

Diagnostic and Treatment Dilemmas in Polymyalgia Rheumatica (PMR)

PMR is a diagnosis made by observing typical symptoms, excluding other diagnoses, and confirming a response to glucocorticoid (GC) therapy¹⁻³



1 Presence of acute, intense pain and stiffness



2 Exclusion of mimicking disorders



3 Observation of rapid improvement with GC therapy

1 PMR symptoms are nonspecific and can present in a variety of other conditions



Demographics^{4,5}

- ≥50 years
- Female



Pain and stiffness^{4,5}

- Bilateral shoulder pain and tenderness
- Pelvic girdle pain
- Restricted range of motion in affected muscles



Physical and mental^{5,6}

- Fever
- Fatigue, malaise
- Loss of appetite
- Weight loss
- Depression

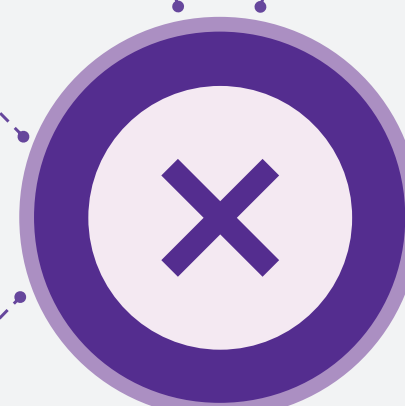
2 Mimicking disorders typically have one or more key differentiation features, which allow them to be distinguished from PMR^{1,7,8}

Headache, temporal tenderness, jaw claudication, visual disturbances*

Inadequate response to GC therapy

Slow-onset, large-joint pain, or pain in joints of hands and feet, or knees; autoantibody positive

Acute-phase reactant laboratory tests within normal range



Multisystem involvement, antinuclear factor and DNA binding; autoantibodies

Elevation of muscle enzymes (CPK, aldolase), muscle weakness

Altered thyroid function tests

Asymmetrical joint pain or swelling

3 Patients with PMR typically show marked response to GC treatment, sometimes as soon as 1 day after initiation²

EULAR/ACR recommend low-dose GC therapy with gradually tailored tapering³

Initial Dose

12.5–25 mg/day

for 2-4 weeks

Initial Tapering

Taper to 10 mg/day

by 4-8 weeks

Relapse Therapy

Increase to pre-relapse dose and taper gradually

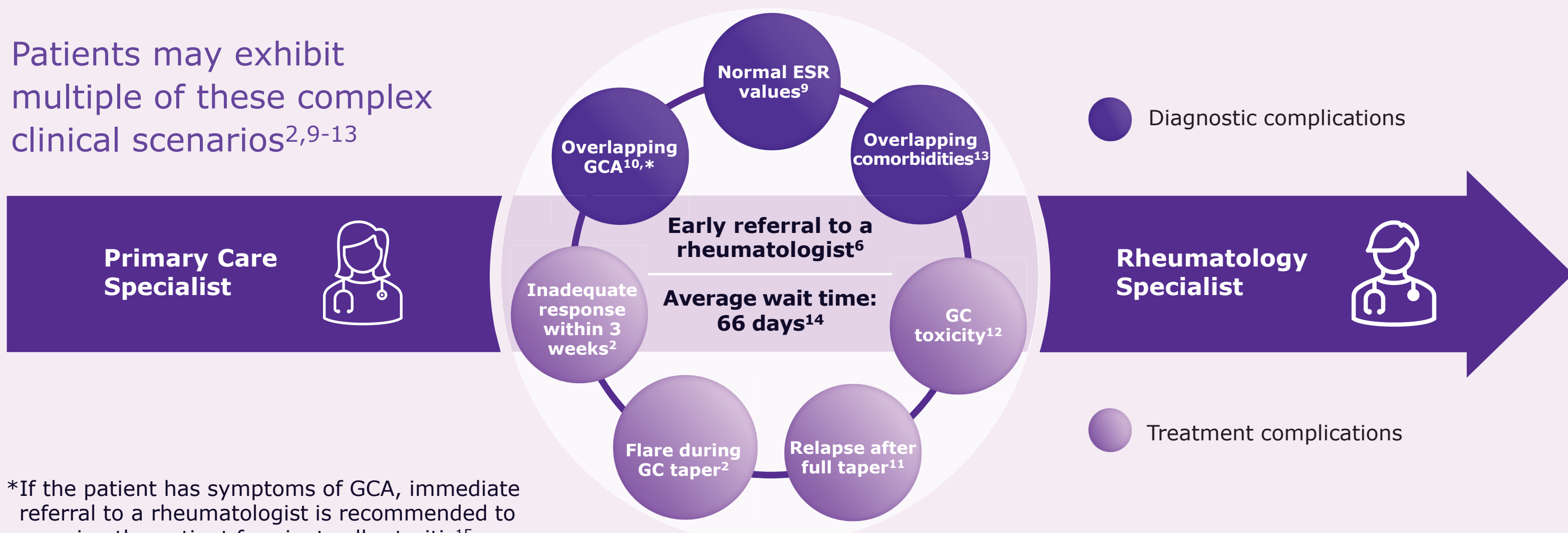
Taper daily by 1 mg every 4 weeks

until discontinuation

Remission

Complex clinical scenarios are common and early referral to a rheumatologist is recommended^{3,7}

Patients may exhibit multiple of these complex clinical scenarios^{2,9-13}



*If the patient has symptoms of GCA, immediate referral to a rheumatologist is recommended to examine the patient for giant cell arteritis¹⁵

Once referred, rheumatologists may recommend the use of alternative treatment options for PMR^{2,16}

References: 1. Nothnagl T, et al. *Drugs Aging*. 2006;23(5):391-402. 2. Leung JL, et al. *J Pharm Pres*. 2019;49:493-500. 3. Dejaco C, et al. *Arthritis Rheumatol*. 2015;67(10):2569-2580. 4. Dasgupta B, et al. *Ann Rheum Dis*. 2012;71(4):484-492. 5. Castañeda S, et al. *Biochem Pharmacol*. 2019;165:221-229. 6. Unwin B, et al. *Am Fam Physician*. 2006;74(9):1547-1554. 7. Dasgupta B, et al. *Rheumatology (Oxford)*. 2010;49(1):186-190. 8. Mahmood SB, et al. *Cleve Clin J Med*. 2020;87(9):549-556. 9. Cantini F, et al. *Semin Arthritis Rheum*. 2000;30(1):17-24. 10. Gonzalez-Gay MA, et al. *Lancet*. 2017;390(10103):1700-1712. 11. Floris A, et al. *Clin Rheumatol*. 2022;41(1):19-31. 12. Cimmino MA, et al. *BMC Musculoskelet Disord*. 2011;12(1):94. 13. Partington R, et al. *Semin Arthritis Rheum*. 2020;50(4):663-672. 14. Widdifield J, et al. *CMAJ Open*. 2016;4(2):E205-E212. 15. Prior JA, et al. *BMC Medicine*. 2017;15:120. 16. Baker E. *Br J Clin Pharmacol*. 2021; 87: 12–22.