## Should We Fear A Wave of Cancers After the COVID-19 Pandemic?

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ancer incidence is increasing globally. It is well documented that the incidence of most cancers increases with age [1]. Due to factors such as improved medical care, better hygiene, healthier life styles, sufficient food and decreased child mortality, human life expectancy is increasing at a rapid rate so that nowadays we can expect to live much longer than our ancestors who lived a few generations back [2]. Given these considerations, It is widely believed that increased life span is the main reason cancer risk overall is rising [3]. A paper published in the Lancet reports that delays in screening, diagnosis, and treatment due to the COVID-19 pandemic could lead to excess cancer deaths, and slow or even reverse the declining trend in mortality projected for some cancers [4].

Furthermore, the report by Harvard Medical School researchers at Dana-Farber Cancer Institute and colleagues from other institutions, suggests that COVID-19 has complicated the treatment for patients with cancer. "In patients with cancer, COVID-19 can be especially harsh. This is likely because many of these patients have a weakened immune system—either as a result of the cancer itself or the therapies used to treat it—and are therefore less able to fight off infection by the new coronavirus" [5].

In 2021, a research team led by Zhou highlighted the clinical and molecular similarities between cancer and COVID-19 and summarized the four major signaling pathways at the intersection of COVID-19 and cancer, namely, cytokine, type I interferon (IFN-I), androgen receptor (AR), and immune checkpoint signaling. They also discussed the advantages and disadvantages of repurposing anticancer treatment for the treatment of COVID-19 [6].

However, Professor Abdollah Jafarzadeh and his research team, in their paper "Review SARS-CoV-2 Infection: A Possible Risk Factor for Incidence and Recurrence of Cancers" explored another aspect of the interplays of COVID and cancer. Their findings which are published in the International Journal of Hematology-Oncology and Stem Cell Research might shed some light on the dark corners of the potential interactions of COVID-19 and cancer development [7].

Jafarzadeh et al. reported that the patients with some types of cancers may be more vulnerable to SARS-CoV-2 infection compared with the

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Received: 30 October 2023 Accepted: 10 November 2023 non-cancerous individuals, due to their immunocompromised state resulted from malignancy, chemotherapy, and other concomitant abnormalities as well as perhaps greater expression of angiotensin-converting enzyme 2. Moreover, they reported that clinically recovered COVID-19 individuals display immune abnormalities that persist several months after discharge [7].

The lymphopenia-related immunosuppression, functional exhaustion of cytotoxic lymphocytes (such as CD8+ cytotoxic T-cells and natural killer cells), hyperinflammatory responses, oxidative stress, downregulation of interferon response, development of the myeloid-derived suppressor cells, downregulation of tumor suppressor proteins and perhaps reactivation of the latent oncogenic viruses may directly and/or indirectly play a role in the cancer development and recurrence in severe COVID-19 [7].

## Conflict of Interest

None

## References

- 1. White MC, Holman DM, Boehm JE, Peipins LA, Grossman M, Jane Henley S. Age and Cancer Risk: A Potentially Modifiable Relationship. *American Journal of Preventive Medicine*. 2014;**46**(3):S7-15. doi: 10.1016/j. amepre.2013.10.029.
- 2. Brown GC. Living too long: The current focus of medical research on increasing the quantity, rather than the quality, of life is damaging our health and harming the economy. *EMBO Reports*. 2015;**16**(2):137-41. doi: 10.15252/embr.201439518.
- 3. Gu X, Zheng R, Xia C, Zeng H, Zhang S, Zou X, et al. Interactions between life expectancy and the incidence and mortality rates of cancer in China: a population-based cluster analysis. *Cancer Commun (Lond)*. 2018;**38**(1):44. doi: 10.1186/s40880-018-0308-x. PubMed PMID: 29970165. PubMed PMCID: PMC6029078.
- 4. Wells CR, Galvani AP. Impact of the COVID-19 pandemic on cancer incidence and mortality. *Lancet Public Health*. 2022;**7**(6):e490-1. doi: 10.1016/S2468-2667(22)00111-6. PubMed PMID: 35660207. PubMed PMCID: PMC9159732.
- 5. Dana-Farber Communications. A Covid-19/Cancer Interplay, Study Offers Global Review of Covid-19's Impact on Cancer Treatment And Research. Dana-Farber Communications; 2020.
- 6. Zong Z, Wei Y, Ren J, Zhang L, Zhou F. The intersection of COVID-19 and cancer: signaling pathways and treatment implications. *Mol Cancer*. 2021;**20**(1):76. doi: 10.1186/s12943-021-01363-1. PubMed PMID: 34001144. PubMed PMCID: PMC8126512.
- 7. Jafarzadeh A, Gosain R, Mortazavi SMJ, Nemati M, Jafarzadeh S, Ghaderi A. SARS-CoV-2 Infection: A Possible Risk Factor for Incidence and Recurrence of Cancers. *Int J Hematol Oncol Stem Cell Res.* 2022;**16**(2):117-27. doi: 10.18502/ijhoscr.v16i2.9205. PubMed PMID: 36304732. PubMed PMCID: PMC9547773.