

COVID-19 and Cardiovascular Disease in the Global Chronic Disease Epidemic

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Letter to Editor

The COVID-19 epidemic and global chronic disease epidemic is expected to cost billions of dollars in the next 20 years. The role of various chronic diseases such as NAFLD, diabetes, cardiovascular disease and neurodegenerative disease research may now be relevant to the COVID-19 pandemic with the anti-aging gene repression connected to mitophagy [1,2] and the severity of the COVID-19 and heart disease. The role of critical anti-aging genes such as Sirtuin 1 (Sirt 1) have attracted interest in cardiovascular disease with a critical role of Sirt 1 in the determination of cell death and survival involved with the severity of cardiovascular disease [3,4]. Sirt 1 is a NAD(+) dependent class III histone deacetylase (HDAC) protein involved in transcriptional regulation to determine gene expression with relevance to insulin resistance and various chronic diseases. Research studies now have reported an association between COVID-19 and cardiovascular disease [5]. Interests in COVID-19 and Sirt 1 have accelerated since Sirt 1 repression may determine the effects on cellular gene expression with effects on mitophagy and cardiovascular disease. Mitophagy and cardiovascular disease are closely connected and are of major relevance to cardiology and cardiac surgery [6,7]. The role of Sirt 1 on mitochondrial survival is now important to cardiovascular disease and the effects on cardiac function and therapy. The effect of COVID-19 on inactivation of Sirt 1 [8] may lead to mitophagy and programmed cell death with relevance to myocardial infarction and ischemic heart disease. Individuals from the global chronic disease epidemic [9-11] with COVID-19 may now be extremely susceptible to cardiovascular disease and heart failure. The consumption of Sirt 1 activators [12,13] in COVID-19 heart disease individuals is critical to delay the severity of cardiovascular disease. Sirt 1 inhibitors should be avoided to improve cardiac function and rehabilitation. Low calorie diets, lifestyle changes and

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exercise will increase plasma Sirt 1 levels [14] to prevent chronic disease and cardiovascular disease in COVID-19 individuals.

References

1. Martins IJ. Anti-aging genes improve appetite regulation and reverse cell senescence and apoptosis in global populations. *Adv Aging Res.* 2016;5:9-26.
2. Martins IJ. Single gene inactivation with implications to diabetes and multiple organ dysfunction syndrome. *J Clin Epigenet.* 2017;3(24):2472-1158.
3. Chong ZZ, Wang S, Shang YC, Maiese K. Targeting cardiovascular disease with novel SIRT1 pathways. *Future Cardiol.* 2012;8(1):89-100. [PubMed](#) | [CrossRef](#)
4. Matsushima S, Sadoshima J. The role of sirtuins in cardiac disease. *Am J Physiol Heart Circ Physiol.* 2015;309(9):H1375-89. [PubMed](#) | [CrossRef](#)
5. Nishiga M, Wang DW, Han Y, Lewis DB, Wu JC. COVID-19 and cardiovascular disease: from basic mechanisms to clinical perspectives. *Nat Rev Cardiol.* 2020;17(9):543-58. [PubMed](#) | [CrossRef](#)
6. Morciano G, Patergnani S, Bonora M, Pedriali G, Tarocco A, Bouhamida E, et al. Mitophagy in cardiovascular diseases. *J Clin Med.* 2020;9(3):892. [PubMed](#) | [CrossRef](#)
7. Yang Y, Li T, Li Z, Liu N, Yan Y, Liu B. Role of mitophagy in cardiovascular disease. *Aging Dis.* 2020;11(2):419. [PubMed](#) | [CrossRef](#)
8. Martins IJ. COVID-19 infection and anti-aging gene inactivation. *Int J Mol Sci.* 2015;16:29554-73.
9. Martins IJ. Appetite Dysregulation and the Apelinergic System are Connected to the Global Chronic Disease Epidemic. *Series of Endocrinol, Diabetes and Metab.* 2020;1(3):67-9. [PubMed](#) | [CrossRef](#)
10. Martins I. Increased Risk for Obesity and Diabetes with Neurodegeneration in Developing Countries. Top 10 Contribution on Genetics. In *Increased Risk for Obesity and Diabetes with Neurodegeneration in Developing Countries. Top 10 Contribution on Genetics.* Avid Science. 2018:1-35.
11. Martins IJ. Insulin Therapy and Autoimmune Disease with Relevance to Non Alcoholic Fatty Liver Disease. In *Nonalcoholic Fatty Liver Disease-An Update.* IntechOpen. 2018.
12. Martins IJ. Nutrition therapy regulates caffeine metabolism with relevance to NAFLD and induction of type 3 diabetes. *J Diabetes Metab Disord.* 2017;4(1):1-9.
13. Dai H, Sinclair DA, Ellis JL, Steegborn C. Sirtuin activators and inhibitors: promises, achievements, and challenges. *Pharmacol Ther.* 2018;188:140-54. [PubMed](#) | [CrossRef](#)
14. Martins IJ. Sirtuin 1, a diagnostic protein marker and its relevance to chronic disease and therapeutic drug interventions. *EC Pharmacol Toxicol.* 2018;6(4):209-15.