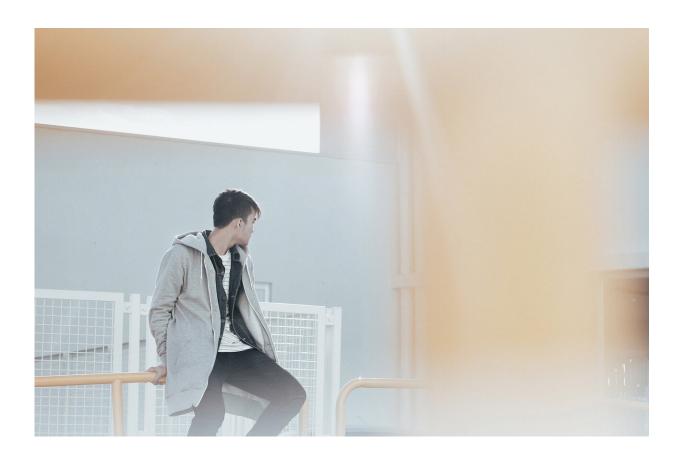


## **Study confirms racial differences in response** to prostate cancer treatment

May 6 2021, by Sarah Avery



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A study designed to enroll an equal number of Black and white men with advanced prostate cancer confirms key findings that have been evident in retrospective analyses and suggest potential new avenues for treating



Black patients who disproportionately die of the disease.

Researchers at Duke Cancer Institute enrolled 50 Black and 50 <u>white</u> <u>men</u> with advanced <u>prostate cancer</u> to test whether there were outcome differences on treatment with the hormone therapy abiraterone acetate plus the steroid prednisone. In retrospective data reviews, the Duke researchers had previously found <u>racial differences</u> in PSA responses among advanced <u>prostate</u> cancer patients.

Publishing online in the journal *Cancer*, the researchers confirmed trends indicating that Black men's PSA levels dropped further and more frequently than those of white men undergoing the therapy. These PSA changes, however, did not result in differences in disease progression or overall survival times.

But the survival finding has an important subtlety, said lead author Daniel George, M.D., professor in the departments of Medicine and Surgery at Duke University School of Medicine. George noted that most drug studies among prostate cancer patients include a small fraction of Black men that is far lower than their numbers in the larger population.

Exclusions typically result because Black men with prostate cancer are more likely to have other illnesses such as diabetes and <u>high blood</u> <u>pressure</u>, which study leaders often fear could put them at higher risk for complications. Additionally, there are deep historic and cultural reasons that Black men tend to decline participation in <u>clinical studies</u>.

For their study, however, the Duke team—including senior author Andrew Armstrong, M.D., professor in the departments of Medicine and Surgery—found that Black men were eager to join the clinical trial.

They were able to enroll a much larger proportion of Black men than what most studies include, in part because the study was addressing a



question pertaining to race. And they did not exclude men with coexisting conditions, asserting that since the treatment is FDA approved for this population, they should be inclusive of the patients they see in practice.

"When you look at the overall survival data for our study, they're equal between Black and white men," George said. "But given the prevalence of coexisting conditions in the Black men we enrolled, mortality should have actually been higher for them.

"Our finding that it was not higher is telling—it suggests Black men with prostate cancer can fare just as well as whites, even with other health issues," George said. "And it signals that future studies should consider enrolling Black men despite these often-disqualifying conditions."

George said the researchers also identified a possible marker of ancestrydependent treatment outcomes that could help explain why Black men respond more readily to hormone therapy, potentially pointing to new ways to address <u>advanced prostate cancer</u> in Black men.

"We need to understand how genetic ancestry might affect treatment outcomes—especially disease responsiveness in prostate <u>cancer</u> —because we are now using and studying these therapies earlier in the disease where we have the opportunity to cure patients," George said.

"If there is a subgroup of patients with an ancestry-based predisposition for potential better response, we need to understand that. But to do so, we will need greater genetic diversity in our future study populations, especially among those with African ancestry. We aren't going to fully understand this genetic complexity by solely enrolling men with European ancestry."

More information: Daniel J. George et al. A prospective trial of



abiraterone acetate plus prednisone in Black and White men with metastatic castrate-resistant prostate cancer, *Cancer* (2021). <u>DOI:</u> 10.1002/cncr.33589

## Provided by Duke University School of Nursing

Citation: Study confirms racial differences in response to prostate cancer treatment (2021, May 6) retrieved 22 November 2023 from <u>https://medicalxpress.com/news/2021-05-racial-differences-response-prostate-cancer.html</u>

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