

Delaying HIV Treatment Linked to Stiffened Blood Vessels

This finding raises concerns that staying off antiretrovirals may raise the risk of more serious cardiovascular problems.

October 29, 2019 By [Benjamin Ryan](#)

Starting antiretroviral (ARV) treatment for HIV is associated with a reduction in endothelial dysfunction, a driver of more serious cardiovascular disease (CVD). Given this new finding, researchers are concerned that delaying HIV treatment initiation may raise the risk of CVD.

Endothelial dysfunction occurs when the layer of cells lining blood vessels stiffen, impeding the widening of such vessels that increases blood flow and lowers blood pressure. Traditionally, endothelial dysfunction is associated with cardiovascular risk factors such as diabetes, high blood pressure, irregular blood lipid levels and smoking.

A research team led by Kelvin N. V. Bush, MD, of the division of cardiology at San Antonio Military Medical Center, sought to analyze endothelial dysfunction's association with HIV and ARV treatment. Publishing their findings in JAMA Network Open, they conducted a cohort study of 61 members of the U.S. Air Force who were diagnosed with HIV between September 2015 and September 2017.

At the study's outset, the participants were evaluated for endothelial dysfunction using a test that expresses results in what's known as natural log-transformed RHI values (lnRHI). A value of less than 0.51 is considered an abnormal level and a value of at least 0.51 is considered normal.

Ninety-five percent of the participants were male, and 57% were Black. The average age at HIV diagnosis was 28.1 years old. The median time since the estimated date of the participants' acquisition of HIV was an estimated 10.6 months. The median CD4 count was 552.

The participants had an average body mass index of 26.2 (25 to 29.9 is considered overweight; 30 and over is considered obese), a median LDL cholesterol of 97 and a median total cholesterol of

163. None had diabetes.

The average InRHI when the participants were diagnosed was 0.70—in the normal range. At the study's outset, 77% of the participants had a normal baseline InRHI result, while 23% had an abnormal result.

At the study's outset, every 10 years of increased age was associated with about double the odds of having an abnormal InRHI level, but this finding was not statistically significant, meaning it may have been driven by chance.

Of the 41 participants who had follow-up InRHI assessments, 40 started ARVs immediately after joining the study. The group had repeated assessments in intervals of a median of 6.4 months.

HIV treatment was associated with a significant increase in the mean InRHI level. The 11 individuals who started the study with an abnormal InRHI level experienced a greater average increase compared with the 29 people who began with a normal level. Among those with an abnormal initial InRHI level, 73% experienced improved endothelial function after starting ARVs.

The one person who did not go on HIV treatment went from having a normal InRHI level of 0.60 to an abnormal level of 0.11 after 8.3 months.

“Endothelial dysfunction identified in clinical practice is managed with modification of cardiovascular risk factors, including weight loss, exercise prescriptions, [ACE] inhibitors [high blood pressure medication], lowering of [LDL] levels with statins, hormonal therapies, antioxidants and smoking cessation,” the study authors noted.

“In our study,” they continued, “none of the patients received specific interventions beyond education and counseling. The patients in our study did not have significant changes in their body mass index, and no exercise prescriptions were ordered to suggest that changes in physical activity influenced the interval InRHI changes. This leads us to conclude that ART initiation was significantly associated with the reversal of endothelial dysfunction observed after approximately six months of therapy.”

To read the study, [click here](#).

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