

Two-Drug Immunotherapy Shows Benefit in Some Patients With Lung Cancer

Adding radiation therapy to Imfinzi and tremelimumab didn't help those with non-small cell lung cancer, according to a new study.

January 26, 2022 By Dana-Farber Cancer Institute

A combination of two drugs that open the floodgates to an immune system attack on cancer curtailed tumor growth in some patients with non-small cell lung cancer (NSCLC) that was resistant to a single immunotherapy agent, results from a recent clinical trial show. The addition of radiation therapy to the two-drug regimen did not improve outcomes, however, researchers at Dana-Farber Brigham Cancer Center and other centers report in a new study.

The paper, published online today by [The Lancet Oncology](#), is part of a broad-based effort to extend the benefits of drugs known as immune checkpoint inhibitors by combining them with other treatments and with one another. The new trial is one of the first to test radiation therapy in combination with immune therapy in patients with cancer. The fact that radiation therapy didn't enhance the effect of the two-drug therapy suggests that future studies should explore different combinations, the study authors say.

"Patients with NSCLC that doesn't respond to checkpoint inhibitors known as PD(L)-1 inhibitors have few treatment options. In this trial, we tested a PD-L1 inhibitor in combination with a drug that targets a different immune checkpoint protein, CTLA-4 — an approach that has shown promise in laboratory research but hasn't been extensively tested in patients whose NSCLC is resistant to PD(L)-1 inhibitors," says study lead author [Jonathan Schoenfeld, MD, MPH](#), of Dana-Farber Brigham Cancer Center.

"We not only found that the combination brought the cancer under control in some patients, but our laboratory work suggests a way of determining which patients are likely to have this benefit," he remarked.

The trial is part of a wave of research into combinations of checkpoint inhibitors and other therapies in patients with cancer. The inhibitors work by stifling "checkpoint" proteins on cancer cells or immune system cells that would otherwise protect tumors from an immune system attack. Such drugs have proved highly effective against certain types of cancer, but they've been less

successful, on their own, in many other types.

One promising solution has been to use checkpoint inhibitors in conjunction with radiation therapy. A phase 2 clinical trial of that combination in patients with metastatic NSCLC produced encouraging results, but it wasn't clear what dose of radiation works best.

The new trial set out to answer that question in patients with NSCLC resistant to PD(L)-1 inhibitors, and to evaluate a combination of PD-L1 and CTLA-4 inhibitors, with and without radiation, in these patients.

The trial involved 78 patients at 18 cancer centers in the U.S. All participants received the drugs durvalumab [Imfinzi] (which targets the PD-L1 checkpoint protein on tumor cells) and tremelimumab [not yet approved] (which targets CTLA-4). One-third of them were also treated with low-dose radiation therapy followed by durvalumab. And one-third were treated with higher-dose, "hypofractionated" radiation therapy followed by durvalumab.

There was no difference in response rates between patients treated with radiation therapy and those in the immunotherapy-only group, investigators found. At the same time, though, about 30% of the patients had their disease brought under control, including 10% who had long-lasting responses to treatment — an impressive outcome considering that all participants had cancers that were resistant to PD(L)-1 inhibitors.

As part of the study, researchers analyzed tumor tissue samples from the participants. They found that those whose tumors teemed with immune system T cells were more likely to respond to the treatment than those with fewer T cells in their tumor tissue.

"Our findings suggest that durvalumab/tremelimumab should be further explored for its ability to benefit NSCLC patients who have progressed on prior PD(L)-1 therapy," Schoenfeld said. "In future trials of these and similar agents, analyzing tumor tissue for T-cell infiltration may be able to identify patients who can most benefit from this type of treatment."

The senior authors of the paper are Catherine J. Wu, MD, and F. Stephen Hodi, MD, of Dana-Farber; Helen X. Chen, MD, of the National Cancer Institute; and Arta M. Monjazeb, MD, of the University of California Davis Comprehensive Cancer Center. Co-authors are: Anita Giobbie-Hurder, MS, Yang Liu, PhD, Aashna Jhaveri, MS, Lin Yang, MS, Jennifer Altreuter, MD, Jason L. Weirather, PhD, Srinika Ranasinghe, PhD, Katrina Z. Kao, Ana Lako, PhD, Ryan C. Brennick, Scott J. Rodig, MD, and Mark M. Awad, MD, of Dana-Farber; Lauren Gunasti and Raymond H. Mak, MD, of Dana-Farber Brigham Cancer Center; Junko Tsuji, PhD, Hayley Lyon, Carrie Cibuskis, and Niall Lennon, PhD, of the Broad Institute of MIT and Harvard; Ryan D Gantzler, MD, of the University of Virginia Cancer Center; Carrie Lee, MD, of the University of North Carolina Lineberger Comprehensive Cancer Center; Joleen Hubbard, MD, of the Mayo Clinic; Susanne M. Arnold, MD, of the University of Kentucky Markey Cancer Center; James L. Abbruzzese, MD, of Duke Cancer Institute; Salma K. Jabbour, MD, of Rutgers Cancer Institute of New Jersey; Nataliya V. Uboha, MD, of the University of Wisconsin Carbone Cancer Center; Kevin L. Stephans, MD, of the Cleveland Clinic; Jennifer M Johnson, MD, of Sidney Kimmel Cancer Center; Haeseong Park, MD, of Siteman Cancer Center, Washington

University, St. Louis, Mo; Liza C. Villaruz, MD, of the University of Pittsburgh Medical Center Hillman Cancer Center; and Elad Sharon, MD, Howard Streicher, MD, and Mansoor M Ahmed, PhD, of the National Cancer Institute.

The study was funded by the National Institutes of Health and Dana-Farber, including grant UM1 CA186709, a Biomarker Supplement to UM1 CA186709. Funding also included grants U24CA224331 and U24CA224316.

[This article](#) was originally published on January 13, 2022, by Dana-Farber Cancer Institute. It is republished with permission.

© 2026 Smart + Strong All Rights Reserved.

<http://beta.docker.cancerhealth.com/article/twodrug-immunotherapy-shows-benefit-patients-lung-cancer>