## ALL RECOMMENDATIONS *P.3*

### PRESENTATION AND DIAGNOSIS OF IRIS

- Clinicians should include IRIS as part of the differential diagnosis when inflammatory signs or symptoms occur following recent initiation of, re-initiation of, or a change to an ART regimen. (A3)
- In assessing patients for IRIS, clinicians should exclude HIV disease progression, new infections, and drug reactions as underlying causes for inflammatory signs or symptoms. (A3)

### MANAGEMENT AND TREATMENT OF IRIS

- Clinicians should initiate appropriate treatment of opportunistic infections (OIs), as well as symptomatic treatment and supportive care according to the severity of IRIS. (A3)
- Clinicians should not interrupt ART except in severe, life-threatening cases of IRIS. (A3)

### Severe IRIS

- Clinicians should consult with an experienced HIV care provider for the management of severe IRIS, including the decision of whether to interrupt ART if IRIS is severe. (A3)
- Clinicians should treat patients with severe IRIS that is not caused by either cryptococcal meningitis or Kaposi's sarcoma (KS) with 1 to 2 mg/kg prednisone, or the equivalent, for 1 to 2 weeks, followed by a period of tapering dose that is individualized. (B3)
- Clinicians should not use corticosteroids for management of cryptococcal meningitis or in patients with KS. (A2)
- Clinicians should closely monitor patients receiving corticosteroids for the development of OIs, including CMV retinitis and TB disease. (A3)

**Turn over for Summary of Recommended Timing of ART Initiation and Major and Minor Presentations of IRIS →**

# HIV CLINICAL RESOURCE ¼-FOLDED GUIDE

VISIT **HIVGUIDELINES.ORG** TO LEARN MORE OR VIEW COMPLETE GUIDE

# MANAGEMENT OF IMMUNE RECONSTITUTION INFLAMMATORY SYNDROME (IRIS)

**NYSDOH AIDS INSTITUTE PrEP CLINICAL GUIDELINE** JUNE 2017

## → KEY POINTS

- **ART should not be interrupted in patients with IRIS except in life-threatening cases, usually associated with CNS-IRIS, in which corticosteroids did not result in improvement.**
- Steroids should not be used routinely as induction therapy in treatment of cryptococcal IRIS.
- Steroids are not effective in reducing intracranial pressure.
- Before initiating ART in patients who have TB meningitis, extrapulmonary TB, CMV retinitis, or cryptococcal infection, clinicians should consult with a care provider who is experienced in managing patients with HIV infection in the setting of active OIs.
- **Finding an experienced HIV care provider:** The Clinical Education Initiative (CEI) line, which is available through the New York State Department of Health, provides access to providers with experience in managing all aspects of HIV infection. **Call 866-637-2342.**

**← Use this code with your phone's QR code reader to go directly to a mobile-friendly version of the guideline.**

**This ¼-Folded Guide is a companion to the New York State Department of Health AIDS Institute guideline *Management and Treatment of IRIS*. The full guideline is available at www.hivguidelines.org.**

**ALL RECOMMENDATIONS** *P.1*

**TIMING OF ART INITIATION IN PATIENTS WITH RECENT OIs AND PREVENTION OF IRIS**

- Clinicians should recommend that patients initiate antiretroviral therapy (ART) within 2 weeks of beginning treatment for active opportunistic infections (OIs), with exceptions to this recommendation noted below. (A1)
- Clinicians should consult with a provider experienced in managing HIV in the setting of active OIs to determine when to initiate ART in patients with tuberculosis (TB) meningitis, extrapulmonary TB, cytomegalovirus (CMV) retinitis, or cryptococcal infection. (A3)
- For patients with CD4 counts <100 cells/mm$^3$ or known concomitant OIs who are initiating ART, clinicians should be vigilant for the signs and symptoms of IRIS and should educate patients about the risk of developing IRIS. (A3)
- For HIV-infected patients who are co-infected with hepatitis B virus (HBV) or hepatitis C virus (HCV), clinicians should:
  - Measure transaminase levels before initiation of ART, at 6 and 12 weeks after initiation, and at least every 6 months thereafter to monitor for possible IRIS. (A3)
  - Refer patients with elevated transaminase levels in conjunction with jaundice, elevated bilirubin levels, or loss of synthetic function for evaluation by a hepatologist. (B3)

**Pulmonary TB**

- For patients with pulmonary TB, clinicians should initiate ART as follows:
  - CD4 counts ≥50 cells/mm$^3$: As soon as patients are clinically stable on anti-TB therapy and no later than 12 weeks after initiating anti-TB therapy. (A1)
  - CD4 counts <50 cells/mm$^3$: Within the first 2 weeks after initiating anti-TB therapy. (A1)
- For patients with pulmonary TB who are ART-naïve, who have a CD4 count <100 cells/mm$^3$, and who started on anti-TB treatment within the last 30 days, clinicians should initiate prednisone 40 mg daily for 14 days, followed by 20 mg daily for 14 days at the time of ART initiation. (B1)

**ALL RECOMMENDATIONS** *P.2*

**TB Meningitis and Extrapulmonary TB**

- For patients with TB meningitis or extrapulmonary TB, clinicians should consult with an experienced HIV care provider to determine the timing of ART initiation. (A3)

**Cryptococcal Meningitis**

- Clinicians should treat ART-naive patients diagnosed with cryptococcal meningitis with standard antifungal therapy and should:
  - Delay ART initiation until the patient has completed at least 2 weeks of antifungal treatment. (A1)
  - Consult with an experienced HIV care provider to determine optimal timing for ART initiation. (A3)
- If the patient initiates ART before completing 10 weeks of antifungal therapy, the clinician should monitor closely for intracranial pressure and other signs and symptoms of IRIS and manage intracranial pressure aggressively. (A2)
- For patients with other types of cryptococcal infection (not meningitis), clinicians should consult with an experienced HIV care provider to determine the timing of ART initiation. (A3)

**CMV Retinitis**

- Clinicians should ensure that HIV-infected patients with CD4 counts <100 cells/mm$^3$ receive dilated ophthalmologic examination to assess for signs of CMV before initiation of ART. (A2)
- Clinicians should not initiate ART immediately in patients with CMV retinitis (A2) but should consult with an experienced HIV care provider to determine the timing of ART initiation. (A3)
- Clinicians should ensure that after initiation of ART, HIV-infected patients with a history of CMV retinitis are monitored by dilated ophthalmologic examination to assess for possible IRIS as follows:
  - Every 3 months for the first year after initiation of ART. (A3)
  - Immediately if there is a change in visual acuity or development of floaters. (A2)

## SUMMARY OF RECOMMENDED TIMING OF ART INITIATION

| Opportunistic Infection (OI) | Timing of ART Initiation after Starting OI Treatment |
|---|---|
| **Cryptosporidiosis**<br>**Microsporidiosis**<br>**Progressive multifocal leukoencephalopathy (PML)**<br>**Kaposi's sarcoma (KS)**<br>**Pneumocystis jiroveci pneumonia (formerly PCP)**<br>**Hepatitis B virus (HBV) infection**<br>**Hepatitis C virus (HCV) infection**<br>**Pulmonary tuberculosis (TB)**<br>**Other serious bacterial infections** | · Within 2 weeks of starting treatment for an OI or as soon as the patient is clinically stable |
| **Pulmonary TB** | · CD4 count ≥50 cells/mm$^3$: Initiate ART as soon as the patient is clinically stable after initiating TB therapy, but no more than 12 weeks later<br>· CD4 count <50 cells/mm$^3$: Initiate ART within the first 2 weeks after initiating TB therapy |
| **Extrapulmonary TB** | · Optimal timing has not been established; consult with an experienced HIV care provider |
| **TB meningitis** | · Optimal timing has not been established; consult with an experienced HIV care provider |
| **Cryptococcal meningitis** | · Delay 2 to 10 weeks after starting antifungal therapy<br>· Optimal timing has not been established; consult with an experienced HIV care provider |
| **Cryptococcal infection other than meningitis** | · Delay at least 2 weeks after starting antifungal therapy<br>· Optimal timing has not been established; consult with an experienced HIV care provider |
| **CMV retinitis** | · Immediate ART is *not recommended*<br>· Optimal timing has not been established; consult with an experienced HIV care provider |

## MAJOR AND MINOR PRESENTATIONS OF IMMUNE RECONSTITUTION INFLAMMATORY SYNDROME (IRIS)

| Opportunistic Infection (OI) | IRIS Signs/Symptoms |
|---|---|
| **Major Presentations** | |
| **Tuberculosis (TB)** | · Patients responding to TB treatment may have worsening of pulmonary symptoms, X-ray findings that suggest worsening of TB disease, enlarging lymph nodes causing airway obstruction, or meningeal symptoms<br>· Enlarging tuberculoma or pericardial effusions have been described<br>· TB-IRIS can also result in hepatotoxicity, which may be difficult to distinguish from medication-induced toxicity<br>· TB-IRIS may occur in patients with undiagnosed multidrug-resistant TB |
| ***Mycobacterium avium* complex (MAC)** | · May present as pulmonary disease or systemic inflammation that is indistinguishable from active MAC<br>· Atypical presentations, such as localized lymphadenitis or endobronchial mass lesions, may occur; osteomyelitis is an atypical late manifestation<br>· Patients with MAC-IRIS may not be bacteremic and may have no known history of MAC diagnosis |
| **Cryptococcal meningitis** | · Usually presents as worsening of meningitis symptoms, including possible rapid hearing and/or vision loss, ataxia, and/or elevated intracranial pressure |
| **Cytomegalovirus (CMV) retinitis** | · Presents as retinitis, vitritis, or uveitis (variable timing, with median time to immune reconstitution vitritis 20 weeks after ART initiation in one study):<br>– Retinitis is inflammation that is usually at the site of previous CMV retinitis lesions<br>– Uveitis and vitritis are the presence of inflammatory cells in the eye as a result of IRIS and may help to distinguish IRIS from active CMV retinitis<br>· CMV-IRIS in the eye can cause rapid and permanent vision loss |
| **Hepatitis B or C virus** | · Transient elevations in transaminases may occur after initiation of ART with immune reconstitution and can be difficult to distinguish from drug-induced hepatitis<br>· Hepatic flares are usually mild and self-limited but can result in decompensation in someone with preexisting cirrhosis |
| **Progressive multifocal leukoencephalopathy (PML)** | · PML lesions may be unmasked or worsen and could appear as new or worsening focal neurologic deficits or lesions on MRI |
| **Kaposi's sarcoma (KS)** | · Presents as worsening of KS<br>· Cutaneous lesions are the most common presentation; other signs include lymphedema and oral, gastric, lung, genital, or conjunctival lesions<br>· Fatal cases of KS-IRIS have been reported |
| **Cerebral toxoplasmosis** | · May present as cerebral abscess (also known as toxoplasmosis encephalitis) or, rarely, diffuse encephalitis or chorioretinitis |
| **Autoimmune diseases** | · Preexisting sarcoidosis may be exacerbated<br>· Late presentations of Grave's disease have been reported 8 to 33 months after ART initiation |
| **Minor Presentations** | |
| **Herpes simplex virus (HSV) and varicella zoster virus (VZV)** | · HSV and VZV can reactivate after initiation of ART, even in patients without previously diagnosed disease<br>· Presentations are usually similar to non-IRIS disease; however, IRIS may worsen a patient's symptoms |
| **Nonspecific dermatologic** | · A number of dermatologic manifestations, such as folliculitis and oral and genital warts, may appear or worsen during immune reconstitution |