pathogen peptides through defensins and has a dual effect due to suppressing superinfection. Evidence suggests vitamin D supplementation may prevent upper respiratory infections.[55] However, there is some controversy as to whether it should be used and the laboratory value that should be achieved. Research suggests that concerns about vitamin D (increased IL-1beta in cell culture) are not seen clinically. The guidance we suggest is that a laboratory range of >50 and <80 ng/mL serum 25-hydroxy vitamin D may help to mitigate morbidity from COVID-19 infection.

<table>
<thead>
<tr>
<th>Intervention</th>
<th>Vitamin D</th>
</tr>
</thead>
<tbody>
<tr>
<td>Suggested Dose</td>
<td>5,000 IU po qd in the absence of serum levels</td>
</tr>
</tbody>
</table>
| **Mechanism(s) of action against non-COVID-19 viruses**[55],[56],[57],[58],[59],[60],[61],[62],[63],[64],[65],[66],[67],[68],[69],[70],[71],[72],[73],[74],[75],[76],[77],[78] | *Favorably modulate cellular defense and repair mechanisms:*  
  • Activation of macrophages Stimulation of anti-microbial peptides  
  • Stimulation of anti-microbial peptides  
  • Modulation of defensins  
  • Modulation of TH17 cells  
  *Favorably modulate viral-induced pathological cellular processes:*  
  • Reduction in cytokine expression  
  • Modulation of TGF beta |
| Outcomes data supporting their mitigating effects on illness from other viral strains | Reduce progression from colonization to illness Reduce the severity and duration of acute symptoms and complications |
| Strength of evidence                                                          | Limited                                                                   |
| Risk of harm:[79],[80],[81],[82]                                              | Mild                                                                     |

**MELATONIN**

Melatonin has been shown to have an inhibitory effect on the NLRP3 inflammasome.[94] This has not gone unnoticed by the COVID-19 research community, with two recent published papers...
<table>
<thead>
<tr>
<th>Intervention</th>
<th>Melatonin</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Suggested Dose</strong></td>
<td><strong>5-20 mg qd</strong></td>
</tr>
</tbody>
</table>
| **Mechanism(s) of action against non-COVID-19 viruses**.\(^{[83],[84]}\) | *Favorably modulate viral-induced pathological cellular processes*  
  - Modulation of NLRP3 inflammasome activation.\(^{[83],[84]}\) |
| **Outcomes data supporting their mitigating effects on illness from other viral strains** | Research in progress |
| **Strength of evidence**                                                | Conditional                    |
| **Risk of harm:**\(^{[86],[87],[88],[89],[90],[91],[92],[93],[94]}\) | Mild                           |

**VITAMIN A**

Vitamin A is a micronutrient that is crucial for maintaining vision, promoting growth and development, and protecting epithelium and mucus integrity in the body. Vitamin A is known as an anti-inflammation vitamin because of its critical role in enhancing immune function. Vitamin A is involved in the development of the immune system and plays regulatory roles in cellular immune responses and humoral immune processes through the modulation of T helper cells, sIgA, and cytokine production. Vitamin A has demonstrated a therapeutic effect in the treatment of various infectious diseases\(^{[95]}\)

<table>
<thead>
<tr>
<th>Intervention</th>
<th>Vitamin A</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Suggested Dose</strong></td>
<td><strong>Up to 10,000-25,000 IU/d</strong></td>
</tr>
</tbody>
</table>
| **Mechanism(s) of action against non-COVID-19 viruses** \(^{[95],[96]}\) | *Favorably modulate cellular defense and repair mechanisms:*  
  - Modulation of T helper cells  
  - Modulation of sIgA  
  *Favorably modulate viral-induced* |
ELDERBERRY

Elderberry (Sambucus nigra) is seen in many medicinal preparations and has widespread historical use as an anti-viral herb.[103] Based on animal research, elderberry is likely most effective in the prevention of and early infection with respiratory viruses. [104] One in-vitro study reported an increase in TNF-alpha levels related to a specific commercial preparation of elderberry[105] leading some to caution that its use could initiate a “cytokine storm.” However, these data were not confirmed when the same group performed similar studies, which were published in 2002.[106] Therefore, these data suggest it is highly implausible that consumption of properly prepared elderberry products (from berries or flowers) would contribute to an adverse outcome related to overproduction of cytokines or lead to an adverse response in someone infected with COVID-19.

<table>
<thead>
<tr>
<th>Intervention</th>
<th>Elderberry</th>
</tr>
</thead>
<tbody>
<tr>
<td>Suggested Dose</td>
<td>500 mg po qd (of USP standard of 17% anthocyanosides)</td>
</tr>
</tbody>
</table>
| Mechanism(s) of action against non-COVID-19 viruses | Favorably modulate cellular defense and repair mechanisms  
Favorably modulate viral-induced pathological cellular processes |
| Outcomes data supporting their mitigating effects on illness from other viral strains | No data available |
| Strength of evidence | Strong |
| Risk of harm | Mild; caution w/autoimmune disease; uncooked/unripe plant parts toxic; USDA GRAS |
PEA is a naturally occurring anti-inflammatory palmitic acid derivative that interfaces with the endocannabinoid system. There was a significantly favorable outcome in five of six double blind placebo-controlled trials looking at acute respiratory disease due to influenza.\textsuperscript{[115]} Dosing was generally 600 mg three times daily for up to three weeks. There are multiple mechanisms of action associated with PEA, from inhibition of TNF-alpha and NF-kB to mast cell stabilization. In influenza, it is thought that PEA works by attenuating the potentially fatal cytokine storm.

<table>
<thead>
<tr>
<th>Intervention</th>
<th>Palmitoylethanolamide (PEA)</th>
</tr>
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<tbody>
<tr>
<td>Suggested Dose</td>
<td>\textit{300 mg po bid to prevent infection, 600 mg po tid x two weeks to treat infection}</td>
</tr>
<tr>
<td>Mechanism(s) of action against non-COVID-19 viruses\textsuperscript{[115]}</td>
<td>\textit{Favorably modulate cellular defense and repair mechanisms  Favoredly modulate viral-induced pathological cellular processes}</td>
</tr>
<tr>
<td>Outcomes data supporting their mitigating effects on illness from other viral strains</td>
<td>No data available</td>
</tr>
<tr>
<td>Strength of evidence</td>
<td>Conditional (treatment) Strong (prevention)</td>
</tr>
<tr>
<td>Risk of harm:\textsuperscript{[116],[117],[118],[119]}</td>
<td>Mild</td>
</tr>
</tbody>
</table>

**VITAMIN C**

Vitamin C contributes to immune defense by supporting various cellular functions of both the innate and adaptive immune system. Vitamin C accumulates in phagocytic cells, such as neutrophils, and can enhance chemotaxis, phagocytosis, generation of reactive oxygen species, and ultimately microbial killing. Supplementation with vitamin C appears to be able to both prevent and treat respiratory and systemic infections.\textsuperscript{[120]} Vitamin C has been used in hospital ICUs to treat COVID-19 infection.
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<table>
<thead>
<tr>
<th>Viruses</th>
<th>Repair mechanisms</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Favorably modulate viral-induced pathological cellular processes</td>
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<table>
<thead>
<tr>
<th>Outcomes data supporting their mitigating effects on illness from other viral strains</th>
<th>No data available</th>
</tr>
</thead>
<tbody>
<tr>
<td>Strength of evidence</td>
<td>Strong</td>
</tr>
<tr>
<td>Risk of harm[121]</td>
<td>Mild</td>
</tr>
</tbody>
</table>

Evaluative Criteria

In the recommendations above, the following criteria are used to identify strength of evidence and risk of harm.

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**Strength of Evidence**

- **Strength of Evidence Conditional**
  
  Clinical experience and/or expert opinion and/or conflicting studies; biological mechanism at least partly explained.

- **Strength of Evidence Limited**
  
  One study showing correlation between intervention and outcome; compelling ATMs and/or PCFs; biological mechanism at least partly explained.

**Risk of Harm**

- **Risk of Harm Mild**
  
  Risk of self-limited symptoms; no risk of loss of function or corrective intervention anticipated; observation only.

- **Risk of Harm Moderate**
  
  Risk of symptoms; no risk of loss of function or quality of life; minor evaluative and/or therapeutic intervention needed.
between intervention and outcome; biological mechanism at least partly explained.

and/or therapeutic intervention needed.

<table>
<thead>
<tr>
<th>Strength of Evidence</th>
<th>Risk of Harm</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Strong</strong></td>
<td><strong>Severe</strong></td>
</tr>
<tr>
<td>Two independent studies (both LOE = 1 or 2) showing correlation between intervention and outcome; biological mechanism fully explained or partly explained and having one additional correlative study.</td>
<td>Risk of permanent symptoms, loss of function, quality of life, or death; long-term evaluative and/or therapeutic intervention needed.</td>
</tr>
</tbody>
</table>

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Joel Evans, MD (Lead), Robert Rountree, MD, Tom Guiliams, PhD, Michael Stone, MD, Robert Luby, MD, Patrick Hanaway, MD, Kirsten Ramsdell, MS, CN, Sam Yanuck, DC, Helen Messier, MD, and Dan Lukaczer, ND,

Click here to see all references

REFERENCES


Sanders KM, Stuart AL, Williamson EJ, et al. Annual high-dose oral vitamin D and falls and fractures in older women: a randomized controlled trial. *JAMA.* 2010;303(18):1815-1822. doi:10.1001/jama.2010.594


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