CLINICAL GUIDELINES PROGRAM

**NEW YORK STATE DEPARTMENT OF HEALTH AIDS INSTITUTE** | HIV · HCV · SUBSTANCE USE · LGBT HEALTH

## Virologic and Immunologic Monitoring in HIV Care

June 2022

Event	HIV RNA Viral Load	CD4 Count	Comments
Entry into care	Baseline viral load (A1)	Baseline CD4 count (A1)	<ul> <li>If a patient is not taking ART, recommend initiation [b] (A1)</li> <li>Monitor as below</li> </ul>
Patients Taking ART			
ART initiation or change to address virologic failure	<ul> <li>Within 4 weeks after ART start or change (A3)</li> <li>At least every 8 weeks until complete virologic suppression is documented (A3)</li> </ul>	<ul> <li>12 weeks after ART initiation</li> <li>Every 4 months until CD4 count &gt;200 cells/mm<sup>3</sup> is obtained on 2 measurements at least 4 months apart (A2), then monitor as below once virologic suppression is achieved</li> </ul>	<ul> <li>Virologic failure occurs when a viral load</li> <li>&lt;200 copies/mL is either not achieved or not maintained</li> <li>Virologic suppression is defined as a viral load</li> <li>&lt;20 to &lt;50 copies/mL obtained with a highly sensitive assay</li> </ul>
ART change for simplification or due to adverse effects	Within 4 weeks after ART change, then as below (A3)	Monitor as below for documented virologic suppression	_
Documented viral suppression	<ul> <li>At least every 4 months (A3)</li> <li>May extend interval to 6 months in patients stable on ART with CD4 count &gt;200 cells/mm<sup>3</sup> and complete viral suppression for 1 year (B2)</li> </ul>	<ul> <li>At least every 6 months if CD4 count is ≤350 cells/mm<sup>3</sup> (B2)</li> <li>Optional if CD4 count is &gt;350 cells/mm<sup>3</sup> (B2)</li> </ul>	_
New HIV RNA ≥500 copies/mL after previous viral suppression	Repeat viral load test 2 weeks after first result (A2)	Obtain CD4 count if previous result is >6 months old (B3)	<ul> <li>Assess for adherence and drug-drug interactions (A3)</li> <li>Obtain resistance testing (A1)</li> </ul>
New HIV RNA level over the limit of detection of sensitive assays, 20 to 50 copies/mL, but <500 copies/mL after previous viral suppression	Repeat viral load test within 4 weeks to differentiate low-level transient viremia ("blip") from virologic failure [c] (A2)	If repeat viral load is detectable, obtain CD4 count if previous result is >6 months old (B3)	<ul> <li>Assess for adherence and drug-drug interactions (A3)</li> <li>If repeat viral load is detectable, consider resistance testing [d] (B3)</li> <li>Patients with low-level viremia ≤200 copies/mL over a period of 12 months without demonstrated failure may continue routine testing intervals of at least every 4 months [e]</li> </ul>

Available at: hivguidelines.org/antiretroviral-therapy/cd4-and-viral-load-monitoring/#tab\_1

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Table 1: Recommended Viral Load and CD4 Count Monitoring in Nonpregnant Patients With HIV [a]					
Event	HIV RNA Viral Load	CD4 Count	Comments		
Patients Not Taking ART					
CD4 count ≤500 cells/mm <sup>3</sup> (A2)	At least every 4 months	At least every 4 months	At every visit, recommend ART initiation [b]		
CD4 count >500 cells/mm <sup>3</sup> (A2)	At least every 6 months	At least every 6 months	At every visit, recommend ART initiation [b]		
Abbreviations ART, antiretroviral therapy.					
Notes:					
<ul> <li>a. For recommendations on virologic monitoring in pregnancy, see DHHS: <u>Recommendations for the Use of Antiretroviral Drugs</u> <u>During Pregnancy and Interventions to Reduce Perinatal HIV Transmission in the United States</u>.</li> <li>b. See the NYSDOH AI guideline <u>When to Initiate ART</u>, <u>With Protocol for Rapid Initiation</u>.</li> <li>c. An ART regimen should not be changed based on a single viral load elevation. The risk of virologic rebound (breakthrough) increases when values are ≥500 copies/mL [Grennan, et al. 2012].</li> <li>d. Standard enerthrough text and the provide resistance results when viral load is law. For energed low level virance are essentiated.</li> </ul>					
<ul> <li>d. Standard genotypic tests may not provide resistance results when viral load is low. For repeated low-level viremia, an assay that detects resistance mutations in archived proviral DNA is available; however, clinical data are insufficient to recommend for or against its use in the patient care setting.</li> <li>e. In patients with low-level viremia, clinicians should consult with an <u>experienced HIV care provider</u>; low-level viremia can be due to multiple causes, and its clinical effect is not clear.</li> </ul>					

## Reference

Grennan JT, Loutfy MR, Su D, et al. Magnitude of virologic blips is associated with a higher risk for virologic rebound in HIV-infected individuals: a recurrent events analysis. *J Infect Dis* 2012;205(8):1230-1238. [PMID: 22438396] https://pubmed.ncbi.nlm.nih.gov/22438396