### **GPwER: Rheumatology competencies**

The following is based in the Joint Royal Colleges of Physicians Training Board curriculum for rheumatology training draft speciality capabilities in practice (CiP) and practical procedures (p29-33), adapted for the proposed GPwER in MSK and rheumatology accreditation.

The JRCPTB curriculum in turn is based on the generic professional capabilities framework (GPC) introduced by the GMC in 2017, in which the patient is at the centre of any consultation and decision making

#### Generic professional capabilities framework - GMC (gmc-uk.org)

The table below details the key presentations and conditions of Rheumatology.

Particular presentations, conditions and issues are listed either because they are common or serious (having high morbidity, mortality and/or serious implications for treatment or public health).

For each condition/presentation, GPwERs will need to be familiar with such aspects as aetiology, epidemiology, clinical features, investigation, management and prognosis. Our approach is to provide general guidance and not exhaustive detail, which would inevitably become out of date.

Treatment care and strategy includes drug treatments or other interventions for a patient. It includes discussions and decisions as to whether care is focused mainly on curative intent or whether the main focus is on symptomatic relief. It also covers broader aspects of care, including involvement of other professionals or services. It is expected that having completed core GP training, that GPwERs will demonstrate core bedside skills, including information gathering through history and physical examination and information sharing with patients, families and colleagues.

The exact treatment care and strategy approach adopted by the GPwER will depend on the service and tier in which the GPwER is working. However, as a requirement for GPwER accreditation the individual will need to demonstrate an awareness of these conditions, the basis on which diagnosis is made and basic first line management. It is also expected that the GPwER will understand when to refer to a rheumatologist and the urgency of referral.

<u>Clinical area</u>	<b>Presentations</b>	<u>Conditions/Issues</u>
Inflammatory arthritis	Monoarthritis	Septic arthritis
-	Polyarthritis	Gout/Pseudogout
		Chronic infectious arthritis – Mycobacterial arthritis, Lyme disease
		Viral arthritis – Parvo, Hepatitis and HIV-associated arthritis
		Reactive arthritis
		Pigmented Villonodular Synovitis
		Psoriatic arthritis
		Rheumatoid arthritis
		Unclassified inflammatory arthritis
		Arthritis associated with immunodeficiency
		Sarcoidosis – Lofgren's syndrome
		Palindromic arthritis
Spondyloarthropathy	Inflammatory back pain	Axial Spondyloarthropathy (AxSpA)
	Oligoarthritis	(Radiographic/NonRadiographic)
	Enthesitis	Peripheral manifestations of AxSpA
	Dactylitis	IBD associated arthropathy/SpA
		Reactive arthritis
		Undifferentiated Spondyloarthropathy
		Whipple's disease

<u>Clinical area</u>	<u>Presentations</u>	<u>Conditions/Issues</u>
Connective tissue diseases	Facial rashes Discoid rash Renal disorders Scleroderma and Raynaud's Haematological disorder Neurological disorders including peripheral and central syndromes Thrombophilia Sicca syndrome Salivary/Lacrimal gland swelling Lymphadenopathy Muscle weakness with or without rash Serositis	SLE Cutaneous LE SLE-associated nephritis Sjogren's syndrome Systemic sclerosis and associated conditions Inflammatory myopathies Overlap syndromes Antiphospholipid antibody syndrome
Vasculitis	Pulmonary-renal syndromes Systemic illness with multiorgan disease Rash and arthritis/nephritis/lung disease Uveitis Scleritis Deafness – sensorineural External ear disease	ANCA-associated vasculitis Granulomatosis with Polyangiitis (GPA), Eosinophilic Granulomatosis with Polyangiitis (EGPA), Microscopic Polyangiitis (MPA) Non-ANCA Vasculitis – Polyarteritis Nodosa (PAN) Behcet's disease Large Vessel Vasculitis -Takayasu's arteritis, Giant Cell Arteritis Leukocytoclastic vasculitisIgA Vasculitis Cryoglobulinemia Relapsing polychondritis
Auto-inflammatory disorders	Pyrexia of unknown origin Fever and rash	Periodic fever syndromes Familial Mediterranean fever Adult-onset Still's disease

	Fever with multi-organ dysfunction Serositis	Macrophage activation syndrome and HLH Amyloidosis Sweet's syndrome
Multi system disease – others	Lymphadenopathy Granulomatous diseases Retroperitoneal fibrosis Immunodeficiency Inflammatory eye disease	Sarcoidosis Castleman's disease/Histiocytic syndromes IgG4 disease Uveitis Scleritis
Bone disease	Pathological fracture Insufficiency fracture Stress fracture Bone pain Laboratory abnormalities of calcium, phosphate, alkaline phosphatase Incidental radiographic abnormalities	Osteoporosis Osteomalacia Postmenopausal osteoporosis Male osteoporosis Paget's disease of the bone Osteonecrosis Atypical femoral fractures Transient regional osteoporosis
Endocrine and metabolic disorders	Complications of diabetes Complications of thyroid disease Calcinosis	Diabetic stiff hand Thyroid acropachy Haemochromatosis-associated arthropathy Alkaptonuria Neuropathic arthropathy
Neoplastic disorders	Soft tissue swelling Imaging abnormalities of bone and soft tissues Cancer therapy associated syndromes Paraneoplastic syndromes	Sarcomas Primary bone tumours Hypertrophic Pulmonary Osteopathy (HPOA) Graft-versus-host disease (GVHD) Aromatase inhibitor-associated disorder Checkpoint inhibitor-associated disorder

Spinal musculoskeletal pain disorders	Neck pain Back pain Sciatica	Osteoarthritis Disc disease Foraminal stenosis Radiculopathy Myelopathy Cauda equina syndrome
Regional musculoskeletal soft tissue disorders	Rotator cuff disease Enthesopathies Bursitis Entrapment neuropathies Occupational and sports- related problems	Osteoarthritis Calcific tendinitis Epicondylitis, plantar fasciitis Knee and elbow bursitis Carpal tunnel syndrome Greater trochanteric pain syndrome
Pain syndromes	Widespread generalised pain Non-specific limb pain Chest wall pain syndromes	Complex regional pain syndromes – algodystrophy Fibromyalgia and related somatoform disorders
Paediatric and adolescent rheumatological disease	Inflammatory arthritis Connective tissue disorders Pain problems specific to childhood	Juvenile Idiopathic Arthritis (JIA subtypes) Differences between juvenile vs adult Connective Tissue Disorders (CTDs) Macrophage Activation Syndrome (MAS) Transitional care Uveitis Joint hypermobility and spectrum disorders Osgood-Schlatter's disease Perthe's disease Chronic non-bacterial osteomyelitis

# **Other Clinical Syndromes**

Rheumatologic problems in pregnancy Physical symptoms unexplained by organic disease

### **3.6 Practical procedures**

There are a number of procedural skills in which a trainee must become proficient.

Trainees must be able to outline the indications for these procedures and recognise the importance of valid consent, aseptic technique, safe use of analgesia and local anaesthetics, minimisation of patient discomfort, and requesting help when appropriate. For all practical procedures the trainee must be able to recognise complications and respond appropriately if they arise, including calling for help from colleagues in other specialties when necessary.

Trainees should receive training in procedural skills in a clinical skills lab if required. Assessment of procedural skills will be made using the direct observation of procedural skills (DOPS) tool. The table below sets out the minimum competency level expected for each of the practical procedures. When a trainee has been signed off as being able to perform a procedure independently, they are not required to have any further assessment (DOPS) of that procedure, unless they or their educational supervisor think that this is required (in line with standard professional conduct).

Procedure Minimum level required Mandatory	ST4	ST5	ST6	ST7
Large joint knee, shoulder	Competent to perform unsupervised	Maintain	Maintain	Maintain
Medium joints wrist, elbow and ankle	Competent to perform unsupervised	Maintain	Maintain	
Small joints metacarpophalangeal MCP, MTP, PIP		Competent to perform unsupervised		
Procedure	ST4	ST5	ST6	ST7
Soft tissue injections — bursa, tendon sheath, plantar fascia, epicondylitis, carpal tunnel	Competent to perform unsupervised	Maintain	Maintain	Maintain

Nail-fold capillaroscopySkills labMaintainPolarising microscopy of synovial fluid for<br/>crystalsSkills labMaintain

## Recommended

Ultrasound-guided joint or soft tissue injections Fluoroscopy-guided injections