

Selenium and the Limits of Lumping All Antioxidants Together

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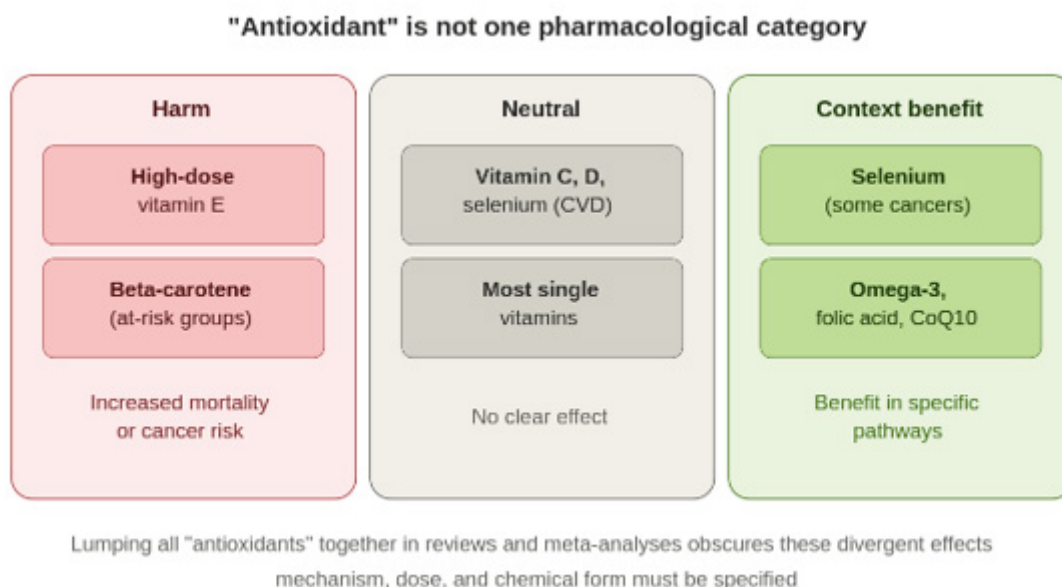
Dear Editor,

Reviews and meta-analyses frequently pool diverse compounds under the single heading of “antioxidants,” as if membership in that category predicted a common biological effect. I write to argue, using selenium as a case in point, that this lumping obscures more than it reveals.

The divergence within the category is substantial. According to PubMed, while high-dose vitamin E has been associated with increased all-cause mortality, and a large meta-analysis found that beta-carotene supplementation increased mortality and stroke risk, the same analysis reported that omega-3 fatty acids, folic acid, and coenzyme Q10 conferred benefits on specific

outcomes—so the label “antioxidant” spans agents that harm, agents that do nothing, and agents that help, depending on the endpoint.

Selenium illustrates the point sharply. According to PubMed, although the epidemiological data are described as somewhat controversial, numerous studies suggest selenium compounds can negatively affect cancer progression, and *in vitro* and animal work has linked selenium supplementation to reduced cell migration, invasion, angiogenic factors, and metastasis in several cancer types, prompting proposals that selenium may act as an anti-metastatic as well as a preventive agent. This is a very different profile from that of high-dose Vitamin E.



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When such heterogeneous agents are combined into a single “antioxidant” effect estimate, the result can be a meaningless average that misrepresents every constituent. A null pooled estimate may hide a harmful agent and a beneficial one canceling out; a harmful pooled estimate may unfairly tarnish compounds with genuine promise. Either way, the category label has done analytical harm.

I would encourage authors and editors to resist treating “antioxidants” as a unified intervention class. Mechanism, chemical form, dose, target tissue, and clinical context all differ across these compounds and should be specified and analyzed

separately. Selenium’s distinctive profile is a reminder that the shared ability to donate an electron in a test tube is not a sound basis for grouping agents in a clinical meta-analysis.

Sincerely,

Mueen Ahmed KK

REFERENCES

- An, P., Wan, S., Luo, Y., Luo, J., Zhang, X., Zhou, S., Xu, T., He, J., Mechanick, J. I., Wu, W.-C., Ren, F., & Liu, S. (2022). Micronutrient supplementation to reduce cardiovascular risk. *Journal of the American College of Cardiology*, 80(24), 2269–2285. <https://doi.org/10.1016/j.jacc.2022.09.048>
- Chen, Y.-C., Prabhu, K. S., & Mastro, A. M. (2013). Is selenium a potential treatment for cancer metastasis? *Nutrients*, 5(4), 1149–1168. <https://doi.org/10.3390/nu5041149>

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