

A new screening method for prostate cancer

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A new study by NYU Langone Medical Center and Northwestern University Feinberg School of Medicine shows novel PSA velocity (PSAV) risk count testing may provide a more effective way for physicians to screen men for clinically significant prostate cancer. The new study, published online by the *British Journal of Urology International* on February 1, 2012, shows the benefits of tracking a man's PSA levels over time to help doctors more accurately assess his risk of life-threatening prostate cancer.

"Risk count could represent a new way to screen for [prostate cancer](#) by focusing on men with the greatest risk of harmful prostate cancers," said lead author Stacy Loeb, MD, an urologist in the Department of Urology and the Joel E. Smilow Comprehensive Prostate Cancer Center at NYU Langone. "The goal of risk count is to help identify the aggressive, clinically significant prostate cancers before advanced symptoms develop, while decreasing the diagnosis of insignificant cancers."

Prostate cancer is the second leading cause of [cancer death](#) in American men, with an estimated 1 in 6 men diagnosed with the disease during their lifetime. Prostate cancer does not present symptoms until advanced stages so screening for the disease is vital. Currently, a [PSA blood test](#) is the standard screening method to evaluate a man's risk of prostate cancer. It measures the amount of prostate-specific antigen in the blood, a substance made only in the prostate gland. An elevated PSA can indicate the presence of disease. However, PSA can also be elevated with [benign prostate enlargement](#) and one high PSA value does not always mean an aggressive prostate cancer is present.

The new PSAV risk count screening works by monitoring fluctuations in PSA levels over time to analyze a man's risk of prostate cancer, instead of relying on just one [PSA test](#) result to assign [prostate cancer risk](#). The risk count is calculated by counting the number of times in a row that the PSA level in the blood increases by 0.4 ng/mL. If PSA goes up by more than 0.4 units multiple years in a row, the risk count rises indicating the patient has an increased risk of aggressive prostate cancer. For example, a man who has a PSA screening for two years in a row would be given a "2" risk count if his serial PSA velocity measurements increased by more than 0.4 units, a "1" risk count if there was only one increase by more than 0.4 units, and a "0" risk count if there was no increase by more than 0.4 units.

In the study, researchers showed PSAV risk count could improve the specificity of screening for prostate cancer and advanced stages of the disease. Researchers evaluated 18, 214 men undergoing prostate cancer screening, 1,125 of which were diagnosed with the disease. The study results show sustained rises in [PSA levels](#) over time indicate a significantly greater risk of prostate cancer and more aggressive disease. In the study, a risk count of "2" was associated with a greater than 8-times risk of prostate cancer and a 5-fold greater risk of aggressive disease. The authors conclude risk count screening may be useful in diagnosing aggressive prostate cancer earlier while possibly reducing unnecessary biopsy, as well as the overdiagnosis and resulting overtreatment of low-risk prostate cancer.

"A persistently rising PSA is a harbinger for life-threatening prostate cancer," said the study's senior author, William Catalona, MD, professor of Urology at Northwestern University. "Our study findings show looking at how much PSA changes over time helps distinguish which cancers are aggressive more so than a single PSA value."

In an accompanying editorial, Dr. H. Ballentine Carter from the Brady

Urological Institute at Johns Hopkins who initially suggested the concept of looking at PSA changes over time, affirms that in order to determine the likelihood of aggressive prostate cancer, "you want to know your patient's risk count, not just their age and PSA level."

Provided by New York University School of Medicine

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