

Induction and Tapering Therapy With Tofacitinib and Glucocorticoid in Patients With Polymyalgia Rheumatica

NCT06172361

Status	RECRUITING
Phase	Phase 3
Sponsor	Zhejiang University
Enrollment	98 participants

Key Eligibility Criteria

Inclusion (5)

- PMR patients who fulfilled the 1982 Chuang criteria or 2012ACR/EULAR criteria for PMR; They did not receive any glucocorticoids or biological agents during the 2 weeks period that preceded their inclusion in the study;
- Patients with high activity rheumatic polymyalgia: disease activity score PMR-AS (Table 3) ≥ 10 ,
- Adults age 50-88, Weight 45-85Kg,
- ESR ≥ 20 mm/h or CRP ≥ 50 mg/L (5mg/dl),
- Informed consent.

Exclusion (15)

- Patients with known allergies to tofacitinib, prednisone or methylprednisolone,
- Patients with identified giant cell arteritis, systemic lupus erythematosus, rheumatoid arthritis, calcium pyrophosphate deposition (CPPD) arthropathy, and other rheumatic diseases,
- Patients with severe osteoarthritis,
- Subjects with any severe acute, chronic or recurrent infection (e.g. pneumonia or pyelonephritis, recurrent pneumonia, chronic bronchiectasis, tuberculosis, etc.),
- Hepatitis B virus carriers or individuals with chronic active hepatitis B or C, other chronic liver diseases, HIV infection,
- ... and 10 more (see full listing online)

Locations (1 total)

Division of Rheumatology, the First Affiliated Hospital, Zhejiang University School of Medicine, Hangzhou, Zhejiang, China